

2017 Aetna Pharmacy Drug Guide - Self Insured
Abilify

Products Affected

- ABILIFY ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Schizophrenia, Bipolar Disorder, Major Depressive Disorder (MDD), Autistic Disorder, Tourette's Disorder
Exclusion Criteria	
Required Medical Information	A Documented diagnosis of Schizophrenia, Bipolar Disorder, Major Depressive Disorder (MDD), Autistic Disorder, or Tourette's Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	FOR A DIAGNOSIS OF BIPOLAR DISORDER OR SCHIZOPHRENIA: A documented contraindication, intolerance, allergy, or failure of one generic medication (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) or Latuda. FOR ALL OTHER DIAGNOSIS: A documented contraindication, intolerance, allergy, or failure of one month of one generic medication (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone)
QL Criteria	1 tab Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Absorica

Products Affected

- ABSORICA

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Abstral

Products Affected

- ABSTRAL

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of the preferred generic alternative, fentanyl transmucosal lozenge, and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 tablets Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Accolate

Products Affected

- ACCOLATE

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Accu-Chek Aviva Plus

Products Affected

- ACCU-CHEK AVIVA PLUS IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Accu-Chek Compact Plus

Products Affected

- ACCU-CHEK COMPACT PLUS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Accu-Chek SmartView

Products Affected

- ACCU-CHEK SMARTVIEW

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Accutrend Glucose

Products Affected

- ACCUTREND GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acetaminophen-Codeine

Products Affected

- *acetaminophen-codeine oral solution*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acetaminophen-Codeine

Products Affected

- *acetaminophen-codeine oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acetaminophen-Codeine #2

Products Affected

- *acetaminophen-codeine #2*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acetaminophen-Codeine #3

Products Affected

- *acetaminophen-codeine #3*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acetaminophen-Codeine #4

Products Affected

- *acetaminophen-codeine #4*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aciphex

Products Affected

- ACIPHEX

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 tab Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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AcipHex Sprinkle

Products Affected

- ACIPHEX SPRINKLE

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required). In addition for approval the following criteria must also be met: Documentation of an inability to swallow tablets/capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 caps Per 1 Day
Notes/References	Annual Review: 02/2017

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Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Acitretin

Products Affected

- *acitretin*

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actemra

Products Affected

- ACTEMRA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Actemra.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Actemra.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actimmune

Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/actimmune.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actiq

Products Affected

- ACTIQ BUCCAL LOZENGE ON A
HANDLE 1200 MCG, 1600 MCG, 400
MCG, 600 MCG, 800 MCG

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 lozenges Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actiq

Products Affected

- ACTIQ BUCCAL LOZENGE ON A
HANDLE 200 MCG

QL Criteria	120 lozenges Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Activella

Products Affected

- ACTIVELLA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actonel

Products Affected

- ACTONEL ORAL TABLET 150 MG

QL Criteria	1 tablet Per 30 Days
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actonel

Products Affected

- ACTONEL ORAL TABLET 30 MG
- ACTONEL ORAL TABLET 5 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actonel

Products Affected

- ACTONEL ORAL TABLET 35 MG

QL Criteria	1 tab Per 7 Days
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adcirca

Products Affected

- ADCIRCA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adderall

Products Affected

- ADDERALL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adderall XR

Products Affected

- ADDERALL XR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adempas

Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adlyxin

Products Affected

- ADLYXIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
QL Criteria	2 pens Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adlyxin Starter Pack

Products Affected

- ADLYXIN STARTER PACK

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
QL Criteria	1 kit Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advair Diskus

Products Affected

- ADVAIR DISKUS

QL Criteria	1 disk Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advair HFA

Products Affected

- ADVAIR HFA

QL Criteria	1 inhaler Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advance Intuition Test

Products Affected

- ADVANCE INTUITION TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advance Micro-Draw Test

Products Affected

- ADVANCE MICRO-DRAW TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advate

Products Affected

- ADVATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advocate Redi-Code

Products Affected

- ADVOCATE REDI-CODE IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advocate Redi-Code+ Test

Products Affected

- ADVOCATE REDI-CODE+ TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advocate Test

Products Affected

- ADVOCATE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adynovate

Products Affected

- *adynovate*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adyphren

Products Affected

- ADYPHREN

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adyphren Amp II

Products Affected

- ADYPHREN AMP II

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adyphren II

Products Affected

- ADYPHREN II

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Adzenys XR-ODT

Products Affected

- ADZENYS XR-ODT

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aerospan

Products Affected

- AEROSPAN

QL Criteria	1 inhaler Per 1 month
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afinitor

Products Affected

- AFINITOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afinitor Disperz

Products Affected

- AFINITOR DISPERZ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afrezza

Products Affected

- AFREZZA INHALATION POWDER 12 UNIT, 8 UNIT

PA Criteria	Criteria Details
Covered Uses	Type 1 Diabetes, Type 2 Diabetes
Exclusion Criteria	
Required Medical Information	Documentation of ALL of the following: (1) In patients with type 1 diabetes, concomitant use of long-acting insulin, (2) In all Patients, no history of chronic lung disease such as asthma or Chronic Obstructive Pulmonary Disease (COPD), and (3) Detailed medical history documenting physical examination and spirometry (FEV1) to identify potential lung disease in all patients.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 24, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afrezza

Products Affected

- AFREZZA INHALATION POWDER 4 & 8 & 12 UNIT, 4 (30) & 8 (60) UNIT, 4 (90) & 8 (90) UNIT, 4 UNIT, 8 (60)& 12 (30) UNIT

PA Criteria	Criteria Details
Covered Uses	Type 1 Diabetes, Type 2 Diabetes
Exclusion Criteria	
Required Medical Information	Documentation of ALL of the following: (1) In patients with type 1 diabetes, concomitant use of long-acting insulin, (2) In all Patients, no history of chronic lung disease such as asthma or Chronic Obstructive Pulmonary Disease (COPD), and (3) Detailed medical history documenting physical examination and spirometry (FEV1) to identify potential lung disease in all patients.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: February 24, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Afrezza

Products Affected

- AFREZZA INHALATION POWDER 4 (60) & 8 (30) UNIT

PA Criteria	Criteria Details
Covered Uses	Type 1 Diabetes, Type 2 Diabetes
Exclusion Criteria	
Required Medical Information	Documentation of ALL of the following: (1) In patients with type 1 diabetes, concomitant use of long-acting insulin, (2) In all Patients, no history of chronic lung disease such as asthma or Chronic Obstructive Pulmonary Disease (COPD), and (3) Detailed medical history documenting physical examination and spirometry (FEV1) to identify potential lung disease in all patients.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 02/2016
Revision Date	Prior Authorization: February 24, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afstyla

Products Affected

- AFSTYLA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AgaMatrix AMP Test

Products Affected

- AGAMATRIX AMP TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AgaMatrix Jazz Test

Products Affected

- AGAMATRIX JAZZ TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AgaMatrix KeyNote Test

Products Affected

- AGAMATRIX KEYNOTE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AgaMatrix Presto Test

Products Affected

- AGAMATRIX PRESTO TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AirDuo RespiClick 113/14

Products Affected

- AIRDUO RESPICLICK 113/14

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Advair, Breo, Dulera, Symbicort, and propionate/salmeterol inhaler (generic Airduo)
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: May 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AirDuo RespiClick 232/14

Products Affected

- AIRDUO RESPICLICK 232/14

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Advair, Breo, Dulera, Symbicort, and propionate/salmeterol inhaler (generic Airduo)
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: May 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AirDuo RespiClick 55/14

Products Affected

- AIRDUO RESPICLICK 55/14

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Advair, Breo, Dulera, Symbicort, and propionate/salmeterol inhaler (generic Airduo)
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: May 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Akynzeo

Products Affected

- AKYNZEO

PA Criteria	Criteria Details
Covered Uses	Prophylaxis of chemotherapy-induced nausea and vomiting
Exclusion Criteria	
Required Medical Information	A documented diagnosis of nausea and vomiting associated with cancer chemotherapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For coverage of additional quantities, a member's treating physician must request prior authorization through the Pharmacy Management Precertification Unit. Additional quantities of Akynzeo will be considered medically necessary for those members who have a documented chemotherapy regimen that requires more than two cycles of antiemetic per 30 days
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of a generic 5-HT3 receptor antagonist, such as granisetron or ondansetron, and one month of aprepitant
QL Criteria	2 capsules Per 1 month
Notes/References	Annual Review: 03/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Albenza

Products Affected

- ALBENZA

QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aldara

Products Affected

- ALDARA

QL Criteria	48 packets Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aldurazyme

Products Affected

- ALDURAZYME

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alecensa

Products Affected

- ALECENSA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alendronate Sodium

Products Affected

- *alendronate sodium oral tablet 10 mg, 40 mg, 5 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alendronate Sodium

Products Affected

- *alendronate sodium oral tablet 35 mg, 70 mg*

QL Criteria	1 tab Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alfuzosin HCl ER

Products Affected

- *alfuzosin hcl er*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alinia

Products Affected

- ALINIA ORAL SUSPENSION
RECONSTITUTED

QL Criteria	180 ml Per 3 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alinia

Products Affected

- ALINIA ORAL TABLET

QL Criteria	6 tablets Per 3 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Almotriptan Malate

Products Affected

- *almotriptan malate*

QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alogliptin Benzoate

Products Affected

- *alogliptin benzoate*

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alogliptin-Metformin HCl

Products Affected

- *alogliptin-metformin hcl*

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alogliptin-Pioglitazone

Products Affected

- *alogliptin-pioglitazone*

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alora

Products Affected

- ALORA

QL Criteria	8 patch Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alosetron HCl

Products Affected

- *alosetron hcl*

PA Criteria	Criteria Details
Covered Uses	Irritable bowel syndrome
Exclusion Criteria	
Required Medical Information	(1)A female patient with a diagnosis of severe* irritable bowel syndrome (IBS) with primary symptom of diarrhea with chronic IBS symptoms (generally lasting 6 months or longer), and (2) anatomic or biochemical abnormalities of the gastrointestinal tract have been excluded, and (3) failure to respond to diphenoxylate/atropine and loperamide for at least one month each.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	*Diarrhea-predominant IBS is severe if it includes diarrhea and one or more of the following: (1) frequent and severe abdominal pain/discomfort, or (2) frequent urgency or fecal incontinence, or (3) disability or restriction of daily activities due to IBS.
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each diphenoxylate/atropine and loperamide
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alphanate/VWF Complex/Human

Products Affected

- ALPHANATE/VWF COMPLEX/HUMAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AlphaNine SD

Products Affected

- ALPHANINE SD

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ALPRAZolam ER

Products Affected

- *alprazolam er*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ALPRAZolam XR

Products Affected

- *alprazolam xr*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alprolix

Products Affected

- ALPROLIX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Altprev

Products Affected

- ALTOPREV ORAL TABLET EXTENDED
RELEASE 24 HOUR 20 MG, 60 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Altprev

Products Affected

- ALTOPREV ORAL TABLET EXTENDED
RELEASE 24 HOUR 40 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alunbrig

Products Affected

- ALUNBRIG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Alunbrig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	6 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: June 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Alvesco

Products Affected

- ALVESCO

QL Criteria	1 inhaler Per 1 month
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ambien

Products Affected

- AMBIEN ORAL TABLET 10 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ambien

Products Affected

- AMBIEN ORAL TABLET 5 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ambien CR

Products Affected

- AMBIEN CR

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amerge

Products Affected

- AMERGE

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amitiza

Products Affected

- AMITIZA

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amlodipine Besylate-Valsartan

Products Affected

- *amlodipine besylate-valsartan*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amlodipine-Olmesartan

Products Affected

- *amlodipine-olmesartan*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amlodipine-Valsartan-HCTZ

Products Affected

- *amlodipine-valsartan-hctz*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amnesteem

Products Affected

- AMNESTEEM

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amphetamine-Dextroamphet ER

Products Affected

- *amphetamine-dextroamphet er*

QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amphetamine-Dextroamphetamine

Products Affected

- *amphetamine-dextroamphetamine*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ampyra

Products Affected

- AMPYRA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Amrix

Products Affected

- AMRIX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of two preferred alternatives (one of which should be cyclobenzaprine or cyclobenzaprine er)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Androderm

Products Affected

- ANDRODERM TRANSDERMAL PATCH
24 HOUR

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	1 patch Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AndroGel

Products Affected

- ANDROGEL TRANSDERMAL GEL 20.25 MG/1.25GM (1.62%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	30 packs Per 30 Days
Notes/References	Annual Review: 02/2017

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Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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AndroGel

Products Affected

- ANDROGEL TRANSDERMAL GEL 25 MG/2.5GM (1%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	2.5 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AndroGel

Products Affected

- ANDROGEL TRANSDERMAL GEL 40.5 MG/2.5GM (1.62%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	60 packs Per 30 Days
Notes/References	Annual Review: 02/2017

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Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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AndroGel

Products Affected

- ANDROGEL TRANSDERMAL GEL 50 MG/5GM (1%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	10 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

AndroGel Pump

Products Affected

- ANDROGEL PUMP TRANSDERMAL
GEL 20.25 MG/ACT (1.62%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 pumps Per 30 Days
Notes/References	Annual Review: 02/2017

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Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Angeliq

Products Affected

- ANGELIQ

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Angeliq

Products Affected

- ANGELIQ

QL Criteria	1 tbalet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Anoro Ellipta

Products Affected

- ANORO ELLIPTA

QL Criteria	2 aerosols Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Antara

Products Affected

- ANTARA ORAL CAPSULE 30 MG, 90 MG

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Anzemet

Products Affected

- ANZEMET ORAL

QL Criteria	5 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

APAP-Caff-Dihydrocodeine

Products Affected

- *apap-caff-dihydrocodeine oral capsule*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

APAP-Caff-Dihydrocodeine

Products Affected

- *apap-caff-dihydrocodeine oral tablet 325-30-16 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Apidra

Products Affected

- APIDRA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Apidra SoloStar

Products Affected

- APIDRA SOLOSTAR SUBCUTANEOUS SOLUTION PEN-INJECTOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aplenzin

Products Affected

- APLENZIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of any of the following medications: bupropion SR/XL, bupropion/SR/XL, citalopram, escitalopram, fluoxetine, fluvoxamine, paroxetine/sr, mirtazapine, selfemra, sertraline, venlafaxine, or venlafaxine sr cap
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aprepitant

Products Affected

- *aprepitant oral capsule 125 mg, 40 mg, 80 mg*

QL Criteria	5 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aprepitant

Products Affected

- *aprepitant oral capsule 80 & 125 mg*

QL Criteria	9 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Apriso

Products Affected

- APRISO

QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aptensio XR

Products Affected

- APTENSIO XR

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 capsule Per 1 Day
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aralast NP

Products Affected

- ARALAST NP INTRAVENOUS SOLUTION RECONSTITUTED 1000 MG, 500 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Alpha-1 Antitrypsin Inhibitor Therapy.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aranesp (Albumin Free)

Products Affected

- ARANESP (ALBUMIN FREE) INJECTION SOLUTION 100 MCG/ML, 200 MCG/ML, 25 MCG/ML, 300 MCG/ML, 40 MCG/ML, 60 MCG/ML SOLUTION PREFILLED SYRINGE 100 MCG/0.5ML, 150 MCG/0.3ML, 200 MCG/0.4ML, 25 MCG/0.42ML, 300 MCG/0.6ML, 40 MCG/0.4ML, 500 MCG/ML, 60 MCG/0.3ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Erythropoiesis_Stimulating_Agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Arcalyst

Products Affected

- ARCALYST

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Arcalyst.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Arcapta Neohaler

Products Affected

- ARCAPTA NEOHALER

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ARIPiprazole

Products Affected

- *aripiprazole oral solution*

QL Criteria	30 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ARIPiprazole

Products Affected

- *aripiprazole oral tablet*
- *aripiprazole oral tablet dispersible*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Armodafinil

Products Affected

- *armodafinil oral tablet 150 mg*
- *armodafinil oral tablet 200 mg, 250 mg*

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with armodafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tablet Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Armodafinil

Products Affected

- *armodafinil oral tablet 50 mg*

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with armodafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tablets Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ArmonAir RespiClick 113

Products Affected

- ARMONAIR RESPICLICK 113

PA Criteria	Criteria Details
Covered Uses	Maintenance treatment of asthma as prophylactic therapy in patients 12 years of age and older.
Exclusion Criteria	Not indicated for the relief of acute bronchospasm
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of QVAR, Asmanex, and Flovent
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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ArmonAir RespiClick 232

Products Affected

- ARMONAIR RESPICLICK 232

PA Criteria	Criteria Details
Covered Uses	Maintenance treatment of asthma as prophylactic therapy in patients 12 years of age and older.
Exclusion Criteria	Not indicated for the relief of acute bronchospasm
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of QVAR, Asmanex, and Flovent
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ArmonAir RespiClick 55

Products Affected

- ARMONAIR RESPICLICK 55

PA Criteria	Criteria Details
Covered Uses	Maintenance treatment of asthma as prophylactic therapy in patients 12 years of age and older.
Exclusion Criteria	Not indicated for the relief of acute bronchospasm
Required Medical Information	A documented diagnosis of Asthma
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of QVAR, Asmanex, and Flovent
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Arnuity Ellipta

Products Affected

- ARNUITY ELLIPTA

QL Criteria	1 blister Per 1 Day
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Arymo ER

Products Affected

- ARYMO ER

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred alternatives which include Hysingla ER, Embeda and Oxycontin
QL Criteria	90 tablets Per 30 days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Asacol HD

Products Affected

- ASACOL HD

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ascomp-Codeine

Products Affected

- ASCOMP-CODEINE

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure 3 Test

Products Affected

- ASSURE 3 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure 4 Test

Products Affected

- ASSURE 4 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure II

Products Affected

- ASSURE II

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure II Check

Products Affected

- ASSURE II CHECK

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure Platinum

Products Affected

- ASSURE PLATINUM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Assure Pro Test

Products Affected

- ASSURE PRO TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

At Last Test

Products Affected

- AT LAST TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atacand

Products Affected

- ATACAND ORAL TABLET 16 MG
- ATACAND ORAL TABLET 4 MG, 8 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atacand

Products Affected

- ATACAND ORAL TABLET 32 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atacand HCT

Products Affected

- ATACAND HCT ORAL TABLET 16-12.5
MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atacand HCT

Products Affected

- ATACAND HCT ORAL TABLET 32-12.5 MG, 32-25 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atelvia

Products Affected

- ATELVIA

QL Criteria	1 tab Per 7 Days
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ativan

Products Affected

- ATIVAN ORAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of lorazepam and two other benzodiazepines
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atomoxetine HCl

Products Affected

- *atomoxetine hcl oral capsule 10 mg, 18 mg, 25 mg, 40 mg, 60 mg*

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atomoxetine HCl

Products Affected

- *atomoxetine hcl oral capsule 100 mg, 80 mg*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atorvastatin Calcium

Products Affected

- *atorvastatin calcium oral*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atovaquone-Proguanil HCl

Products Affected

- *atovaquone-proguanil hcl oral tablet 250-100 mg*

PA Criteria	Criteria Details
Covered Uses	Malaria
Exclusion Criteria	
Required Medical Information	A documented diagnosis of malaria
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Maximum Approval for Malaria: 30 days, Maximum Approval for all other indications: 1 year
Other Criteria	Malaria medications are Certificate of Coverage (COC) Excluded for travel prophylaxis
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Atralin

Products Affected

- ATRALIN

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of generic Atralin and Retin-A
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Atrovent HFA

Products Affected

- ATROVENT HFA

QL Criteria	2 inhalers Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aubagio

Products Affected

- AUBAGIO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Austedo

Products Affected

- AUSTEDO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Austedo.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Austedo.html
QL Criteria	4 tablets Per 1 day
Notes/References	
Revision Date	Prior Authorization: May 09, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Auvi-Q

Products Affected

- AUVI-Q INJECTION SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Covered Uses	Emergency treatment of allergic reactions
Exclusion Criteria	
Required Medical Information	A documented diagnosis of an allergic reaction in patients who are at risk for or have a history of anaphylactic reaction and the individual or their caregiver requires an auto injector with audio or visual cues for administration
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic epinephrine (Adrenaclick), generic epinephrine (Epipen), and Epipen
QL Criteria	4 pens Per 1 month
Notes/References	
Revision Date	Prior Authorization: February 07, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Avalide

Products Affected

- AVALIDE ORAL TABLET 150-12.5 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avalide

Products Affected

- AVALIDE ORAL TABLET 300-12.5 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avapro

Products Affected

- AVAPRO

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avapro

Products Affected

- AVAPRO

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avita

Products Affected

- AVITA

PA Criteria	Criteria Details
Covered Uses	acne vulgaris
Exclusion Criteria	
Required Medical Information	Documented diagnosis of acne vulgaris for members greater than 35 years old
Age Restrictions	Greater than 35. Members under age 35 can obtain medication without authorization.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avodart

Products Affected

- AVODART

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avonex

Products Affected

- AVONEX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avonex Pen

Products Affected

- AVONEX PEN INTRAMUSCULAR
AUTO-INJECTOR KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avonex Prefilled

Products Affected

- AVONEX PREFILLED
INTRAMUSCULAR PREFILLED
SYRINGE KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Axert

Products Affected

- AXERT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan
QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Azilect

Products Affected

- AZILECT

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Azor

Products Affected

- AZOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives, which include Exforge and any of the following medications used in combination with amlodipine: candesartan, eprosartan, irbesartan, losartan, valsartan, or telmisartan
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Azulfidine

Products Affected

- AZULFIDINE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	8 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Azulfidine EN-tabs

Products Affected

- AZULFIDINE EN-TABS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	8 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bactroban

Products Affected

- BACTROBAN EXTERNAL CREAM

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Balsalazide Disodium

Products Affected

- *balsalazide disodium*

QL Criteria	9 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Banzel

Products Affected

- BANZEL ORAL TABLET

QL Criteria	8 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Baxdela

Products Affected

- BAXDELA ORAL

PA Criteria	Criteria Details
Covered Uses	Treatment of acute bacterial skin and skin structure infections (ABSSSI) caused by designated susceptible bacteria
Exclusion Criteria	Known hypersensitivity to Baxdela or other fluoroquinolones
Required Medical Information	A documented diagnosis of acute bacterial skin and skin structure infections (ABSSSI) caused by one the following susceptible pathogens: Gram-positive organisms include Staphylococcus aureus (including methicillin-resistant [MRSA] and methicillinsusceptible [MSSA] isolates), Staphylococcus haemolyticus, Staphylococcus lugdunensis, Streptococcus agalactiae, Streptococcus anginosus Group (including Streptococcus anginosus, Streptococcus intermedius, and Streptococcus constellatus), Streptococcus pyogenes, or Enterococcus faecalis. Gram-negative organisms include: Escherichia coli, Enterobacter cloacae, Klebsiella pneumoniae, and Pseudomonas aeruginosa.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	28 tablets Per 1 fill
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: November 15, 2017

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Bayer Breeze 2 Test

Products Affected

- BAYER BREEZE 2 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bayer Contour Next Test

Products Affected

- BAYER CONTOUR NEXT TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bayer Contour Test

Products Affected

- BAYER CONTOUR TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bebulin

Products Affected

- BEBULIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Beconase AQ

Products Affected

- BECONASE AQ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Belbuca

Products Affected

- BELBUCA

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	2 films Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Belsomra

Products Affected

- BELSOMRA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of one generic alternative: zolpidem, zolpidem er, eszopiclone, or zaleplon
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Benicar

Products Affected

- BENICAR ORAL TABLET 20 MG
- BENICAR ORAL TABLET 5 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Benicar HCT

Products Affected

- BENICAR HCT

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Benicar HCT

Products Affected

- BENICAR HCT

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Benlysta

Products Affected

- BENLYSTA INTRAVENOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benlysta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benlysta.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Benlysta

Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benlysta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benlysta.html
QL Criteria	4 injections Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

BenzEFoamUltra

Products Affected

- BENZEFOAMULTRA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic benzoyl peroxide foam
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Berinert

Products Affected

- BERINERT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Betamethasone Dipropionate Aug

Products Affected

- *betamethasone dipropionate aug external gel ointment*
- *betamethasone dipropionate aug external*

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Betamethasone Dipropionate Aug

Products Affected

- *betamethasone dipropionate aug external lotion*

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Betaseron

Products Affected

- BETASERON SUBCUTANEOUS KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bevespi Aerosphere

Products Affected

- BEVESPI AEROSPHERE

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of either Anoro Ellipta or Stiolto
QL Criteria	1 inhaler Per 30 Days
Notes/References	
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bevyxxa

Products Affected

- BEVYXXA

PA Criteria	Criteria Details
Covered Uses	Prophylaxis of venous thromboembolism (VTE) in adult patients hospitalized for an acute medical illness who are at risk for thromboembolic complications due to moderate or severe restricted mobility and other risk factors for VTE.
Exclusion Criteria	Active pathological bleeding, severe hypersensitivity reaction to Bevyxxa, or for anyone with prosthetic heart valves.
Required Medical Information	Member is requesting product for use of prophylaxis of VTE and is currently taking Bevyxxa during hospitalization and will be continuing therapy following discharge from the hospital.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of enoxaparin or dalteparin, or heparin
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: October 04, 2017 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

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Bicalutamide

Products Affected

- *bicalutamide*

PA Criteria	Criteria Details
Covered Uses	Covered for FDA approved indications in male members without Prior Authorization
Exclusion Criteria	Pregnancy
Required Medical Information	For coverage in female members, there must be a documented diagnosis of hirsutism secondary to ovarian or adrenal dysfunction (e.g., polycystic ovary syndrome, adrenal or ovarian tumor).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: March 04, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Binosto

Products Affected

- BINOSTO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg or Atelvia
QL Criteria	1 tab Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bioscanner Glucose Test

Products Affected

- BIOSCANNER GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bivigam

Products Affected

- BIVIGAM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Blood Glucose Test

Products Affected

- *blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Boniva

Products Affected

- BONIVA ORAL TABLET 150 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	1 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bosulif

Products Affected

- BOSULIF

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Botox

Products Affected

- BOTOX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/botulinum_toxin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/botulinum_toxin.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bravelle

Products Affected

- BRAVELLE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Breo Ellipta

Products Affected

- BREO ELLIPTA

QL Criteria	2 blisters Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Brilinta

Products Affected

- BRILINTA

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Brilinta

Products Affected

- BRILINTA

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Brisdelle

Products Affected

- BRISDELLE

PA Criteria	Criteria Details
Covered Uses	Moderate to severe vasomotor symptoms associated with menopause
Exclusion Criteria	
Required Medical Information	A documented diagnosis of moderate to severe vasomotor symptoms associated with menopause
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 28, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Briviact

Products Affected

- BRIVIACT ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Partial-onset seizures
Exclusion Criteria	
Required Medical Information	A documented diagnosis of partial-onset seizures and documented concurrent therapy with one of the following: carbamazepine, divalproex dr/er/sprinkle, gabapentin, lamotrigine, levetiracetam/ER, oxcarbazepine, phenytoin, topiramate, valproic acid, or zonisamide
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Brovana

Products Affected

- BROVANA

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Serevent (Step Therapy will not apply to members who have a documented inability to use an inhaler)
QL Criteria	60 vials (120ml) Per 1 fill
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Budesonide

Products Affected

- *budesonide inhalation*

QL Criteria	4 ml Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Budesonide

Products Affected

- *budesonide oral*

QL Criteria	3 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bunavail

Products Affected

- BUNAVAIL BUCCAL FILM 2.1-0.3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	6 films Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bunavail

Products Affected

- BUNAVAIL BUCCAL FILM 4.2-0.7 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	3 films Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bunavail

Products Affected

- BUNAVAIL BUCCAL FILM 6.3-1 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	2 films Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Buphenyl

Products Affected

- BUPHENYL ORAL POWDER 3 GM/TSP
- BUPHENYL ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Buprenorphine

Products Affected

- *buprenorphine*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	4 patches Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Buprenorphine HCl

Products Affected

- *buprenorphine hcl sublingual*

QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Buprenorphine HCl-Naloxone HCl

Products Affected

- *buprenorphine hcl-naloxone hcl*

QL Criteria	90 tab Per 30 Days
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

BuPROPion HCl

Products Affected

- *bupropion hcl oral*

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

BuPROPion HCl ER (Smoking Det)

Products Affected

- *bupropion hcl er (smoking det)*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

BuPROPion HCl ER (SR)

Products Affected

- *bupropion hcl er (sr)*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

BuPROPion HCl ER (XL)

Products Affected

- *bupropion hcl er (xl)*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Butalbital-APAP-Caff-Cod

Products Affected

- *butalbital-apap-caff-cod*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Butalbital-ASA-Caff-Codeine

Products Affected

- *butalbital-asa-caff-codeine*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Butorphanol Tartrate

Products Affected

- *butorphanol tartrate nasal*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	2 bottle Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Butrans

Products Affected

- BUTRANS

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	4 patches Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bydureon

Products Affected

- BYDUREON SUBCUTANEOUS PEN-INJECTOR

QL Criteria	4 pens Per 1 month
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Byetta 10 MCG Pen

Products Affected

- BYETTA 10 MCG PEN SUBCUTANEOUS SOLUTION PEN-INJECTOR

QL Criteria	1 pen Per 30 Days
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Byetta 5 MCG Pen

Products Affected

- BYETTA 5 MCG PEN SUBCUTANEOUS SOLUTION PEN-INJECTOR

QL Criteria	1 pen Per 30 Days
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bystolic

Products Affected

- BYSTOLIC ORAL TABLET 10 MG, 5 MG • BYSTOLIC ORAL TABLET 2.5 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bystolic

Products Affected

- BYSTOLIC ORAL TABLET 20 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Byvalson

Products Affected

- BYVALSON

QL Criteria	30 tablets Per 30 Days
Notes/ References	Annual Review: 08/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cabometyx

Products Affected

- CABOMETYX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Caduet

Products Affected

- CADUET ORAL TABLET 10-10 MG, 10-20 MG, 10-40 MG, 10-80 MG, 5-10 MG, 5-20 MG, 5-40 MG, 5-80 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Calcipotriene-Betameth Diprop

Products Affected

- *calcipotriene-betameth diprop*

QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Calcitonin (Salmon)

Products Affected

- *calcitonin (salmon)*

QL Criteria	1 bottle Per 1 month
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cambia

Products Affected

- CAMBIA

QL Criteria	9 pack Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Canasa

Products Affected

- CANASA

QL Criteria	1 EA Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Candesartan Cilexetil

Products Affected

- *candesartan cilexetil oral tablet 16 mg, 4 mg, 8 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Candesartan Cilexetil

Products Affected

- *candesartan cilexetil oral tablet 32 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Candesartan Cilexetil-HCTZ

Products Affected

- *candesartan cilexetil-hctz oral tablet 16-12.5 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Candesartan Cilexetil-HCTZ

Products Affected

- *candesartan cilexetil-hctz oral tablet 32-12.5 mg, 32-25 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Capecitabine

Products Affected

- *capecitabine*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Capex

Products Affected

- CAPEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic fluocinolone
QL Criteria	120 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Caprelsa

Products Affected

- CAPRELSA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Carac

Products Affected

- CARAC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Carac (fluorouracil) and either Efudex or Aldara
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Carbaglu

Products Affected

- CARBAGLU

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cardizem CD

Products Affected

- CARDIZEM CD ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 120 MG,
180 MG, 240 MG, 360 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of diltiazem ER and two other calcium channel blockers
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cardura XL

Products Affected

- CARDURA XL

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CareOne Blood Glucose Test

Products Affected

- CAREONE BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CareSens N Glucose Test

Products Affected

- CARESENS N GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Carimune NF

Products Affected

- CARIMUNE NF INTRAVENOUS SOLUTION RECONSTITUTED 12 GM, 6 GM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CaroSpir

Products Affected

- CAROSPIR

PA Criteria	Criteria Details
Covered Uses	Treatment of NYHA Class III and Class IV heart failure and reduced ejection fraction, use as an add-on therapy for the treatment of hypertension, and for the management of edema in adult cirrhotic patients when edema is not responsive to fluid and sodium restrictions
Exclusion Criteria	Hyperkalemia, Addisons disease, concomitant use of eplerenone
Required Medical Information	A documented diagnosis of severe heart failure (NYHA class III-IV) and has a left ventricular ejection fraction less than or equal to 35%, a documented diagnosis of hypertension and requested drug is being used as add on therapy, or a documented diagnosis of Edema associated with Cirrhosis, and there is documentation of an inability to swallow tablets/capsules
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic spironolactone tablets
QL Criteria	80 milliliters Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: November 06, 2017 Quantity Limits: November 16, 2017

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Casodex

Products Affected

- CASODEX

PA Criteria	Criteria Details
Covered Uses	Covered for FDA approved indications in male members without Prior Authorization
Exclusion Criteria	Pregnancy
Required Medical Information	For coverage in female members, there must be a documented diagnosis of hirsutism secondary to ovarian or adrenal dysfunction (e.g., polycystic ovary syndrome, adrenal or ovarian tumor).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: March 04, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CeleBREX

Products Affected

- CELEBREX

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CeleBREX

Products Affected

- CELEBREX

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Celecoxib

Products Affected

- *celecoxib oral*

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CeleXA

Products Affected

- CELEXA ORAL TABLET

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Centany

Products Affected

- CENTANY

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cerdelga

Products Affected

- CERDELGA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: ?http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gaucher_disease.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cerezyme

Products Affected

- CEREZYME INTRAVENOUS SOLUTION
RECONSTITUTED 400 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: ?http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gaucher_disease.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cesamet

Products Affected

- CESAMET

QL Criteria	20 caps Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cetrotide

Products Affected

- CETROTIDE SUBCUTANEOUS KIT 0.25 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Chantix

Products Affected

- CHANTIX

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Chantix Continuing Month Pak

Products Affected

- CHANTIX CONTINUING MONTH PAK

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Chantix Starting Month Pak

Products Affected

- CHANTIX STARTING MONTH PAK

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cholbam

Products Affected

- CHOLBAM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Cholbam.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Chorionic Gonadotropin

Products Affected

- *chorionic gonadotropin intramuscular*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cialis

Products Affected

- CIALIS ORAL TABLET 2.5 MG
- CIALIS ORAL TABLET 5 MG

PA Criteria	Criteria Details
Covered Uses	Benign prostatic hyperplasia
Exclusion Criteria	
Required Medical Information	Member has diagnosis of BPH (Benign Prostatic Hyperplasia) and of the following: (1)Member is not currently on nitrite/nitrate therapy, (2)is not currently on another phosphodiesterase-5 inhibitor, or (3) has a documented contraindication, intolerance, allergy, or failure of a one month trial of one of the preferred drugs alfuzosin, finasteride, tamsulosin, Avodart, Jalyn or Rapaflo
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cimzia

Products Affected

- CIMZIA SUBCUTANEOUS KIT 2 X 200
MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cimzia Prefilled

Products Affected

- CIMZIA PREFILLED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cimzia Starter Kit

Products Affected

- CIMZIA STARTER KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cinqair

Products Affected

- CINQAIR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/RESP/Cinqair.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cinryze

Products Affected

- CINRYZE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Citalopram Hydrobromide

Products Affected

- *citalopram hydrobromide oral tablet*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Claravis

Products Affected

- CLARAVIS

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clarinet

Products Affected

- CLARINEX ORAL SYRUP

QL Criteria	10 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clarinet

Products Affected

- CLARINEX ORAL TABLET

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clarinet-D 12 Hour

Products Affected

- CLARINEX-D 12 HOUR

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clever Chek Auto-Code Test

Products Affected

- CLEVER CHEK AUTO-CODE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clever Chek Auto-Code Voice

Products Affected

- CLEVER CHEK AUTO-CODE VOICE IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clever Chek Test

Products Affected

- CLEVER CHEK TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clever Choice Auto-Code Test

Products Affected

- CLEVER CHOICE AUTO-CODE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clever Choice Micro Test

Products Affected

- CLEVER CHOICE MICRO TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Climara

Products Affected

- CLIMARA

QL Criteria	1 patch Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Climara Pro

Products Affected

- CLIMARA PRO

QL Criteria	1 patch Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate

Products Affected

- *clobetasol propionate external cream*
- *clobetasol propionate external ointment*
- *clobetasol propionate external gel*

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate

Products Affected

- *clobetasol propionate external foam*
- *clobetasol propionate external solution*

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate

Products Affected

- *clobetasol propionate external liquid*

QL Criteria	125 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate

Products Affected

- *clobetasol propionate external lotion*
- *clobetasol propionate external shampoo*

QL Criteria	236 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate E

Products Affected

- *clobetasol propionate e*

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobetasol Propionate Emulsion

Products Affected

- *clobetasol propionate emulsion*

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobex

Products Affected

- CLOBEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of generic clobetasol lotion or clobetasol shampoo
QL Criteria	236 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clobex Spray

Products Affected

- CLOBEX SPRAY

QL Criteria	125 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clodan

Products Affected

- CLODAN EXTERNAL SHAMPOO

QL Criteria	236 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cloderm

Products Affected

- CLODERM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred alternative generic medication (any dosage form): fluocinolone acetonide, hydrocortisone valerate, mometasone furoate, or triamcinolone acetonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cloderm Pump

Products Affected

- CLODERM PUMP

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred alternative generic medication (any dosage form): fluocinolone acetonide, hydrocortisone valerate, mometasone furoate, or triamcinolone acetonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CloNIDine HCl ER

Products Affected

- *clonidine hcl er*

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clopidogrel Bisulfate

Products Affected

- *clopidogrel bisulfate oral tablet 75 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CloZAPine

Products Affected

- *clozapine oral tablet 100 mg*

QL Criteria	9 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CloZAPine

Products Affected

- *clozapine oral tablet 200 mg*
- *clozapine oral tablet dispersible 200 mg*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CloZAPine

Products Affected

- *clozapine oral tablet 25 mg, 50 mg*

QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CloZAPine

Products Affected

- *clozapine oral tablet dispersible 150 mg*

QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clozaril

Products Affected

- CLOZARIL ORAL TABLET 100 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	9 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clozaril

Products Affected

- CLOZARIL ORAL TABLET 25 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Coagadex

Products Affected

- COAGADEX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Coartem

Products Affected

- COARTEM

PA Criteria	Criteria Details
Covered Uses	Malaria
Exclusion Criteria	
Required Medical Information	A documented diagnosis of malaria
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Maximum Approval for Malaria: 30 days, Maximum Approval for all other indications: 1 year
Other Criteria	Malaria medications are Certificate of Coverage (COC) Excluded for travel prophylaxis
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Codeine Sulfate

Products Affected

- *codeine sulfate oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Colazal

Products Affected

- COLAZAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	9 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Colcrlys

Products Affected

- COLCRYS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Mitigare and generic colchicine
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

CombiPatch

Products Affected

- COMBIPATCH

QL Criteria	8 patch Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Combivent Respimat

Products Affected

- COMBIVENT RESPIMAT

QL Criteria	2 inhalers Per 1 month
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cometriq (100 mg Daily Dose)

Products Affected

- COMETRIQ (100 MG DAILY DOSE)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cometriq (140 mg Daily Dose)

Products Affected

- COMETRIQ (140 MG DAILY DOSE)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cometriq (60 mg Daily Dose)

Products Affected

- COMETRIQ (60 MG DAILY DOSE)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Concerta

Products Affected

- CONCERTA ORAL TABLET EXTENDED
RELEASE 18 MG, 27 MG, 54 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Concerta

Products Affected

- CONCERTA ORAL TABLET EXTENDED
RELEASE 36 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Copaxone

Products Affected

- COPAXONE SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 20 MG/ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Copaxone

Products Affected

- COPAXONE SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 40 MG/ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cordran

Products Affected

- CORDRAN EXTERNAL TAPE

QL Criteria	1 tape roll Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Corifact

Products Affected

- CORIFACT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Corlanor

Products Affected

- CORLANOR

PA Criteria	Criteria Details
Covered Uses	FDA labeled use for heart failure (see required medical information section)
Exclusion Criteria	
Required Medical Information	Documentation of stable, symptomatic chronic heart failure with left ventricular ejection fraction less than or equal to 35%, who are in sinus rhythm with resting heart rate greater or equal to 70 beats per minute and are on maximally tolerated doses of beta-blockers (such as bisoprolol/bisoprolol-HCTZ, carvedilol, carvedilol CR, metoprolol succinate, metoprolol succinate-HCTZ, or nebivolol) or have a documented contraindication to beta-blocker use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step Therapy
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one of the following drug classes: ACE Inhibitor, ACE Inhibitor/HCTZ combination, Angiotensin-Receptor Blocker, or Angiotensin-Receptor Blocker/HCTZ combination
QL Criteria	2 tablets Per 1 Day
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Cormax Scalp Application

Products Affected

- CORMAX SCALP APPLICATION

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cosentyx

Products Affected

- COSENTYX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cosentyx.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cosentyx.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cosentyx Sensoready Pen

Products Affected

- COSENTYX SENSOREADY PEN
SUBCUTANEOUS SOLUTION AUTO-
INJECTOR 150 MG/ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cosentyx.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cosentyx.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cotellic

Products Affected

- COTELLIC

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	63 tablets Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cotempla XR-ODT

Products Affected

- COTEMPLA XR-ODT

PA Criteria	Criteria Details
Covered Uses	Treatment of Attention Deficit Hyperactivity Disorder (ADHD) in pediatric patients 6 to 17 years of age.
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention Deficit Hyperactivity Disorder (ADHD)
Age Restrictions	Approved for patients 6 to 17 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cozaar

Products Affected

- COZAAR ORAL TABLET 25 MG
- COZAAR ORAL TABLET 50 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Crestor

Products Affected

- CRESTOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic statin medication (atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, or simvastatin)
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cuprimine

Products Affected

- CUPRIMINE ORAL CAPSULE 250 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cutivate

Products Affected

- CUTIVATE EXTERNAL CREAM
- CUTIVATE EXTERNAL LOTION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred generic alternative: betamethasone benzoate, betamethasone dipropionate, betamethasone valerate, desonide, desoximetasone, fluocinolone acetonide, fluticasone, fluocinonide, hydrocortisone butyrate, hydrocortisone valerate, prednicarbate, or triamcinolone acetonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cuvitru

Products Affected

- CUVITRU

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cycloset

Products Affected

- CYCLOSET

QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cymbalta

Products Affected

- CYMBALTA ORAL CAPSULE DELAYED
RELEASE PARTICLES 20 MG

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cymbalta

Products Affected

- CYMBALTA ORAL CAPSULE DELAYED
RELEASE PARTICLES 30 MG

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cymbalta

Products Affected

- CYMBALTA ORAL CAPSULE DELAYED
RELEASE PARTICLES 60 MG

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cystadane

Products Affected

- CYSTADANE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cystaran

Products Affected

- CYSTARAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/EYE/opthalmic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 ml (40 drops) Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Daklinza

Products Affected

- DAKLINZA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Daliresp

Products Affected

- DALIRESP

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disease (COPD)
Exclusion Criteria	
Required Medical Information	A Documented diagnosis of severe COPD associated with chronic bronchitis and a history of exacerbations.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two of the following: Breo, Symbicort, Advair, Anoro, Stiolto, Incruse, or Spiriva
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: July 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dapsone

Products Affected

- *dapsone external*

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: November 16, 2017

Daxbia

Products Affected

- DAXBIA

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Daytrana

Products Affected

- DAYTRANA

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 patch Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Delzicol

Products Affected

- DELZICOL

QL Criteria	12 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Demerol

Products Affected

- DEMEROL ORAL

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Depen Titratabs

Products Affected

- DEPEN TITRATABS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Descovy

Products Affected

- DESCOVY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviral_hiv.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Desloratadine

Products Affected

- *desloratadine*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Desonate

Products Affected

- DESONATE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic desonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Desoxyn

Products Affected

- DESOXYN

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tab Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Desvenlafaxine ER

Products Affected

- *desvenlafaxine er*

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Desvenlafaxine Succinate ER

Products Affected

- *desvenlafaxine succinate er*

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Detrol

Products Affected

- DETROL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Detrol LA

Products Affected

- DETROL LA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dexedrine

Products Affected

- DEXEDRINE ORAL CAPSULE
EXTENDED RELEASE 24 HOUR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	3 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dexilant

Products Affected

- DEXILANT

QL Criteria	1 caps Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dexmethylphenidate HCl

Products Affected

- *dexmethylphenidate hcl*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dexmethylphenidate HCl ER

Products Affected

- *dexmethylphenidate hcl er*

QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dextroamphetamine Sulfate

Products Affected

- *dextroamphetamine sulfate oral solution*

QL Criteria	40 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dextroamphetamine Sulfate

Products Affected

- *dextroamphetamine sulfate oral tablet*

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dextroamphetamine Sulfate ER

Products Affected

- *dextroamphetamine sulfate er*

QL Criteria	3 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diastat AcuDial

Products Affected

- DIASTAT ACUDIAL

QL Criteria	1 pack Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diastat Pediatric

Products Affected

- DIASTAT PEDIATRIC

QL Criteria	1 pack Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

DiaTrue Plus Test

Products Affected

- *diatruue plus test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dibenzyline

Products Affected

- DIBENZYLINE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diclegis

Products Affected

- DICLEGIS

PA Criteria	Criteria Details
Covered Uses	Nausea and vomiting in pregnant women
Exclusion Criteria	
Required Medical Information	A documented diagnosis of nausea and vomiting in a pregnant woman who does not respond to conservative management (i.e. trigger avoidance, small frequent meals, etc) and a documented contraindication, intolerance, allergy, or failure of an adequate trial of one week of any of the following: otc doxylamine, or otc pyridoxine (vit B6), or metoclopramide, or promethazine, or ondansetron
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	4 tab Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 01, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diclofenac Sodium

Products Affected

- *diclofenac sodium transdermal gel 1 %*

QL Criteria	200 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diclofenac Sodium

Products Affected

- *diclofenac sodium transdermal gel 3 %*

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Differin

Products Affected

- DIFFERIN EXTERNAL GEL 0.3 %

PA Criteria	Criteria Details
Covered Uses	acne vulgaris
Exclusion Criteria	
Required Medical Information	Documented diagnosis of acne vulgaris for members greater than 35 years old
Age Restrictions	Greater than 35. Members under age 35 can obtain medication without authorization.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dificid

Products Affected

- DIFICID

QL Criteria	20 tab Per 30 Days
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dihydroergotamine Mesylate

Products Affected

- *dihydroergotamine mesylate nasal*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of generic Migranal and two of the following: naratriptan, rizatriptan, sumatriptan, zolmitriptan
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dilaudid

Products Affected

- DILAUDID ORAL LIQUID

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dilaudid

Products Affected

- DILAUDID ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diovan

Products Affected

- DIOVAN ORAL TABLET 160 MG, 40 MG, 80 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diovan HCT

Products Affected

- DIOVAN HCT

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diovan HCT

Products Affected

- DIOVAN HCT

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dipentum

Products Affected

- DIPENTUM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diprolene

Products Affected

- DIPROLENE EXTERNAL LOTION

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diprolene

Products Affected

- DIPROLENE EXTERNAL OINTMENT

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ditropan XL

Products Affected

- DITROPAN XL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dolophine

Products Affected

- DOLOPHINE ORAL TABLET 5 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	180 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Doxepin HCl

Products Affected

- *doxepin hcl external*

QL Criteria	45 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Doxercalciferol

Products Affected

- *doxercalciferol oral*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Doxycycline

Products Affected

- *doxycycline*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Doxycycline Monohydrate

Products Affected

- *doxycycline monohydrate oral capsule 75 mg*

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dronabinol

Products Affected

- dronabinol*

PA Criteria	Criteria Details
Covered Uses	Anorexia associated with weight loss in patients with AIDS, Chemotherapy-induced nausea and vomiting
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Anorexia associated with weight loss in patients with AIDS, or Chemotherapy-induced nausea and vomiting
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: July 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duavee

Products Affected

- DUAVEE

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duexis

Products Affected

- DUEXIS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred generic non steroidal anti-inflammatory agent
QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dulera

Products Affected

- DULERA

QL Criteria	1 inhaler Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

DULoxetine HCl

Products Affected

- *duloxetine hcl oral capsule delayed release particles 20 mg*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

DULoxetine HCl

Products Affected

- *duloxetine hcl oral capsule delayed release particles 30 mg*

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

DULoxetine HCl

Products Affected

- *duloxetine hcl oral capsule delayed release particles 40 mg*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

DULoxetine HCl

Products Affected

- *duloxetine hcl oral capsule delayed release particles 60 mg*

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duo-Care Test

Products Affected

- DUO-CARE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duopa

Products Affected

- DUOPA ENTERAL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/antiparkinsons.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/antiparkinsons.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dupixent

Products Affected

- DUPIXENT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Dupixent.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 injections Per 1 month
Notes/References	
Revision Date	Prior Authorization: May 09, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duragesic-100

Products Affected

- DURAGESIC-100

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duragesic-12

Products Affected

- DURAGESIC-12

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duragesic-25

Products Affected

- DURAGESIC-25

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duragesic-50

Products Affected

- DURAGESIC-50

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duragesic-75

Products Affected

- DURAGESIC-75

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Durolane

Products Affected

- DUROLANE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dutasteride

Products Affected

- *dutasteride*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dutoprol

Products Affected

- DUTOPROL ORAL TABLET EXTENDED
RELEASE 24 HOUR 100-12.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of metoprolol-hydrochlorothiazide tablet or metoprolol succinate er tablet and hydrochlorothiazide tablet
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dutoprol

Products Affected

- DUTOPROL ORAL TABLET EXTENDED
RELEASE 24 HOUR 25-12.5 MG, 50-12.5
MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of metoprolol-hydrochlorothiazide tablet or metoprolol succinate er tablet and hydrochlorothiazide tablet
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Duzallo

Products Affected

- DUZALLO

PA Criteria	Criteria Details
Covered Uses	Treatment of hyperuricemia associated with gout in patients who have not achieved target serum uric acid levels with a medically appropriate daily dose of allopurinol alone.
Exclusion Criteria	For the treatment of asymptomatic hyperuricemia, severe renal impairment, end stage renal disease, kidney transplant recipients, or patients on dialysis, tumor lysis syndrome or Lesch-Nyhan syndrome, or for anyone with a known hypersensitivity to allopurinol, including previous occurrence of skin rash.
Required Medical Information	A documented diagnosis of hyperuricemia associated with gout and the member has a documented trial of allopurinol and has not achieved target serum uric acid levels.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of allopurinol or febuxostat
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: October 03, 2017 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

Dyanavel XR

Products Affected

- DYANA VEL XR

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	240 ml Per 30 Days
Notes/References	
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dysport

Products Affected

- DYSPOORT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/botulinum_toxin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Plus Blood Glucose Test

Products Affected

- *easy plus blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Plus II Glucose Test

Products Affected

- *easy plus ii glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Step Test

Products Affected

- EASY STEP TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Talk Blood Glucose Test

Products Affected

- *easy talk blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Touch Test

Products Affected

- EASY TOUCH TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Easy Trak Blood Glucose Test

Products Affected

- *easy trak blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EasyGluco

Products Affected

- EASYGLUCO IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EasyMax 15 Test

Products Affected

- EASYMAX 15 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EASYMax Test

Products Affected

- EASYMAX TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EasyPlus Blood Glucose Test

Products Affected

- *easyplus blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EasyPRO Blood Glucose Test

Products Affected

- EASYPRO BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EasyPRO Plus

Products Affected

- EASYPRO PLUS IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Econazole Nitrate

Products Affected

- *econazole nitrate external*

QL Criteria	85 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Edarbi

Products Affected

- EDARBI

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Edarbyclor

Products Affected

- EDARBYCLOR

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Edluar

Products Affected

- EDLUAR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 7 days (one week) of the preferred generic alternative zolpidem or zolpidem er.
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Effexor XR

Products Affected

- EFFEXOR XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 150 MG

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Effexor XR

Products Affected

- EFFEXOR XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 37.5 MG
- EFFEXOR XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 75 MG

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Effient

Products Affected

- EFFIENT

PA Criteria	Criteria Details
Covered Uses	Acute coronary syndrome (ACS) managed with percutaneous coronary intervention which includes unstable angina or non-ST elevation myocardial infarction or ST elevation myocardial infarction (MI)
Exclusion Criteria	History of Stroke or transient ischemic attack (TIA)
Required Medical Information	Member has a documented diagnosis of acute coronary syndrome (ACS) and is managed by percutaneous coronary intervention (PCI), which includes unstable angina, non-ST-elevation myocardial infarction (NSTEMI), or ST -elevation myocardial infarction (STEMI) managed with primary or delayed PCI and member has no prior history of stroke or transient ischemic attack (TIA)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tab Per 1 Day
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: May 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Efudex

Products Affected

- EFUDEX EXTERNAL CREAM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic fluorouracil 5% cream
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Elaprase

Products Affected

- ELAPRASE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Elelyso

Products Affected

- ELELYSO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: ?http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gaucher_disease.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Element Compact Test

Products Affected

- *element compact test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Element Test

Products Affected

- ELEMENT TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Elestrin

Products Affected

- ELESTRIN

QL Criteria	52 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Eletriptan Hydrobromide

Products Affected

- *eletriptan hydrobromide*

QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Elidel

Products Affected

- ELIDEL

PA Criteria	Criteria Details
Covered Uses	Atopic dermatitis
Exclusion Criteria	
Required Medical Information	FOR MEMBERS LESS THAN 2 YEARS OF AGE: Covered for the treatment of mild to moderate atopic dermatitis (eczema) for short-term use (up to 3 months). FOR MEMBERS OVER 2 YEARS OF AGE: A documented diagnosis of atopic dermatitis (eczema) and has a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for their condition, or they are being treated for atopic dermatitis (eczema) in an area at high risk for skin atrophy such as face, eyelids, or genital areas.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Less than 2 years of age: 3 months. Over 2 years of age: 1 year.
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patients condition
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Eligard

Products Affected

- ELIGARD

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Elmiron

Products Affected

- ELMIRON

QL Criteria	90 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Eloctate

Products Affected

- ELOCTATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Embeda

Products Affected

- EMBEDA

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Embrace Blood Glucose Test

Products Affected

- EMBRACE BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Embrace Evo Blood Glucose Test

Products Affected

- EMBRACE EVO BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Embrace Pro Glucose Test

Products Affected

- EMBRACE PRO GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emend

Products Affected

- EMEND ORAL CAPSULE 125 MG
- EMEND ORAL CAPSULE 40 MG

QL Criteria	5 caps Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emend

Products Affected

- EMEND ORAL CAPSULE 80 MG

QL Criteria	5 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emflaza

Products Affected

- EMFLAZA ORAL SUSPENSION

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Emflaza.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: April 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emflaza

Products Affected

- EMFLAZA ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Emflaza.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: April 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emsam

Products Affected

- EMSAM

QL Criteria	1 patch Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Emverm

Products Affected

- EMVERM

QL Criteria	6 tablets Per 3 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enablex

Products Affected

- ENABLEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enbrel

Products Affected

- ENBREL SUBCUTANEOUS SOLUTION
PREFILLED SYRINGE 25 MG/0.5ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
QL Criteria	8 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enbrel

Products Affected

- ENBREL SUBCUTANEOUS SOLUTION
PREFILLED SYRINGE 50 MG/ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
QL Criteria	4 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enbrel Mini

Products Affected

- ENBREL MINI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
QL Criteria	8 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enbrel SureClick

Products Affected

- ENBREL SURECLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Enbrel.html
QL Criteria	4 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Endocet

Products Affected

- ENDOCET ORAL TABLET 10-325 MG, 5-325 MG
- ENDOCET ORAL TABLET 2.5-325 MG, 7.5-325 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Enstilar

Products Affected

- ENSTILAR

QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Entocort EC

Products Affected

- ENTOCORT EC ORAL CAPSULE
DELAYED RELEASE PARTICLES

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic budesonide SR
QL Criteria	3 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Entresto

Products Affected

- ENTRESTO

PA Criteria	Criteria Details
Covered Uses	Chronic heart failure (NYHA Class II-IV) and reduced ejection fraction
Exclusion Criteria	
Required Medical Information	A documented diagnosis of chronic heart failure (NYHA Class II-IV) and reduced ejection fraction
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	Annual Review: 08/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Entyvio

Products Affected

- ENTYVIO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Entyvio.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Entyvio.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Epaned

Products Affected

- EPANED ORAL SOLUTION

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented inability to swallow
Age Restrictions	13 years or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Covered without prior authorization for those 12 years of age or younger
QL Criteria	1 bottle Per 30 Days
Notes/References	Annual Review: 08/2017
Revision Date	Prior Authorization: June 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Epclusa

Products Affected

- EPCLUSA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EPINEPHrine

Products Affected

- *epinephrine injection solution auto-injector*

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EpiPen 2-Pak

Products Affected

- EPIPEN 2-PAK INJECTION SOLUTION
AUTO-INJECTOR

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EpiPen Jr 2-Pak

Products Affected

- EPIPEN JR 2-PAK INJECTION
SOLUTION AUTO-INJECTOR

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EPIsnap

Products Affected

- EPISNAP

QL Criteria	4 injections Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Epogen

Products Affected

- EPOGEN INJECTION SOLUTION 10000 UNIT/ML, 2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML, 4000 UNIT/ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/Erythroipoiesis_Stimulating_Agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Epoprostenol Sodium

Products Affected

- *epoprostenol sodium*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Eprosartan Mesylate

Products Affected

- *eprosartan mesylate*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Erivedge

Products Affected

- ERIVEDGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ertaczo

Products Affected

- ERTACZO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Esbriet

Products Affected

- ESBRIET ORAL CAPSULE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/Idiopathic_Pulmonary_Fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	9 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Esbriet

Products Affected

- ESBRIET ORAL TABLET 267 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/Idiopathic_Pulmonary_Fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	9 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Esbriet

Products Affected

- ESBRIET ORAL TABLET 801 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Idiopathic_Pulmonary_Fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Escitalopram Oxalate

Products Affected

- *escitalopram oxalate oral solution*

QL Criteria	20 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Escitalopram Oxalate

Products Affected

- *escitalopram oxalate oral tablet*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Esomeprazole Magnesium

Products Affected

- *esomeprazole magnesium oral capsule
delayed release 40 mg*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Esomeprazole Strontium

Products Affected

- *esomeprazole strontium oral capsule delayed release 49.3 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 caps Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Estradiol

Products Affected

- *estradiol transdermal patch twice weekly*

QL Criteria	8 patches Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Estradiol

Products Affected

- *estradiol transdermal patch weekly*

QL Criteria	4 patches Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Estrogel

Products Affected

- ESTROGEL

QL Criteria	1 pump Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Eszopiclone

Products Affected

- *eszopiclone*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Eucrisa

Products Affected

- EUCRISA

PA Criteria	Criteria Details
Covered Uses	Mild to moderate atopic dermatitis
Exclusion Criteria	Not covered for members under 2 years old
Required Medical Information	A documented diagnosis of mild to moderate atopic dermatitis
Age Restrictions	Not covered for members under 2 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patients condition
Notes/References	
Revision Date	Prior Authorization: January 30, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Euflexxa

Products Affected

- EUFLEXXA INTRA-ARTICULAR SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Evamist

Products Affected

- EVAMIST

QL Criteria	2 bottles Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Evekeo

Products Affected

- EVEKEO

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD), Narcolepsy
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD) OR Narcolepsy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	120 tablets Per 30 Days
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: January 25, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EvenCare + Blood Glucose Test

Products Affected

- EVENCARE + BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EvenCare Blood Glucose Test

Products Affected

- EVENCARE BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EvenCare G2 Test

Products Affected

- EVENCARE G2 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

EvenCare G3 Test

Products Affected

- EVENCARE G3 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Evolution Autocode

Products Affected

- EVOLUTION AUTOCODE IN VITRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Evzio

Products Affected

- EVZIO INJECTION SOLUTION AUTO-INJECTOR 0.4 MG/0.4ML

PA Criteria	Criteria Details
Covered Uses	Emergency treatment of known or suspected opioid overdose
Exclusion Criteria	
Required Medical Information	Medication is being used for emergency treatment of opioid overdose
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Narcan nasal spray
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exalgo

Products Affected

- EXALGO ORAL TABLET ER 24 HOUR
ABUSE-DETERRENT 12 MG, 8 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exalgo

Products Affected

- EXALGO ORAL TABLET ER 24 HOUR
ABUSE-DETERRENT 16 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exalgo

Products Affected

- EXALGO ORAL TABLET ER 24 HOUR
ABUSE-DETERRENT 32 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exelderm

Products Affected

- EXELDERM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exelderm

Products Affected

- EXELDERM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 ml Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exforge

Products Affected

- EXFORGE

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exforge HCT

Products Affected

- EXFORGE HCT

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exjade

Products Affected

- EXJADE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Anitdotes.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Extavia

Products Affected

- EXTAVIA SUBCUTANEOUS KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Extina

Products Affected

- EXTINA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	50 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ez Smart Blood Glucose Test

Products Affected

- EZ SMART BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ez Smart Plus Glucose Test

Products Affected

- EZ SMART PLUS GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ezetimibe

Products Affected

- *ezetimibe*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ezetimibe-Simvastatin

Products Affected

- *ezetimibe-simvastatin*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fabrazyme

Products Affected

- FABRAZYME

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Famciclovir

Products Affected

- *famciclovir oral tablet 125 mg, 250 mg*

QL Criteria	60 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Famciclovir

Products Affected

- *famciclovir oral tablet 500 mg*

QL Criteria	21 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Famvir

Products Affected

- FAMVIR ORAL TABLET 500 MG

QL Criteria	21 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fanapt

Products Affected

- FANAPT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fanapt Titration Pack

Products Affected

- FANAPT TITRATION PACK

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	8 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Farxiga

Products Affected

- FARXIGA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Farydak

Products Affected

- FARYDAK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	12 capsules Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Faslodex

Products Affected

- FASLODEX INTRAMUSCULAR SOLUTION 250 MG/5ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FazaClo

Products Affected

- FAZACLO ORAL TABLET DISPERSIBLE
100 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	9 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FazaClo

Products Affected

- FAZACLO ORAL TABLET DISPERSIBLE
12.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FazaClo

Products Affected

- FAZACLO ORAL TABLET DISPERSIBLE
150 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FazaClo

Products Affected

- FAZACLO ORAL TABLET DISPERSIBLE
200 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FazaClo

Products Affected

- FAZACLO ORAL TABLET DISPERSIBLE
25 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Feiba

Products Affected

- FEIBA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Femhrt Low Dose

Products Affected

- FEMHRT LOW DOSE

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fenofibrate

Products Affected

- *fenofibrate oral*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fenofibrate

Products Affected

- *fenofibrate oral*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fenofibrate Micronized

Products Affected

- *fenofibrate micronized*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fenoglide

Products Affected

- FENOGLIDE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of three preferred fenofibrates (Trilipix, Fibracor, Lipofen, Tricor, Lofibra, Triglide/Lofibra, Antara)
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FentaNYL

Products Affected

- *fentanyl*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

QL Criteria	20 patch Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FentaNYL

Products Affected

- *fentanyl*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	20 patches Per 30 Days
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FentaNYL Citrate

Products Affected

- *fentanyl citrate buccal*

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 lozenges Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fentora

Products Affected

- FENTORA BUCCAL TABLET 100 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of the preferred generic alternative, fentanyl transmucosal lozenge, and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 tablets Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ferriprox

Products Affected

- FERRIPROX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Anitdotes.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fetzima

Products Affected

- FETZIMA

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 capsule Per 1 Day
Notes/References	Annual Review: 05/2017

Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Fetzima Titration

Products Affected

- FETZIMA TITRATION

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 titration pack Per 28 dayss
Notes/References	Annual Review: 05/2017

Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Fiasp

Products Affected

- FIASP

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fiasp FlexTouch

Products Affected

- FIASP FLEXTOUCH

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fifty50 Glucose Test 2.0

Products Affected

- FIFTY50 GLUCOSE TEST 2.0

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Finasteride

Products Affected

- *finasteride oral tablet 5 mg*

PA Criteria	Criteria Details
Covered Uses	Benign prostatic hyperplasia
Exclusion Criteria	
Required Medical Information	Member is greater than 50 years old or has diagnosis of BPH (Benign Prostatic Hyperplasia). For female members, must have a documented diagnosis of hirsutism secondary to ovarian or adrenal dysfunction (for example, polycystic ovary syndrome, adrenal or ovarian tumor)and must not be pregnant.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: October 25, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fioricet/Codeine

Products Affected

- FIORICET/CODEINE ORAL CAPSULE
50-300-40-30 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fiorinal/Codeine #3

Products Affected

- FIORINAL/CODEINE #3

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Firazyr

Products Affected

- FIRAZYR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Firmagon

Products Affected

- FIRMAGON

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Flebogamma DIF

Products Affected

- FLEBOGAMMA DIF

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Flector

Products Affected

- FLECTOR

QL Criteria	2 patch Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Flolan

Products Affected

- FLOLAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FloLipid

Products Affected

- *flolipid oral suspension 20 mg/5ml*

PA Criteria	Criteria Details
Covered Uses	Hyperlipidemia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Hyperlipidemia and documentation of an inability to swallow tablets/capsules
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of each of the following generic products: atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, and simvastatin
QL Criteria	5 milliliters Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 13, 2017 Step Therapy: November 06, 2017 Quantity Limits: November 15, 2017

FloLipid

Products Affected

- *flo lipid oral suspension 40 mg/5ml*

PA Criteria	Criteria Details
Covered Uses	Hyperlipidemia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Hyperlipidemia and documentation of an inability to swallow tablets/capsules
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of each of the following generic products: atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, and simvastatin
QL Criteria	10 milliliters Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 13, 2017 Step Therapy: November 06, 2017 Quantity Limits: August 25, 2015

Flovent Diskus

Products Affected

- FLOVENT DISKUS

QL Criteria	1 inhaler Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Flovent HFA

Products Affected

- FLOVENT HFA

QL Criteria	1 inhaler Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluocinonide

Products Affected

- *fluocinonide external*

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluoroplex

Products Affected

- FLUOROPLEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic fluorouracil 5% cream
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral capsule 10 mg*

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral capsule 20 mg*

QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral capsule 40 mg*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral capsule delayed release*

QL Criteria	1 caps Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral solution*

QL Criteria	10 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral tablet 10 mg, 60 mg*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral tablet 20 mg*

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluticasone-Salmeterol

Products Affected

- *fluticasone-salmeterol*

QL Criteria	1 inhaler Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluvastatin Sodium

Products Affected

- *fluvastatin sodium*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluvastatin Sodium ER

Products Affected

- *fluvastatin sodium er*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluvoxamine Maleate

Products Affected

- *fluvoxamine maleate oral tablet 100 mg*

QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluvoxamine Maleate

Products Affected

- *fluvoxamine maleate oral tablet 25 mg*
- *fluvoxamine maleate oral tablet 50 mg*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluvoxamine Maleate ER

Products Affected

- *fluvoxamine maleate er*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Focalin

Products Affected

- FOCALIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Focalin XR

Products Affected

- FOCALIN XR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Follistim AQ

Products Affected

- FOLLISTIM AQ INJECTION SOLUTION 75 UNT/0.5ML
- FOLLISTIM AQ SUBCUTANEOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA D15g Blood Glucose Test

Products Affected

- FORA D15G BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA D20 Blood Glucose Test

Products Affected

- FORA D20 BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA G20 Blood Glucose Test

Products Affected

- FORA G20 BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA G30a Blood Glucose Test

Products Affected

- FORA G30A BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fora GD20 Test

Products Affected

- FORA GD20 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA V10 Blood Glucose Test

Products Affected

- FORA V10 BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA V12 Blood Glucose Test

Products Affected

- FORA V12 BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA V20 Blood Glucose Test

Products Affected

- FORA V20 BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FORA V30a Blood Glucose Test

Products Affected

- FORA V30A BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ForaCare GD40 Test

Products Affected

- FORACARE GD40 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ForaCare premium V10 Test

Products Affected

- FORACARE PREMIUM V10 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ForaCare Test N Go Test

Products Affected

- FORACARE TEST N GO TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Forfivo XL

Products Affected

- FORFIVO XL

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fortamet

Products Affected

- FORTAMET ORAL TABLET EXTENDED
RELEASE 24 HOUR 1000 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Glucophage/Glucophage XR
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fortamet

Products Affected

- FORTAMET ORAL TABLET EXTENDED
RELEASE 24 HOUR 500 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Glucophage/Glucophage XR
QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Forteo

Products Affected

- FORTEO SUBCUTANEOUS SOLUTION
600 MCG/2.4ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fortesta

Products Affected

- FORTESTA

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	4 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fosamax

Products Affected

- FOSAMAX ORAL TABLET 70 MG

QL Criteria	1 tab Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fosamax Plus D

Products Affected

- FOSAMAX PLUS D

QL Criteria	1 tab Per 7 Days
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FreeStyle InsuLinx Test

Products Affected

- FREESTYLE INSULINX TEST

QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FreeStyle Lite Test

Products Affected

- FREESTYLE LITE TEST

QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

FreeStyle Test

Products Affected

- FREESTYLE TEST

QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Frova

Products Affected

- FROVA

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Frovatriptan Succinate

Products Affected

- *frovatriptan succinate*

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fuzeon

Products Affected

- FUZEON SUBCUTANEOUS SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviral_hiv.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fycompa

Products Affected

- FYCOMPA ORAL TABLET

QL Criteria	1 tab Per 1 Day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabapentin

Products Affected

- *gabapentin oral capsule*

QL Criteria	6 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabapentin

Products Affected

- *gabapentin oral solution 250 mg/5ml*

QL Criteria	40 ML Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabapentin

Products Affected

- *gabapentin oral tablet*

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabril

Products Affected

- GABITRIL ORAL TABLET 12 MG
- GABITRIL ORAL TABLET 4 MG

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabril

Products Affected

- GABITRIL ORAL TABLET 16 MG

QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gabitril

Products Affected

- GABITRIL ORAL TABLET 2 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gammagard

Products Affected

- GAMMAGARD

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gammagard S/D Less IgA

Products Affected

- GAMMAGARD S/D LESS IGA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gammaked

Products Affected

- GAMMAKED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gammaplex

Products Affected

- GAMMAPLEX INTRAVENOUS SOLUTION 10 GM/200ML, 20 GM/400ML, 5 GM/100ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gamunex-C

Products Affected

- GAMUNEX-C

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ganirelix Acetate

Products Affected

- *ganirelix acetate*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/inferility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gattex

Products Affected

- GATTEX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gattex.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

GE100 Blood Glucose Test

Products Affected

- *ge100 blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gelnique

Products Affected

- GELNIQUE TRANSDERMAL GEL 10 %

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gel-One

Products Affected

- GEL-ONE INTRA-ARTICULAR
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Gelsyn-3

Products Affected

- GELSYN-3

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Genotropin

Products Affected

- GENOTROPIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Genotropin MiniQuick

Products Affected

- GENOTROPIN MINIQUICK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

GenStrip 50

Products Affected

- GENSTRIP 50

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Genvoya

Products Affected

- GENVOYA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviral_hiv.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Geodon

Products Affected

- GEODON ORAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Giazo

Products Affected

- GIAZO

PA Criteria	Criteria Details
Covered Uses	Mild to moderate ulcerative colitis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of mild to moderate ulcerative colitis in male patients 18 years and older
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Per Product Labeling, Giazo effectiveness was not demonstrated in female patients
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Apriso, Delzicol, Lialda, or Pentasa
QL Criteria	6 tab Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gilenya

Products Affected

- GILENYA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gilotrif

Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glassia

Products Affected

- GLASSIA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Alpha-1 Antitrypsin Inhibitor Therapy.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glatopa

Products Affected

- GLATOPA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gleevec

Products Affected

- GLEEVEC ORAL TABLET 100 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gleevec

Products Affected

- GLEEVEC ORAL TABLET 400 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gluco Perfect 3 Test

Products Affected

- GLUCO PERFECT 3 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glucocard 01 Sensor Plus

Products Affected

- GLUCOCARD 01 SENSOR PLUS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glucocard Expression Test

Products Affected

- GLUCOCARD EXPRESSION TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glucocard Shine Test

Products Affected

- GLUCOCARD SHINE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glucocard Vital Test

Products Affected

- GLUCOCARD VITAL TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glucocard X-Sensor

Products Affected

- GLUCOCARD X-SENSOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

GlucoCom Test

Products Affected

- GLUCOCOM TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

GlucoNavii Blood Glucose Test

Products Affected

- GLUCONAVII BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glumetza

Products Affected

- GLUMETZA ORAL TABLET EXTENDED
RELEASE 24 HOUR 1000 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Glucophage/Glucophage XR
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glumetza

Products Affected

- GLUMETZA ORAL TABLET EXTENDED
RELEASE 24 HOUR 500 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Glucophage/Glucophage XR
QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Glyxambi

Products Affected

- GLYXAMBI

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gocovri

Products Affected

- GOCOVRI

PA Criteria	Criteria Details
Covered Uses	Treatment of dyskinesia in patients with Parkinsons disease receiving levodopa based therapy, with or without concomitant dopaminergic medications
Exclusion Criteria	Contraindicated in patients with end stage renal disease
Required Medical Information	A documented diagnosis of dyskinesia associated with Parkinsons disease and member is currently receiving levodopa based therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of amantadine
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: November 06, 2017 Quantity Limits: November 15, 2017

Gonal-f

Products Affected

- GONAL-F

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/inferility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gonal-f RFF

Products Affected

- GONAL-F RFF

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infer-tility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gonal-f RFF Rediject

Products Affected

- GONAL-F RFF REDIJECT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/inferility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gralise

Products Affected

- GRALISE ORAL TABLET 300 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of gabapentin
QL Criteria	5 tab Per 1 Day
Notes/ References	Annual Review: 02/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gralise

Products Affected

- GRALISE ORAL TABLET 600 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of gabapentin
QL Criteria	3 tab Per 1 Day
Notes/ References	Annual Review: 02/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Gralise Starter

Products Affected

- GRALISE STARTER

ST Criteria	A documented contraindication, intolerance, allergy, or failure of gabapentin
QL Criteria	1 pack Per 365 Days
Notes/ References	Annual Review: 02/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Granix

Products Affected

- GRANIX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/G-CSF.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

GuanFACINE HCl ER

Products Affected

- *guanfacine hcl er*

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Haegarda

Products Affected

- HAEGARDA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
QL Criteria	16 kits Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Halobetasol Propionate

Products Affected

- *halobetasol propionate*

QL Criteria	50 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Harvoni

Products Affected

- HARVONI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hectorol

Products Affected

- HECTOROL ORAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of doxercalciferol and calcitriol
QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Helixate FS

Products Affected

- HELIXATE FS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hemangeol

Products Affected

- HEMANGEOL

PA Criteria	Criteria Details
Covered Uses	Proliferating infantile hemangioma
Exclusion Criteria	History of asthma or bronchospasms
Required Medical Information	A documented diagnosis of proliferating infantile hemangioma requiring systemic therapy and documented all of the following: (1) Member was not born prematurely with a corrected age of less than 5 weeks, (2) Member does not weigh less than 2kg, have sustained heart rate less than 80 beats per minute, have greater than first degree heart block, or have decompensated heart failure, and (3) Member does not have sustained blood pressure less than 50/ 30mmHg.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: July 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hemofil M

Products Affected

- HEMOFIL M INTRAVENOUS SOLUTION
RECONSTITUTED 1000 UNIT, 1700
UNIT, 250 UNIT, 500 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hetlioz

Products Affected

- HETLIOZ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/sedative-hypnotics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hizentra

Products Affected

- HIZENTRA SUBCUTANEOUS SOLUTION 1 GM/5ML, 10 GM/50ML, 2 GM/10ML, 4 GM/20ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Horizant

Products Affected

- HORIZANT ORAL TABLET EXTENDED RELEASE

PA Criteria	Criteria Details
Covered Uses	Post-herpetic neuralgia and Restless leg syndrome
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Restless Leg Syndrome (RLS) or Post Herpetic Neuralgia (shingles)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	FOR POST-HERPETIC NEURALGIA: A documented contraindication, intolerance, allergy, or failure of two weeks of gabapentin. FOR RESTLESS LEG SYNDROME: A documented contraindication, intolerance, allergy, or failure of two weeks of pramipexole or ropinirole.
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: February 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

HP Acthar

Products Affected

- HP ACTHAR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/acthar.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humate-P

Products Affected

- HUMATE-P INTRAVENOUS SOLUTION
RECONSTITUTED 1000-2400 UNIT, 250-
600 UNIT, 500-1200 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humatrope

Products Affected

- HUMATROPE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira

Products Affected

- HUMIRA SUBCUTANEOUS PREFILLED SYRINGE KIT 10 MG/0.2ML, 20 MG/0.4ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	2 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira

Products Affected

- HUMIRA SUBCUTANEOUS PREFILLED SYRINGE KIT 40 MG/0.8ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	6 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira Pediatric Crohns Start

Products Affected

- HUMIRA PEDIATRIC CROHNS START
SUBCUTANEOUS PREFILLED SYRINGE
KIT 40 MG/0.8ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	6 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira Pen

Products Affected

- HUMIRA PEN SUBCUTANEOUS PEN-INJECTOR KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	6 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira Pen-Crohns Starter

Products Affected

- HUMIRA PEN-CROHNS STARTER
SUBCUTANEOUS PEN-INJECTOR KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	6 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Humira Pen-Psoriasis Starter

Products Affected

- HUMIRA PEN-PSORIASIS STARTER
SUBCUTANEOUS PEN-INJECTOR KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Humira.html
QL Criteria	6 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hyalgan

Products Affected

- HYALGAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hycamtin

Products Affected

- HYCAMTIN ORAL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hycet

Products Affected

- HYCET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hydrocodone-Acetaminophen

Products Affected

- *hydrocodone-acetaminophen oral solution*
2.5-108 mg/5ml, 5-217 mg/10ml, 7.5-325 mg/15ml

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hydrocodone-Acetaminophen

Products Affected

- *hydrocodone-acetaminophen oral tablet 10-300 mg, 10-325 mg, 2.5-325 mg, 5-300 mg, 5-325 mg, 7.5-300 mg, 7.5-325 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hydrocodone-Ibuprofen

Products Affected

- *hydrocodone-ibuprofen oral tablet 10-200 mg*
- *hydrocodone-ibuprofen oral tablet 5-200 mg, 7.5-200 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

HYDRomorphone HCl

Products Affected

- *hydromorphone hcl oral liquid*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

HYDRomorphone HCl

Products Affected

- *hydromorphone hcl oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

HYDRomorphone HCl ER

Products Affected

- *hydromorphone hcl er oral tablet er 24 hour abuse-deterrent 12 mg, 32 mg, 8 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

HYDRomorphone HCl ER

Products Affected

- *hydromorphone hcl er oral tablet er 24 hour abuse-deterrent 16 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hymovis

Products Affected

- HYMOVIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hyqvia

Products Affected

- HYQVIA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Hysingla ER

Products Affected

- HYSINGLA ER

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ibandronate Sodium

Products Affected

- *ibandronate sodium oral*

QL Criteria	1 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ibrance

Products Affected

- IBRANCE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	21 capsules Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ibudone

Products Affected

- IBUDONE

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Iclusig

Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Idelvion

Products Affected

- IDELVION

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

IDHIFA

Products Affected

- IDHIFA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/I/dhifa.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ilaris

Products Affected

- ILARIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Ilaris.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ilaris (150mg Delivered)

Products Affected

- ILARIS (150MG DELIVERED)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Ilaris.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imatinib Mesylate

Products Affected

- *imatinib mesylate oral tablet 100 mg*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imatinib Mesylate

Products Affected

- *imatinib mesylate oral tablet 400 mg*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imbruvica

Products Affected

- IMBRUVICA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imiquimod

Products Affected

- *imiquimod external*

QL Criteria	48 packets Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imitrex

Products Affected

- IMITREX NASAL

QL Criteria	6 sprays Per 30 days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imitrex

Products Affected

- IMITREX ORAL

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imitrex

Products Affected

- IMITREX SUBCUTANEOUS

QL Criteria	10 vials Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Impavido

Products Affected

- IMPAVIDO

PA Criteria	Criteria Details
Covered Uses	Leishmaniasis
Exclusion Criteria	Known or suspected pregnancy
Required Medical Information	A documented diagnosis of any of the following leishmaniasis infections: Visceral leishmaniasis due to <i>Leishmania donovani</i> , Cutaneous leishmaniasis due to <i>Leishmania braziliensis</i> , <i>Leishmania guyanensis</i> , and <i>Leishmania panamensis</i> , or Mucosal leishmaniasis due to <i>Leishmania braziliensis</i>
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	28 days
Other Criteria	
QL Criteria	84 capsules Per 28 Days
Notes/References	
Revision Date	Prior Authorization: August 16, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Increlex

Products Affected

- INCRELEX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Increlex.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Incruse Ellipta

Products Affected

- INCRUSE ELLIPTA

QL Criteria	1 blister Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Inderal LA

Products Affected

- INDERAL LA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic propranolol SR
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Indomethacin

Products Affected

- *indomethacin oral*

QL Criteria	3 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Infinity Blood Glucose Test

Products Affected

- INFINITY BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Inflectra

Products Affected

- INFLECTRA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Inflectra.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Inflectra.html
Notes/References	
Revision Date	Prior Authorization: December 13, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ingrezza

Products Affected

- INGREZZA ORAL CAPSULE 40 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/Ingrezza.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: June 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ingrezza

Products Affected

- INGREZZA ORAL CAPSULE 80 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/Ingrezza.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: June 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: November 16, 2017

Inlyta

Products Affected

- INLYTA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Intermezzo

Products Affected

- INTERMEZZO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 7 days (one week) of the preferred generic alternative zolpidem or zolpidem er.
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Intrarosa

Products Affected

- INTRAROSA

QL Criteria	1 insert Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Intron A

Products Affected

- INTRON A

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Introvale

Products Affected

- INTROVALE

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Intuniv

Products Affected

- INTUNIV

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: clonidine/sr, guanfacine, amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine, or Vyvanse
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Invega

Products Affected

- INVEGA ORAL TABLET EXTENDED
RELEASE 24 HOUR 1.5 MG, 3 MG, 6 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Invega

Products Affected

- INVEGA ORAL TABLET EXTENDED
RELEASE 24 HOUR 9 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Invokamet

Products Affected

- INVOKAMET

QL Criteria	1 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Invokamet XR

Products Affected

- INVOKAMET XR

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Invokana

Products Affected

- INVOKANA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Irbesartan

Products Affected

- *irbesartan*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Irbesartan

Products Affected

- *irbesartan*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Irbesartan-Hydrochlorothiazide

Products Affected

- *irbesartan-hydrochlorothiazide oral tablet*
150-12.5 mg

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Iressa

Products Affected

- IRESSA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Isentress HD

Products Affected

- ISENTRESS HD

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ixinity

Products Affected

- IXINITY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jadenu

Products Affected

- JADENU

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Anitdotes.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jadenu Sprinkle

Products Affected

- JADENU SPRINKLE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Anitdotes.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jakafi

Products Affected

- JAKAFI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Janumet

Products Affected

- JANUMET

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Janumet XR

Products Affected

- JANUMET XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 100-
1000 MG, 50-500 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Janumet XR

Products Affected

- JANUMET XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 50-1000
MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Januvia

Products Affected

- JANUVIA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jardiance

Products Affected

- JARDIANCE

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jentaduetto

Products Affected

- JENTADUETO

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 05/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jentaduetto XR

Products Affected

- JENTADUETO XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 2.5-1000
MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 05/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jentaduetto XR

Products Affected

- JENTADUETO XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 5-1000
MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jetrea

Products Affected

- JETREA INTRAVITREAL SOLUTION
0.375 MG/0.3ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/EYE/ophtalmic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jolessa

Products Affected

- JOLESSA

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Juxtapid

Products Affected

- JUXTAPID ORAL CAPSULE 10 MG, 5 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
QL Criteria	1 capsule Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Juxtapid

Products Affected

- JUXTAPID ORAL CAPSULE 20 MG, 30 MG, 40 MG, 60 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kadian

Products Affected

- KADIAN ORAL CAPSULE EXTENDED RELEASE 24 HOUR 10 MG, 100 MG, 20 MG, 30 MG, 50 MG, 60 MG, 80 MG
- KADIAN ORAL CAPSULE EXTENDED RELEASE 24 HOUR 200 MG, 40 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kalbitor

Products Affected

- KALBITOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kalydeco

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/cystic_fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 packets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kanuma

Products Affected

- KANUMA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kapvay

Products Affected

- KAPVAY ORAL TABLET EXTENDED RELEASE 12 HOUR

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tablets Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kazano

Products Affected

- KAZANO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentaducto; and generic alogliptin, alogliptin/pioglitazone, alogliptin/metformin
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kcentra

Products Affected

- KCENTRA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Keppra XR

Products Affected

- KEPPRA XR ORAL TABLET EXTENDED
RELEASE 24 HOUR 500 MG

QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Keppra XR

Products Affected

- KEPPRA XR ORAL TABLET EXTENDED
RELEASE 24 HOUR 750 MG

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ketoconazole

Products Affected

- *ketoconazole external foam*

QL Criteria	50 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ketorolac Tromethamine

Products Affected

- *ketorolac tromethamine oral*

QL Criteria	20 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Keveyis

Products Affected

- KEVEYIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/carb onic_anhydrase_inhibitor.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kevzara

Products Affected

- KEVZARA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Kevzara.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Kevzara.html
QL Criteria	2 injections Per 1 month
Notes/References	
Revision Date	Prior Authorization: June 23, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Khedezla

Products Affected

- KHEDEZLA

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 05/2017

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Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Kineret

Products Affected

- KINERET SUBCUTANEOUS SOLUTION
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Kineret.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Kineret.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali 200 Dose

Products Affected

- KISQALI 200 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali 400 Dose

Products Affected

- KISQALI 400 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali 600 Dose

Products Affected

- KISQALI 600 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali Femara 200 Dose

Products Affected

- KISQALI FEMARA 200 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali Femara 400 Dose

Products Affected

- KISQALI FEMARA 400 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kisqali Femara 600 Dose

Products Affected

- KISQALI FEMARA 600 DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Kisqali.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 box Per 1 month
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Koate

Products Affected

- KOATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Koate-DVI

Products Affected

- KOATE-DVI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kogenate FS

Products Affected

- KOGENATE FS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kogenate FS Bio-Set

Products Affected

- KOGENATE FS BIO-SET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kombiglyze XR

Products Affected

- KOMBIGLYZE XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 2.5-1000
MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kombiglyze XR

Products Affected

- KOMBIGLYZE XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 5-1000
MG, 5-500 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Korlym

Products Affected

- KORLYM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/anti-diabetic-agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kovaltry

Products Affected

- KOVALTRY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kroger Blood Glucose Test

Products Affected

- *kroger blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kroger Premium Glucose Test

Products Affected

- *kroger premium glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kroger Test

Products Affected

- *kroger test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Krystexxa

Products Affected

- KRYSTEXXA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gout.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gout.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kuvan

Products Affected

- KUVAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kynamro

Products Affected

- KYNAMRO SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipidemic_Agents_HOFH.html
QL Criteria	4 injections Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET
DISPERSIBLE 100 MG, 200 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET
DISPERSIBLE 25 MG

QL Criteria	6 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET
DISPERSIBLE 50 MG

QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 100 MG,
25 MG, 50 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 200 MG

QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 250 MG,
300 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIGine

Products Affected

- *lamotrigine oral tablet dispersible 100 mg, 200 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRigine

Products Affected

- *lamotrigine oral tablet dispersible 25 mg*

QL Criteria	6 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIGine

Products Affected

- *lamotrigine oral tablet dispersible 50 mg*

QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIGine ER

Products Affected

- *lamotrigine er oral tablet extended release*
24 hour 100 mg, 25 mg, 50 mg

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIGine ER

Products Affected

- *lamotrigine er oral tablet extended release*
24 hour 200 mg

QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIGine ER

Products Affected

- *lamotrigine er oral tablet extended release*
24 hour 250 mg, 300 mg

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lansoprazole

Products Affected

- *lansoprazole oral capsule delayed release 30 mg*

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Larin Fe 1/20

Products Affected

- LARIN FE 1/20

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Latuda

Products Affected

- LATUDA ORAL TABLET 120 MG, 20 MG, 40 MG, 60 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Latuda

Products Affected

- LATUDA ORAL TABLET 80 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lazanda

Products Affected

- LAZANDA

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of the preferred generic alternative, fentanyl transmucosal lozenge, and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	4 bottle Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lazanda

Products Affected

- LAZANDA

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of the preferred generic alternative, fentanyl transmucosal lozenge, and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	4 bottles Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Lemtrada

Products Affected

- LEMTRADA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
QL Criteria	6 vials Per 365 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 10 MG Daily Dose

Products Affected

- LENVIMA 10 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 14 MG Daily Dose

Products Affected

- LENVIMA 14 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 18 MG Daily Dose

Products Affected

- LENVIMA 18 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 20 MG Daily Dose

Products Affected

- LENVIMA 20 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 24 MG Daily Dose

Products Affected

- LENVIMA 24 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lenvima 8 MG Daily Dose

Products Affected

- LENVIMA 8 MG DAILY DOSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lescol

Products Affected

- LESCOL ORAL CAPSULE 20 MG

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lescol XL

Products Affected

- LESCOL XL

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Letairis

Products Affected

- LETAIRIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Leukine

Products Affected

- LEUKINE INTRAVENOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/G-CSF.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Leuprolide Acetate

Products Affected

- *leuprolide acetate injection*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LevETIRAcetam ER

Products Affected

- *levetiracetam er oral tablet extended release*
24 hour 500 mg

QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LevETIRAcetam ER

Products Affected

- *levetiracetam er oral tablet extended release*
24 hour 750 mg

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Levonorgest-Eth Estrad 91-Day

Products Affected

- *levonorgest-eth estrad 91-day oral tablet*
0.1-0.02 & 0.01 mg, 0.15-0.03 mg

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Levorphanol Tartrate

Products Affected

- *levorphanol tartrate oral*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Levulan Kerastick

Products Affected

- LEVULAN KERASTICK

QL Criteria	1 stick Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lexapro

Products Affected

- LEXAPRO ORAL TABLET

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lialda

Products Affected

- LIALDA

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Liberty Next Generation Test

Products Affected

- LIBERTY NEXT GENERATION TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Liberty Test

Products Affected

- *liberty test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidocaine

Products Affected

- *lidocaine external ointment*

PA Criteria	Criteria Details
Covered Uses	***AUTHORIZATION IS NOT REQUIRED FOR LESS THAN 50 GRAMS OF LIDOCAINE EVERY 30 DAYS*** For quantities over 50 grams every 30 days, there must be a documented temporary need for anesthesia for any of the following: Accessible mucous membranes of the oropharynx, skin and mucous membranes or stomatitis, or pain associated with a minor burns, including sunburn, abrasions of the skin, and insect bites.
Exclusion Criteria	Documentation of any of the following: Planned area of application includes non-intact skin, sensitivity to amide-type local anesthetics or any other component of the product, planned use on large surface area of the body as this can lead to increased toxicity, planned area of application includes severely traumatized skin (e.g., mucosal or skin abrasion, eczema, burns), the medication is being used in conjunction with a cosmetic procedure (i.e. hair removal), or if the product will be compounded with other products that would alter the total dose/dosage form being administered
Required Medical Information	A documented need for temporary anesthesia for any of the following: Accessible mucous membranes of the oropharynx, skin and mucous membranes or stomatitis, or pain associated with a minor burns, including sunburn, abrasions of the skin, and insect bites.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months

PA Criteria	Criteria Details
Other Criteria	<p>*Topical lidocaine ointment is used for temporary anesthesia. Prescription renewals for longer than 3 months require clinical documentation of medical necessity. Due to Safety Concerns higher quantities and prolonged use are not recommended. Renewal Duration: 3 months *Approval can made up to an additional 50gms per 30 days. Higher additional quantities are not approvable *FOR ADULTS: A single application should not exceed 5 g of Lidocaine Ointment 5%, containing 250 mg of lidocaine base (equivalent chemically to approximately 300 mg of lidocaine hydrochloride). This is roughly equivalent to squeezing a six (6) inch length of ointment from the tube. In a 70 kg adult this dose equals 3.6 mg/kg (1.6 mg/lb) lidocaine base. No more than one-half tube, approximately 17-20 g of ointment or 850-1000 mg lidocaine base, should be administered in any one day. FOR CHILDREN: For children less than ten years who have a normal lean body mass and a normal lean body development, the maximum dose may be determined by the application of one of the standard pediatric drug formulas (e.g., Clark's rule). For example a child of five years weighing 50 lbs., the dose of lidocaine should not exceed 75-100 mg when calculated according to Clark's rule. In any case, the maximum amount of lidocaine administered should not exceed 4.5 mg/kg (2.0 mg/lb) of body weight ***Lidocaine toxicity resulting from transcutaneous absorption is theoretically possible. Signs and symptoms of systemic lidocaine toxicity include CNS excitation and/or depression, nervousness, confusion, dizziness, tinnitus, blurred or double vision, vomiting, twitching, tremors, seizures, unconsciousness, respiratory depression, bradycardia, hypotension, and cardiopulmonary arrest. If there is suspicion of lidocaine-related systemic toxicity, check lidocaine blood concentrations</p>
QL Criteria	50 grams Per 30 Days
Notes/References	
Revision Date	Prior Authorization: October 03, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidocaine

Products Affected

- *lidocaine external patch 5 %*

PA Criteria	Criteria Details
Covered Uses	Neuropathic pain (i.e. post-herpetic neuralgia)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of neuropathic pain (i.e. post-herpetic neuralgia)
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	3 patches Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidocaine PAK

Products Affected

- *lidocaine pak*

QL Criteria	50 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidocaine-Prilocaine

Products Affected

- *lidocaine-prilocaine external cream*

PA Criteria	Criteria Details
Covered Uses	***AUTHORIZATION IS NOT REQUIRED FOR LESS THAN 50 GRAMS OF LIDOCAINE EVERY 30 DAYS*** For quantities over 50 grams every 30 days, there must be a documented temporary need for topical anesthetic in either of the following situations: Normal, intact skin for local analgesia, or Genital mucous membranes for superficial minor surgery and as pretreatment for infiltration anesthesia
Exclusion Criteria	Documentation of any of the following: Planned area of application includes non-intact skin, Sensitivity to amide-type local anesthetics or any other component of the product, Planned use on large surface area of the body or for a period of time over 3 hours as this can lead to increased toxicity, the medication is being used in conjunction with a cosmetic procedure (i.e. hair removal), Use in situations where the drug may migrate into the middle ear, beyond the tympanic membrane, History of methemoglobinemia, or if the product will be compounded with other products that would alter the total dose/dosage form being administered
Required Medical Information	A documented need for topical anesthetic in either of the following situations: Normal, intact skin for local analgesia, or Genital mucous membranes for superficial minor surgery and as pretreatment for infiltration anesthesia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months

PA Criteria	Criteria Details
Other Criteria	*Topical lidocaine/prilocaine cream is used for temporary anesthesia. Prescription renewals for longer than 3 months require clinical documentation of medical necessity. Due to Safety Concerns higher quantities and prolonged use are not recommended. Renewal Duration: 3 months *Up to an additional 30 grams per 30 days. Higher additional quantities are not approvable.
QL Criteria	30 grams Per 30 Days
Notes/References	
Revision Date	Prior Authorization: October 03, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidocaine-Tetracaine

Products Affected

- *lidocaine-tetracaine*

QL Criteria	30 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lidoderm

Products Affected

- LIDODERM

PA Criteria	Criteria Details
Covered Uses	Neuropathic pain (i.e. post-herpetic neuralgia)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of neuropathic pain (i.e. post-herpetic neuralgia)
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic gabapentin or Lyrica
QL Criteria	3 patches Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Linzess

Products Affected

- LINZESS

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lipitor

Products Affected

- LIPITOR

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lipofen

Products Affected

- LIPOFEN

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Livalo

Products Affected

- LIVALO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic statin medication (atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, or simvastatin)
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Locoid

Products Affected

- LOCOID

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred generic alternative from the following medications: betamethasone benzoate, betamethasone dipropionate, betamethasone valerate, desonide, desoximetasone, fluocinolone acetonide, fluticasone, fluocinonide, hydrocortisone butyrate, hydrocortisone valerate, prednicarbate, or triamcinolone acetonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Locoid Lipocream

Products Affected

- LOCOID LIPOCREAM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred generic alternative from the following medications: betamethasone benzoate, betamethasone dipropionate, betamethasone valerate, desonide, desoximetasone, fluocinolone acetonide, fluticasone, fluocinonide, hydrocortisone butyrate, hydrocortisone valerate, prednicarbate, or triamcinolone acetonide
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lofibra

Products Affected

- LOFIBRA ORAL CAPSULE 134 MG, 67 MG

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lofibra

Products Affected

- LOFIBRA ORAL TABLET 54 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lonsurf

Products Affected

- LONSURF ORAL TABLET 15-6.14 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	100 tablets Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lonsurf

Products Affected

- LONSURF ORAL TABLET 20-8.19 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	80 tablets Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lorcet

Products Affected

- LORCET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lorcet HD

Products Affected

- LORCET HD

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lorcet Plus

Products Affected

- LORCET PLUS ORAL TABLET 7.5-325 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lortab

Products Affected

- LORTAB ORAL ELIXIR 10-300 MG/15ML

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Losartan Potassium

Products Affected

- *losartan potassium oral tablet 25 mg, 50 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LoSeasonique

Products Affected

- LOSEASONIQUE

QL Criteria	90 days maximum Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lotronex

Products Affected

- LOTRONEX

PA Criteria	Criteria Details
Covered Uses	Irritable bowel syndrome
Exclusion Criteria	
Required Medical Information	(1)A female patient with a diagnosis of severe* irritable bowel syndrome (IBS) with primary symptom of diarrhea with chronic IBS symptoms (generally lasting 6 months or longer), and (2) anatomic or biochemical abnormalities of the gastrointestinal tract have been excluded, and (3) failure to respond to diphenoxylate/atropine and loperamide for at least one month each.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	*Diarrhea-predominant IBS is severe if it includes diarrhea and one or more of the following: (1) frequent and severe abdominal pain/discomfort, or (2) frequent urgency or fecal incontinence, or (3) disability or restriction of daily activities due to IBS.
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each diphenoxylate/atropine and loperamide
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lovastatin

Products Affected

- *lovastatin*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lovaza

Products Affected

- LOVAZA

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lucentis

Products Affected

- LUCENTIS INTRAVITREAL SOLUTION
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/EYE/ophtalmic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lumizyme

Products Affected

- LUMIZYME

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lunesta

Products Affected

- LUNESTA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupaneta Pack

Products Affected

- LUPANETA PACK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot (1-Month)

Products Affected

- LUPRON DEPOT (1-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot (3-Month)

Products Affected

- LUPRON DEPOT (3-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot (4-Month)

Products Affected

- LUPRON DEPOT (4-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot (6-Month)

Products Affected

- LUPRON DEPOT (6-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot-Ped (1-Month)

Products Affected

- LUPRON DEPOT-PED (1-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lupron Depot-Ped (3-Month)

Products Affected

- LUPRON DEPOT-PED (3-MONTH)

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Luxiq

Products Affected

- LUXIQ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of a preferred generic betamethasone alternative
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Luzu

Products Affected

- LUZU

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lynparza

Products Affected

- LYNPARZA ORAL CAPSULE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lynparza

Products Affected

- LYNPARZA ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lysteda

Products Affected

- LYSTEDA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic tranexamic acid
QL Criteria	30 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Macugen

Products Affected

- MACUGEN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/EYE/opthalmic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Makena

Products Affected

- MAKENA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/hydroxyprogesterone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	5 vials Per 365 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Malarone

Products Affected

- MALARONE

PA Criteria	Criteria Details
Covered Uses	Malaria
Exclusion Criteria	
Required Medical Information	A documented diagnosis of malaria
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Maximum Approval for Malaria: 30 days, Maximum Approval for all other indications: 1 year
Other Criteria	Malaria medications are Certificate of Coverage (COC) Excluded for travel prophylaxis
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Marinol

Products Affected

- MARINOL

PA Criteria	Criteria Details
Covered Uses	Anorexia associated with weight loss in patients with AIDS, Chemotherapy-induced nausea and vomiting
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Anorexia associated with weight loss in patients with AIDS, or Chemotherapy-induced nausea and vomiting
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: July 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mavyret

Products Affected

- MAVYRET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Maxalt

Products Affected

- MAXALT

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Maxalt-MLT

Products Affected

- MAXALT-MLT

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mefloquine HCl

Products Affected

- *mefloquine hcl*

PA Criteria	Criteria Details
Covered Uses	Malaria
Exclusion Criteria	
Required Medical Information	A documented diagnosis of malaria
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Maximum Approval for Malaria: 30 days, Maximum Approval for all other indications: 1 year
Other Criteria	Malaria medications are Certificate of Coverage (COC) Excluded for travel prophylaxis
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Meijer Blood Glucose Test

Products Affected

- *meijer blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Meijer Premium Glucose Test

Products Affected

- *meijer premium glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Meijer TRUEtest Test

Products Affected

- MEIJER TRUETEST TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Meijer TRUEtrack Test

Products Affected

- MEIJER TRUETRACK TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mekinist

Products Affected

- MEKINIST

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Menopur

Products Affected

- MENOPUR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infer-tility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Menostar

Products Affected

- MENOSTAR

QL Criteria	1 patch Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Meperidine HCl

Products Affected

- *meperidine hcl oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mephyton

Products Affected

- MEPHYTON

QL Criteria	25 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mesalamine

Products Affected

- *mesalamine oral tablet delayed release 1.2 gm*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mesalamine

Products Affected

- *mesalamine oral tablet delayed release 800 mg*

QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metadate ER

Products Affected

- METADATE ER ORAL TABLET
EXTENDED RELEASE 20 MG

QL Criteria	3 tab Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MetFORMIN HCl ER (MOD)

Products Affected

- *metformin hcl er (mod) oral tablet extended release 24 hour 1000 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MetFORMIN HCl ER (MOD)

Products Affected

- *metformin hcl er (mod) oral tablet extended release 24 hour 500 mg*

QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MetFORMIN HCl ER (OSM)

Products Affected

- *metformin hcl er (osm) oral tablet extended release 24 hour 1000 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MetFORMIN HCl ER (OSM)

Products Affected

- *metformin hcl er (osm) oral tablet extended release 24 hour 500 mg*

QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadone HCl

Products Affected

- *methadone hcl oral concentrate*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. (4) FOR A DOCUMENTED DIAGNOSIS OF OPIOID ADDICTION (heroin or other morphine-like drugs): medication must be dispensed by a treatment program certified by SAMHSA (Substance Abuse and Mental Health Services Administration) and the patient will be monitored during therapy for signs and symptoms of abuse/misuse as well as compliance and the potential diversion to others. (Initiation/detoxification treatment = 1 month approval, continuation of therapy/maintenance treatment = 6 month approval).</p>

PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	See required medical information
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: November 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadone HCl

Products Affected

- *methadone hcl oral solution 10 mg/5ml*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. (4) FOR A DOCUMENTED DIAGNOSIS OF OPIOID ADDICTION (heroin or other morphine-like drugs): medication must be dispensed by a treatment program certified by SAMHSA (Substance Abuse and Mental Health Services Administration) and the patient will be monitored during therapy for signs and symptoms of abuse/misuse as well as compliance and the potential diversion to others. (Initiation/detoxification treatment = 1 month approval, continuation of therapy/maintenance treatment = 6 month approval).</p>

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PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	See required medical information
Other Criteria	
QL Criteria	30 ml Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadone HCl

Products Affected

- *methadone hcl oral solution 5 mg/5ml*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. (4) FOR A DOCUMENTED DIAGNOSIS OF OPIOID ADDICTION (heroin or other morphine-like drugs): medication must be dispensed by a treatment program certified by SAMHSA (Substance Abuse and Mental Health Services Administration) and the patient will be monitored during therapy for signs and symptoms of abuse/misuse as well as compliance and the potential diversion to others. (Initiation/detoxification treatment = 1 month approval, continuation of therapy/maintenance treatment = 6 month approval).</p>

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PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	See required medical information
Other Criteria	
QL Criteria	60 ml Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadone HCl

Products Affected

- *methadone hcl oral tablet*
- *methadone hcl oral tablet soluble*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. (4) FOR A DOCUMENTED DIAGNOSIS OF OPIOID ADDICTION (heroin or other morphine-like drugs): medication must be dispensed by a treatment program certified by SAMHSA (Substance Abuse and Mental Health Services Administration) and the patient will be monitored during therapy for signs and symptoms of abuse/misuse as well as compliance and the potential diversion to others. (Initiation/detoxification treatment = 1 month approval, continuation of therapy/maintenance treatment = 6 month approval).</p>

PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	See required medical information
Other Criteria	
QL Criteria	6 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadone HCl Intensol

Products Affected

- METHADONE HCL INTENSOL

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. (4) FOR A DOCUMENTED DIAGNOSIS OF OPIOID ADDICTION (heroin or other morphine-like drugs): medication must be dispensed by a treatment program certified by SAMHSA (Substance Abuse and Mental Health Services Administration) and the patient will be monitored during therapy for signs and symptoms of abuse/misuse as well as compliance and the potential diversion to others. (Initiation/detoxification treatment = 1 month approval, continuation of therapy/maintenance treatment = 6 month approval).</p>

PA Criteria	Criteria Details
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	See required medical information
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: November 27, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadose

Products Affected

- METHADOSE ORAL CONCENTRATE 10 MG/ML

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methadose Sugar-Free

Products Affected

- METHADOSE SUGAR-FREE

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methamphetamine HCl

Products Affected

- *methamphetamine hcl*

QL Criteria	4 tablets Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methergine

Products Affected

- METHERGINE ORAL

QL Criteria	28 tablets Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylin

Products Affected

- METHYLIN ORAL SOLUTION 10 MG/5ML

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	30 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylin

Products Affected

- METHYLIN ORAL SOLUTION 5
MG/5ML

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	60 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl

Products Affected

- *methylphenidate hcl oral solution 10 mg/5ml*

QL Criteria	30 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl

Products Affected

- *methylphenidate hcl oral solution 5 mg/5ml*

QL Criteria	60 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl

Products Affected

- *methylphenidate hcl oral tablet*

QL Criteria	6 tablets Per 1 Day
Notes/ References	Annual Review: 10/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl

Products Affected

- *methylphenidate hcl oral tablet chewable*

QL Criteria	6 tablets Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 10 mg*

QL Criteria	3 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 18 mg, 27 mg, 54 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 20 mg*

QL Criteria	3 tab Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 36 mg*

QL Criteria	4 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 24 hour 18 mg, 27 mg, 54 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER

Products Affected

- *methylphenidate hcl er oral tablet extended release 24 hour 36 mg*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER (CD)

Products Affected

- *methylphenidate hcl er (cd)*

QL Criteria	1 caps Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER (LA)

Products Affected

- *methylphenidate hcl er (la) oral capsule extended release 24 hour 20 mg*
- *methylphenidate hcl er (la) oral capsule extended release 24 hour 40 mg*

QL Criteria	1 caps Per 1 Day
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER (LA)

Products Affected

- *methylphenidate hcl er (la) oral capsule
extended release 24 hour 30 mg*

QL Criteria	2 caps Per 1 Day
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Methylphenidate HCl ER (LA)

Products Affected

- *methylphenidate hcl er (la) oral capsule
extended release 24 hour 60 mg*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metoprolol Succinate ER

Products Affected

- *metoprolol succinate er oral tablet extended release 24 hour 100 mg, 50 mg*

QL Criteria	1.5 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metoprolol Succinate ER

Products Affected

- *metoprolol succinate er oral tablet extended release 24 hour 200 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metoprolol Succinate ER

Products Affected

- *metoprolol succinate er oral tablet extended release 24 hour 25 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metoprolol-HCTZ ER

Products Affected

- *metoprolol-hctz er oral tablet extended release 24 hour 100-12.5 mg*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of metoprolol-hydrochlorothiazide tablet or metoprolol succinate er tablet and hydrochlorothiazide tablet
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Metoprolol-HCTZ ER

Products Affected

- *metoprolol-hctz er oral tablet extended release 24 hour 25-12.5 mg, 50-12.5 mg*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of metoprolol-hydrochlorothiazide tablet or metoprolol succinate er tablet and hydrochlorothiazide tablet
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mevacor

Products Affected

- MEVACOR ORAL TABLET 40 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Miacalcin

Products Affected

- MIACALCIN INJECTION

ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Miacalcin

Products Affected

- MIACALCIN NASAL

ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
QL Criteria	1 bottle Per 1 month
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Micardis

Products Affected

- MICARDIS

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Micardis HCT

Products Affected

- MICARDIS HCT

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MiCort-HC

Products Affected

- MICORT-HC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alclometasone cream/oint or Hydrocort
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Microdot Test

Products Affected

- MICRODOT TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Microgestin FE 1/20

Products Affected

- MICROGESTIN FE 1/20

QL Criteria	1.5 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Migranal

Products Affected

- MIGRANAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of generic Migranal and two of the following: naratriptan, rizatriptan, sumatriptan, zolmitriptan
QL Criteria	1 pack Per 30 days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mimvey

Products Affected

- MIMVEY

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Minivelle

Products Affected

- MINIVELLE

QL Criteria	8 patches Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mirapex ER

Products Affected

- MIRAPEX ER

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mircera

Products Affected

- MIRCERA INJECTION SOLUTION
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Erythropoiesis_Stimulating_Agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mirtazapine

Products Affected

- *mirtazapine oral tablet 15 mg, 30 mg, 45 mg* • *mirtazapine oral tablet dispersible*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mitigare

Products Affected

- MITIGARE

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Modafinil

Products Affected

- *modafinil*

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with modafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tab Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mondoxyne NL

Products Affected

- MONDOXYNE NL ORAL CAPSULE 75
MG

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Monoclalte-P

Products Affected

- MONOCLATE-P INTRAVENOUS KIT
1000 UNIT, 1500 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Monodox

Products Affected

- MONODOX ORAL CAPSULE 75 MG

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mononine

Products Affected

- MONONINE INTRAVENOUS SOLUTION
RECONSTITUTED 1000 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Monovisc

Products Affected

- MONOVISC

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Montelukast Sodium

Products Affected

- *montelukast sodium oral*

QL Criteria	1 pack Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Montelukast Sodium

Products Affected

- *montelukast sodium oral*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MorphaBond ER

Products Affected

- MORPHABOND ER

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred alternatives which include Hysingla ER, Embeda and Oxycontin
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Morphine Sulfate

Products Affected

- *morphine sulfate oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Morphine Sulfate ER

Products Affected

- *morphine sulfate er oral capsule extended release 24 hour*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Morphine Sulfate ER

Products Affected

- *morphine sulfate er oral tablet extended release*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Morphine Sulfate ER Beads

Products Affected

- *morphine sulfate er beads*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Movantik

Products Affected

- MOVANTIK

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 03/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MS Contin

Products Affected

- MS CONTIN ORAL TABLET EXTENDED RELEASE

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mupirocin

Products Affected

- *mupirocin external*

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mupirocin Calcium

Products Affected

- *mupirocin calcium*

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Myalept

Products Affected

- MYALEPT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/myalept.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	15 vials Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mydayis

Products Affected

- MYDAYIS

PA Criteria	Criteria Details
Covered Uses	Treatment of Attention Deficit Hyperactivity Disorder (ADHD) in patients 13 years and older
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention Deficit Hyperactivity Disorder (ADHD)
Age Restrictions	13 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 capsule Per 1 day
Notes/References	
Revision Date	Prior Authorization: July 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

MyGlucoHealth Test

Products Affected

- MYGLUCOHEALTH TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Myorisan

Products Affected

- MYORISAN

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Myorisan

Products Affected

- MYORISAN

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mytesi

Products Affected

- MYTESI

PA Criteria	Criteria Details
Covered Uses	Non-infectious diarrhea in patients with HIV/AIDS on anti-retroviral therapy
Exclusion Criteria	
Required Medical Information	Covered for adult members who have a documented diagnosis of noninfectious diarrhea associated with HIV/AIDS infection that has lasted at least for one month and who are currently stable on anti-retroviral therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of at least one anti-motility agent (loperamide, diphenoxylate/atropine, bismuth subsalicylate)
QL Criteria	2 tablets Per 1 Day
Notes/References	Annual Review: 03/2017
Revision Date	Prior Authorization: September 12, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Naftifine HCl

Products Affected

- *naftifine hcl external cream 2 %*

QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Naftin

Products Affected

- NAFTIN EXTERNAL CREAM 2 %
- NAFTIN EXTERNAL GEL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic naftifine 1% cream
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Naglazyme

Products Affected

- NAGLAZYME

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Naprelan

Products Affected

- NAPRELAN ORAL TABLET EXTENDED
RELEASE 24 HOUR 375 MG, 500 MG, 750
MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic immediate release naproxen
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Naratriptan HCl

Products Affected

- *naratriptan hcl*

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nascobal

Products Affected

- NASCOBAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic cyanocobalamin injectable
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nasonex

Products Affected

- NASONEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Natesto

Products Affected

- NATESTO

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, Gender Dysphoria, Gender reassignment surgery
Exclusion Criteria	Excluded in patients with carcinoma of the breast or suspected carcinoma of the prostate or if the patient will be using therapy for muscle building purposes
Required Medical Information	Documented diagnosis of gender dysphoria, documentation of undergoing gender reassignment surgery, or a diagnoses of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available) Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	3 pumps Per 30 Days

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Natpara

Products Affected

- NATPARA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 cartridges Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nefazodone HCl

Products Affected

- *nefazodone hcl oral tablet 250 mg, 50 mg*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one of the following medications: bupropion SR/XL, bupropion/SR/XL, citalopram, escitalopram, fluoxetine, fluvoxamine, paroxetine/sr, mirtazapine, selfemra, sertraline, venlafaxine, venlafaxine er tablet, or venlafaxine sr cap
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nerlynx

Products Affected

- NERLYNX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Nerlynx.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	6 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 02, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nesina

Products Affected

- NESINA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentaducto; and generic alogliptin, alogliptin/pioglitazone, alogliptin/metformin
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neulasta

Products Affected

- NEULASTA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/G-CSF.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neupogen

Products Affected

- NEUPOGEN INJECTION SOLUTION 300 MCG/ML, 480 MCG/1.6ML
- NEUPOGEN INJECTION SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/G-CSF.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neupro

Products Affected

- NEUPRO

QL Criteria	1 patch Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neurontin

Products Affected

- NEURONTIN ORAL CAPSULE

QL Criteria	6 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neurontin

Products Affected

- NEURONTIN ORAL TABLET

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Neutek 2Tek Test

Products Affected

- NEUTEK 2TEK TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NexAVAR

Products Affected

- NEXAVAR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NexGen Test

Products Affected

- NEXGEN TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NexIUM

Products Affected

- NEXIUM ORAL CAPSULE DELAYED
RELEASE 40 MG

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NexIUM

Products Affected

- NEXIUM ORAL PACKET

QL Criteria	1 packet Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Next Choice One Dose

Products Affected

- NEXT CHOICE ONE DOSE

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nicoderm CQ

Products Affected

- NICODERM CQ

QL Criteria	1 patch Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nicotine

Products Affected

- *nicotine*

QL Criteria	1 patch Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nicotine Polacrilex

Products Affected

- *nicotine polacrilex mouth/throat*

QL Criteria	24 pieces Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nicotrol

Products Affected

- NICOTROL

QL Criteria	16 cartridges Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nicotrol NS

Products Affected

- NICOTROL NS

QL Criteria	12 bottles Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ninlaro

Products Affected

- NINLARO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	3 capsules Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nitrostat

Products Affected

- NITROSTAT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of nitroglycerin
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nityr

Products Affected

- NITYR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Norco

Products Affected

- NORCO

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Norditropin FlexPro

Products Affected

- NORDITROPIN FLEXPPO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Norethindrone-Eth Estradiol

Products Affected

- *norethindrone-eth estradiol oral tablet 0.5-2.5 mg-mcg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Northera

Products Affected

- NORTHERA ORAL CAPSULE 100 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northera.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northera.html
QL Criteria	3 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Northera

Products Affected

- NORTHERA ORAL CAPSULE 200 MG, 300 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northera.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northera.html
QL Criteria	6 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nova Max Glucose Test

Products Affected

- NOVA MAX GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Novarel

Products Affected

- *novarel intramuscular solution reconstituted*
10000 unit

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infertility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Novoeight

Products Affected

- NOVOEIGHT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN 70/30

Products Affected

- NOVOLIN 70/30

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN 70/30 ReliOn

Products Affected

- NOVOLIN 70/30 RELION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN N

Products Affected

- NOVOLIN N

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN N ReliOn

Products Affected

- NOVOLIN N RELION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN R

Products Affected

- NOVOLIN R

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLIN R ReliOn

Products Affected

- NOVOLIN R RELION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLOG

Products Affected

- NOVOLOG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLOG FlexPen

Products Affected

- NOVOLOG FLEXPEN SUBCUTANEOUS SOLUTION PEN-INJECTOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLOG Mix 70/30

Products Affected

- NOVOLOG MIX 70/30

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLOG Mix 70/30 FlexPen

Products Affected

- NOVOLOG MIX 70/30 FLEXPEN
SUBCUTANEOUS SUSPENSION PEN-
INJECTOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoLOG PenFill

Products Affected

- NOVOLOG PENFILL SUBCUTANEOUS SOLUTION CARTRIDGE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of one preferred alternative insulin, Humulin or Humalog
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NovoSeven RT

Products Affected

- NOVOSEVEN RT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nplate

Products Affected

- NPLATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Neu mega.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nucala

Products Affected

- NUCALA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/RESP/Interleukin Antagonist.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 injection Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nucynta

Products Affected

- NUCYNTA

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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ST Criteria	A documented contraindication, intolerance, allergy, or failure of two days of immediate release oxycodone or morphine
QL Criteria	120 tablets Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nucynta ER

Products Affected

- NUCYNTA ER

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	60 tablets Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuedexta

Products Affected

- NUEDEXTA

QL Criteria	2 capsules Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuplazid

Products Affected

- NUPLAZID

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/Nuplazid.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nutropin AQ NuSpin 10

Products Affected

- NUTROPIN AQ NUSPIN 10

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nutropin AQ NuSpin 20

Products Affected

- NUTROPIN AQ NUSPIN 20

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nutropin AQ NuSpin 5

Products Affected

- NUTROPIN AQ NUSPIN 5

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuvigil

Products Affected

- NUVIGIL ORAL TABLET 150 MG, 250 MG

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with armodafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tab Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuvigil

Products Affected

- NUVIGIL ORAL TABLET 200 MG

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with armodafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tablet Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuvigil

Products Affected

- NUVIGIL ORAL TABLET 50 MG

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with armodafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tab Per 1 Day

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Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nuwiq

Products Affected

- NUWIQ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nymalize

Products Affected

- NYMALIZE ORAL SOLUTION 60
MG/20ML

QL Criteria	2520 ML Per 21 days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ocaliva

Products Affected

- OCALIVA ORAL TABLET 5 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/Primary_Biliary_Cholangitis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/Primary_Biliary_Cholangitis.html
QL Criteria	1 tablet Per 1 day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Octagam

Products Affected

- OCTAGAM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Octreotide Acetate

Products Affected

- *octreotide acetate injection solution 100 mcg/ml, 1000 mcg/ml, 200 mcg/ml, 50 mcg/ml, 500 mcg/ml*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Sandostatin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Odefsey

Products Affected

- ODEFSEY

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Odomzo

Products Affected

- ODOMZO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ofev

Products Affected

- OFEV

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Idiopathic_Pulmonary_Fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OLANZapine

Products Affected

- *olanzapine oral tablet 10 mg, 15 mg, 20 mg, 5 mg, 7.5 mg*
- *olanzapine oral tablet dispersible*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OLANZapine

Products Affected

- *olanzapine oral tablet 2.5 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OLANZapine-FLUoxetine HCl

Products Affected

- *olanzapine-fluoxetine hcl*

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olmesartan Medoxomil

Products Affected

- *olmesartan medoxomil oral tablet 20 mg, 5 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olmesartan Medoxomil-HCTZ

Products Affected

- *olmesartan medoxomil-hctz*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olmesartan-Amlodipine-HCTZ

Products Affected

- *olmesartan-amlodipine-hctz*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olux

Products Affected

- OLUX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of a generic clobetasol alternative
QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olux-E

Products Affected

- OLUX-E

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of a generic clobetasol alternative
QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Olysio

Products Affected

- OLYSIO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omega-3-acid Ethyl Esters

Products Affected

- *omega-3-acid ethyl esters*

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omeprazole-Sodium Bicarbonate

Products Affected

- *omeprazole-sodium bicarbonate oral capsule*
40-1100 mg

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omeprazole-Sodium Bicarbonate

Products Affected

- *omeprazole-sodium bicarbonate oral packet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required). In addition for approval the following criteria must also be met: Documentation of an inability to swallow tablets/capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 packet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omnaris

Products Affected

- OMNARIS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omnitrope

Products Affected

- OMNITROPE SUBCUTANEOUS SOLUTION

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omnitrope

Products Affected

- OMNITROPE SUBCUTANEOUS SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

On Call Express Blood Glucose

Products Affected

- ON CALL EXPRESS BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

On Call Plus Blood Glucose

Products Affected

- ON CALL PLUS BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

On Call Vivid Blood Glucose

Products Affected

- ON CALL VIVID BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OneTouch Ultra Blue

Products Affected

- ONETOUCH ULTRA BLUE

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OneTouch Verio

Products Affected

- ONETOUCH VERIO IN VITRO STRIP

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Onfi

Products Affected

- ONFI ORAL TABLET 10 MG, 20 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Onglyza

Products Affected

- ONGLYZA

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Onzetra Xsail

Products Affected

- ONZETRA XSAIL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of sumatriptan nasal spray
QL Criteria	1 kit Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Opana

Products Affected

- OPANA ORAL

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Opana ER

Products Affected

- OPANA ER ORAL TABLET ER 12 HOUR ABUSE-DETERRENT

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Opsumit

Products Affected

- OPSUMIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OptumRx Blood Glucose Test

Products Affected

- OPTUMRX BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oracea

Products Affected

- ORACEA

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oravig

Products Affected

- ORAVIG

QL Criteria	14 tab Per 30 Days
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orencia

Products Affected

- ORENCIA INTRAVENOUS PREFILLED SYRINGE 125 MG/ML
- ORENCIA SUBCUTANEOUS SOLUTION

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MUSC/Orencia.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnnonmedicare/data/2017/MUSC/Orencia.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orencia

Products Affected

- ORENCIA SUBCUTANEOUS SOLUTION
 PREFILLED SYRINGE 50 MG/0.4ML, 87.5
 MG/0.7ML

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Orencia.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Orencia.html
QL Criteria	4 syringes Per 1 month
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orencia ClickJect

Products Affected

- ORENCIA CLICKJECT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Orencia.html
QL Criteria	4 syringes Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orenitram

Products Affected

- ORENITRAM ORAL TABLET
EXTENDED RELEASE 0.125 MG, 0.25
MG, 1 MG, 2.5 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orenitram

Products Affected

- ORENITRAM ORAL TABLET
EXTENDED RELEASE 5 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orfadin

Products Affected

- ORFADIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Orkambi

Products Affected

- ORKAMBI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/cystic_fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OrthoVisc

Products Affected

- ORTHOVISC INTRA-ARTICULAR SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oseltamivir Phosphate

Products Affected

- *oseltamivir phosphate oral capsule*

QL Criteria	20 capsules Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oseni

Products Affected

- OSENI

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentaducto; and generic alogliptin, alogliptin/pioglitazone, alogliptin/metformin
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Osphena

Products Affected

- OSPHENA

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Otezla

Products Affected

- OTEZLA ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otezla.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otezla.html
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Otezla

Products Affected

- OTEZLA ORAL TABLET THERAPY PACK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otezla.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otezla.html
QL Criteria	1 pack Per 1 year
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Otrexup

Products Affected

- OTREXUP SUBCUTANEOUS SOLUTION
AUTO-INJECTOR 10 MG/0.4ML, 12.5
MG/0.4ML, 15 MG/0.4ML, 17.5
MG/0.4ML, 20 MG/0.4ML, 22.5
MG/0.4ML, 25 MG/0.4ML

ST Criteria	http://www.aetna.com/products/rxnnonmedicare/data/2017/MUSC/Otrexup_Rasuvo.html
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ovidrel

Products Affected

- OVIDREL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/inferility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxaydo

Products Affected

- OXAYDO

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxiconazole Nitrate

Products Affected

- *oxiconazole nitrate*

QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxistat

Products Affected

- OXISTAT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxistat

Products Affected

- OXISTAT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	60 ml Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxtellar XR

Products Affected

- OXTELLAR XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 150 MG,
300 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxtellar XR

Products Affected

- OXTELLAR XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 600 MG

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyCODONE HCl

Products Affected

- *oxycodone hcl oral capsule*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Last Update 12/2017

QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyCODONE HCl

Products Affected

- *oxycodone hcl oral concentrate 100 mg/5ml*
- *oxycodone hcl oral solution*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyCODONE HCl

Products Affected

- *oxycodone hcl oral tablet*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

2017 Aetna Pharmacy Drug Guide - Self Insured
Last Update 12/2017

QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyCODONE HCl ER

Products Affected

- *oxycodone hcl er oral tablet er 12 hour abuse-deterrent 10 mg, 20 mg, 40 mg, 80 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxycodone-Acetaminophen

Products Affected

- *oxycodone-acetaminophen oral solution*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxycodone-Acetaminophen

Products Affected

- *oxycodone-acetaminophen oral tablet 10-325 mg, 2.5-325 mg, 5-325 mg, 7.5-325 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxycodone-Aspirin

Products Affected

- *oxycodone-aspirin oral tablet 4.8355-325 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxycodone-Ibuprofen

Products Affected

- *oxycodone-ibuprofen*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyCONTIN

Products Affected

- OXYCONTIN ORAL TABLET ER 12 HOUR ABUSE-DETERRENT

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxymorphone HCl

Products Affected

- *oxymorphone hcl*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

OxyMORphone HCl ER

Products Affected

- *oxymorphone hcl er*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paliperidone ER

Products Affected

- *paliperidone er oral tablet extended release*
24 hour 1.5 mg, 3 mg, 6 mg

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paliperidone ER

Products Affected

- *paliperidone er oral tablet extended release*
24 hour 9 mg

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pancreaze

Products Affected

- PANCREAZE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Creon and Zenpep
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paricalcitol

Products Affected

- *paricalcitol oral*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PARoxetine HCl

Products Affected

- *paroxetine hcl oral tablet 10 mg, 20 mg*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PARoxetine HCl

Products Affected

- *paroxetine hcl oral tablet 30 mg, 40 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PARoxetine HCl ER

Products Affected

- *paroxetine hcl er*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PARoxetine Mesylate

Products Affected

- *paroxetine mesylate*

PA Criteria	Criteria Details
Covered Uses	Moderate to severe vasomotor symptoms associated with menopause
Exclusion Criteria	
Required Medical Information	A documented diagnosis of moderate to severe vasomotor symptoms associated with menopause
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 capsule Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 28, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paxil

Products Affected

- PAXIL ORAL SUSPENSION

QL Criteria	30 pen Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paxil

Products Affected

- PAXIL ORAL TABLET 10 MG, 20 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paxil

Products Affected

- PAXIL ORAL TABLET 30 MG, 40 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Paxil CR

Products Affected

- PAXIL CR

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pegasys

Products Affected

- PEGASYS SUBCUTANEOUS SOLUTION

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pegasys ProClick

Products Affected

- PEGASYS PROCLICK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PegIntron

Products Affected

- PEGINTRON

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pennsaid

Products Affected

- PENNSAID TRANSDERMAL SOLUTION
2 %

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of Voltaren Gel
QL Criteria	4 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pentasa

Products Affected

- PENTASA ORAL CAPSULE EXTENDED
RELEASE 250 MG

QL Criteria	16 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pentasa

Products Affected

- PENTASA ORAL CAPSULE EXTENDED
RELEASE 500 MG

QL Criteria	8 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pentazocine-Naloxone HCl

Products Affected

- *pentazocine-naloxone hcl*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Percocet

Products Affected

- PERCO CET ORAL TABLET 10-325 MG, 2.5-325 MG, 5-325 MG, 7.5-325 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Perforomist

Products Affected

- PERFOROMIST

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Serevent (Step Therapy will not apply to members who have a documented inability to use an inhaler)
QL Criteria	60 vials (120ml) Per 1 fill
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pertzye

Products Affected

- PERTZYE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Creon and Zenpep
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pexeva

Products Affected

- PEXEVA ORAL TABLET 10 MG, 20 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of paroxetine
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pexeva

Products Affected

- PEXEVA ORAL TABLET 30 MG, 40 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of paroxetine
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pharmacist Choice Autocode

Products Affected

- PHARMACIST CHOICE AUTOCODE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Phenadoz

Products Affected

- PHENADOZ

PA Criteria	Criteria Details
Covered Uses	Administration of analgesic (prophylaxis), Allergic condition, Motion sickness, Nausea and vomiting, Postoperative pain, Sedation
Exclusion Criteria	
Required Medical Information	Member is less than 6 years of age and the member's physician provides documentation (controlled clinical trial) from the peer reviewed medical literature that supports use in specified indication for this age group.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Note: FDA alert ⁶ - Use of Phenergan/promethazine is contraindicated in infants and children less than 2 years of age due to risks for fatal respiratory depression. In addition the use of promethazine and codeine (with or without phenylephrine) is contraindicated in pediatric patients less than 6 years of age.
Notes/References	
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Phenoxybenzamine HCl

Products Affected

- *phenoxybenzamine hcl oral*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Picato

Products Affected

- PICATO

QL Criteria	1 tube Per 60 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Plavix

Products Affected

- PLAVIX ORAL TABLET 75 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Plegridy

Products Affected

- PLEGRIDY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Plegridy Starter Pack

Products Affected

- PLEGRIDY STARTER PACK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Plexion Cleansing Cloth

Products Affected

- PLEXION CLEANSING CLOTH
EXTERNAL PAD

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred generic sulfacetamide sodium with sulfur products
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PocketChem EZ Test

Products Affected

- POCKETCHEM EZ TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pomalyst

Products Affected

- POMALYST

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Praluent

Products Affected

- PRALUENT SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
QL Criteria	2 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pramipexole Dihydrochloride ER

Products Affected

- *pramipexole dihydrochloride er oral tablet extended release 24 hour 2.25 mg, 3.75 mg, 4.5 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prasugrel HCl

Products Affected

- *prasugrel hcl*

PA Criteria	Criteria Details
Covered Uses	Acute coronary syndrome (ACS) managed with percutaneous coronary intervention which includes unstable angina or non-ST elevation myocardial infarction or ST elevation myocardial infarction (MI)
Exclusion Criteria	History of Stroke or transient ischemic attack (TIA)
Required Medical Information	Member has a documented diagnosis of acute coronary syndrome (ACS) and is managed by percutaneous coronary intervention (PCI), which includes unstable angina, non-ST-elevation myocardial infarction (NSTEMI), or ST -elevation myocardial infarction (STEMI) managed with primary or delayed PCI and member has no prior history of stroke or transient ischemic attack (TIA)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tablet Per 1 day
Notes/References	
Revision Date	Prior Authorization: May 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pravachol

Products Affected

- PRAVACHOL ORAL TABLET 20 MG, 40 MG, 80 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pravastatin Sodium

Products Affected

- *pravastatin sodium*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision PCx

Products Affected

- PRECISION PCX

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision PCX Plus Test

Products Affected

- PRECISION PCX PLUS TEST

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision Point of Care Test

Products Affected

- PRECISION POINT OF CARE TEST

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision QID Test

Products Affected

- PRECISION QID TEST

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision Sof-Tact Test

Products Affected

- PRECISION SOF-TACT TEST

QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Precision Xtra Blood Glucose

Products Affected

- PRECISION XTRA BLOOD GLUCOSE

QL Criteria	300 strips Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prefest

Products Affected

- PREFEST

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pregnyl

Products Affected

- *pregnyl*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/infer-tility.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prevacid

Products Affected

- PREVACID ORAL CAPSULE DELAYED RELEASE 30 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 caps Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Prevacid SoluTab

Products Affected

- PREVACID SOLUTAB

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required). In addition for approval the following criteria must also be met: Documentation of an inability to swallow tablets/capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 tab Per 1 Day
Notes/References	Annual Review: 02/2017

Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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PriLOSEC

Products Affected

- PRILOSEC ORAL PACKET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required). In addition for approval the following criteria must also be met: Documentation of an inability to swallow tablets/capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	2 pack Per 1 Day
Notes/References	Annual Review: 02/2017

Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Primlev

Products Affected

- PRIMLEV

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pristiq

Products Affected

- PRISTIQ

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 tab Per 1 Day
Notes/References	Annual Review: 05/2017

Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Pristiq

Products Affected

- PRISTIQ

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 05/2017

Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Privigen

Products Affected

- PRIVIGEN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/ivig.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ProAir RespiClick

Products Affected

- PROAIR RESPICLICK

QL Criteria	2 inhalers Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ProCentra

Products Affected

- PROCENTRA

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD), Narcolepsy
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD) OR Narcolepsy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	40 ml Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: January 25, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Procrit

Products Affected

- PROCRIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Erythropoiesis_Stimulating_Agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Proctocort

Products Affected

- PROCTOCORT RECTAL CREAM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic hydrocortisone rectal cream
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Procysbi

Products Affected

- PROCYSBI ORAL CAPSULE DELAYED
RELEASE 25 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
QL Criteria	4 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Procysbi

Products Affected

- PROCYSBI ORAL CAPSULE DELAYED
RELEASE 75 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
QL Criteria	25 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prodigy No Coding Blood Gluc

Products Affected

- PRODIGY NO CODING BLOOD GLUC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Profilnine SD

Products Affected

- PROFILNINE SD

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prolastin-C

Products Affected

- PROLASTIN-C INTRAVENOUS
SOLUTION RECONSTITUTED 1000 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Alpha-1 Antitrypsin Inhibitor Therapy.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prolia

Products Affected

- PROLIA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Promacta

Products Affected

- PROMACTA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Promacta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Promethazine HCl

Products Affected

- *promethazine hcl oral* 25 mg
- *promethazine hcl rectal suppository 12.5 mg,*

PA Criteria	Criteria Details
Covered Uses	Administration of analgesic (prophylaxis), Allergic condition, Motion sickness, Nausea and vomiting, Postoperative pain, Sedation
Exclusion Criteria	
Required Medical Information	Member is less than 6 years of age and the member's physician provides documentation (controlled clinical trial) from the peer reviewed medical literature that supports use in specified indication for this age group.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Note: FDA alert ⁶ - Use of Phenergan/promethazine is contraindicated in infants and children less than 2 years of age due to risks for fatal respiratory depression. In addition the use of promethazine and codeine (with or without phenylephrine) is contraindicated in pediatric patients less than 6 years of age.
Notes/References	
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Promethegan

Products Affected

- PROMETHEGAN

PA Criteria	Criteria Details
Covered Uses	Administration of analgesic (prophylaxis), Allergic condition, Motion sickness, Nausea and vomiting, Postoperative pain, Sedation
Exclusion Criteria	
Required Medical Information	Member is less than 6 years of age and the member's physician provides documentation (controlled clinical trial) from the peer reviewed medical literature that supports use in specified indication for this age group.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Note: FDA alert6- Use of Phenergan/promethazine is contraindicated in infants and children less than 2 years of age due to risks for fatal respiratory depression. In addition the use of promethazine and codeine (with or without phenylephrine) is contraindicated in pediatric patients less than 6 years of age.
Notes/References	
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Proscar

Products Affected

- PROSCAR

PA Criteria	Criteria Details
Covered Uses	Benign prostatic hyperplasia
Exclusion Criteria	
Required Medical Information	Member is greater than 50 years old or has diagnosis of BPH (Benign Prostatic Hyperplasia). For female members, must have a documented diagnosis of hirsutism secondary to ovarian or adrenal dysfunction (for example, polycystic ovary syndrome, adrenal or ovarian tumor)and must not be pregnant.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: October 25, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Protopic

Products Affected

- PROTOPIC

PA Criteria	Criteria Details
Covered Uses	Atopic dermatitis, Vitiligo
Exclusion Criteria	
Required Medical Information	FOR PROTOPIC 0.1%: A documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or an adolescent 16 years of age or older with either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas. FOR PROTOPIC 0.03%: A documented diagnosis of mild to moderate atopic dermatitis (eczema) in patients less than 2 years of age for short-term use (up to 3 months)(Note: requirement of a trial of topical corticosteroid is not required) or a documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or child 2 years of age or older and either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	

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ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks (14 days) of one preferred alternative topical corticosteroid (triamcinolone acetonide, fluocinonide cream, augmented betamethasone gel, betamethasone dipropionate, hydrocortisone valerate, or fluticasone propionate ointment)
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 26, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Proventil HFA

Products Affected

- PROVENTIL HFA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 week each of Ventolin HFA AND Proair
Notes/ References	Annual Review: 03/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Provigil

Products Affected

- PROVIGIL

PA Criteria	Criteria Details
Covered Uses	Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder
Exclusion Criteria	
Required Medical Information	FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with modafinil therapy, and the daytime fatigue is significantly impacting, impairing, or compromising the patient's ability to function normally, and the prescribing physician has established a patient care plan to treat the cause of OSAHS in conjunction with treating the daily fatigue, and patient must be compliant with recommendations for OSAHS treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tab Per 1 Day

Notes/ References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PROzac

Products Affected

- PROZAC ORAL CAPSULE 10 MG

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PROzac

Products Affected

- PROZAC ORAL CAPSULE 20 MG

QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PROzac

Products Affected

- PROZAC ORAL CAPSULE 40 MG

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prudoxin

Products Affected

- PRUDOXIN

QL Criteria	45 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PTS Panels Glucose Test

Products Affected

- PTS PANELS GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pulmicort

Products Affected

- PULMICORT

QL Criteria	4 ml Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pulmicort Flexhaler

Products Affected

- PULMICORT FLEXHALER

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Asmanex, QVAR, or Flovent
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pulmozyme

Products Affected

- PULMOZYME

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/cystic_fibrosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Purixan

Products Affected

- PURIXAN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	100 ml Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Qbrelis

Products Affected

- QBRELIS

PA Criteria	Criteria Details
Covered Uses	Hypertension, Heart Failure, Myocardial Infarction
Exclusion Criteria	
Required Medical Information	A documented diagnosis of hypertension (Approved only for ages 6 and older), Heart failure, or Myocardial Infarction AND must have a documented inability to swallow tablets/capsules
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 09, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Quasense

Products Affected

- QUASENSE

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Qudexy XR

Products Affected

- QUDEXY XR

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate

Products Affected

- *quetiapine fumarate oral tablet 100 mg, 50 mg*

QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate

Products Affected

- *quetiapine fumarate oral tablet 200 mg*

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate

Products Affected

- *quetiapine fumarate oral tablet 25 mg*

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate

Products Affected

- *quetiapine fumarate oral tablet 300 mg, 400 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate ER

Products Affected

- *quetiapine fumarate er oral tablet extended release 24 hour 150 mg, 200 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QUetiapine Fumarate ER

Products Affected

- *quetiapine fumarate er oral tablet extended release 24 hour 300 mg, 400 mg, 50 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QuickTek Test

Products Affected

- QUICKTEK TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

QuilliChew ER

Products Affected

- QUILICHEW ER ORAL TABLET
CHEWABLE EXTENDED RELEASE 20
MG, 40 MG

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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QuilliChew ER

Products Affected

- QUILICHEW ER ORAL TABLET
CHEWABLE EXTENDED RELEASE 30
MG

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Quillivant XR

Products Affected

- QUILLIVANT XR

PA Criteria	Criteria Details
Covered Uses	Attention deficit hyperactivity disorder (ADHD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Attention deficit hyperactivity disorder (ADHD)
Age Restrictions	For Quillivant Only- 17 years of age and older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	12 ML Per 1 Day
Notes/References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Quintet AC Blood Glucose Test

Products Affected

- QUINTET AC BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Quintet Blood Glucose Test

Products Affected

- QUINTET BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RA TRUEtest Test

Products Affected

- RA TRUETEST TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RABEprazole Sodium

Products Affected

- *rabeprazole sodium*

QL Criteria	1 tab Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ranexa

Products Affected

- RANEXA ORAL TABLET EXTENDED
RELEASE 12 HOUR 1000 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ranexa

Products Affected

- RANEXA ORAL TABLET EXTENDED
RELEASE 12 HOUR 500 MG

QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rasagiline Mesylate

Products Affected

- *rasagiline mesylate oral*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rasuvo

Products Affected

- RASUVO SUBCUTANEOUS SOLUTION
AUTO-INJECTOR 10 MG/0.2ML, 12.5
MG/0.25ML, 15 MG/0.3ML, 17.5
MG/0.35ML, 20 MG/0.4ML, 22.5
MG/0.45ML, 25 MG/0.5ML, 30 MG/0.6ML,
7.5 MG/0.15ML

ST Criteria	<a href="http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otr
exup_Rasuvo.html">http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Otr exup_Rasuvo.html
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ravicti

Products Affected

- RAVICTI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rayaldee

Products Affected

- RAYALDEE

PA Criteria	Criteria Details
Covered Uses	Treatment of secondary hyperparathyroidism in adult patients with stage 3 or 4 chronic kidney disease (CKD)
Exclusion Criteria	Patients with stage 5 CKD or in patients with end stage renal disease (ESRD) on dialysis
Required Medical Information	A documented diagnosis of secondary hyperparathyroidism and Stage 3 or 4 chronic kidney disease (CKD) and serum total 25-hydroxyvitamin D level is less than 30 ng/mL
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of calcitriol
QL Criteria	1 capsule Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 13, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rayos

Products Affected

- RAYOS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of prednisone
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rebif

Products Affected

- REBIF SUBCUTANEOUS SOLUTION
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rebif Rebidose

Products Affected

- REBIF REBIDOSE SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rebif Rebidoose Titration Pack

Products Affected

- REBIF REBIDOSE TITRATION PACK
SUBCUTANEOUS SOLUTION AUTO-
INJECTOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rebif Titration Pack

Products Affected

- REBIF TITRATION PACK
SUBCUTANEOUS SOLUTION
PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Reclast

Products Affected

- RECLAST

QL Criteria	1 bottle Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Recombinate

Products Affected

- RECOMBINATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RefuAH Plus Blood Glucose Test

Products Affected

- REFUAH PLUS BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Regranex

Products Affected

- REGRANEX

PA Criteria	Criteria Details
Covered Uses	Treatment of lower extremity diabetic neuropathic ulcers
Exclusion Criteria	Documentation that the patient has NONE of the following: Neoplasm(s) at the sites(s) of application, will not be using in pressure ulcers, venous stasis ulcers, or ischemic diabetic ulcers, exposed joints, tendons, ligaments, and bone (at application site), or will not be using in wounds that close by primary intention (such as suturing or gluing)
Required Medical Information	A documented diagnosis of diabetes with lower extremity neuropathic ulcers that extend into the subcutaneous tissue or beyond with adequate blood supply
Age Restrictions	16 years or older
Prescriber Restrictions	
Coverage Duration	20 weeks
Other Criteria	NOTE: The safety and efficacy of treatment beyond 20 weeks have not been determined.
QL Criteria	30 grams Per 30 Days
Notes/References	
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: November 06, 2017

Relenza Diskhaler

Products Affected

- RELENZA DISKHALER

QL Criteria	2 EA Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Relistor

Products Affected

- RELISTOR ORAL

PA Criteria	Criteria Details
Covered Uses	Opioid-induced constipation (OIC) in adults with chronic non-cancer pain
Exclusion Criteria	
Required Medical Information	A documented diagnosis of opioid induced constipation due to non-cancer pain and documented concomitant use of opioid therapy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	3 tablets Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Relistor

Products Affected

- RELISTOR SUBCUTANEOUS
SOLUTION 12 MG/0.6ML

QL Criteria	0.6 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Relistor

Products Affected

- RELISTOR SUBCUTANEOUS
SOLUTION 8 MG/0.4ML

QL Criteria	0.4 ml Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Relpax

Products Affected

- RELPAX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan
QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Remeron

Products Affected

- REMERON

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Remeron SolTab

Products Affected

- REMERON SOLTAB

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Remicade

Products Affected

- REMICADE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Remicade.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Remicade.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Remodulin

Products Affected

- REMODULIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Renflexis

Products Affected

- RENFLEXIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Renflexis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Renflexis.html
Notes/References	
Revision Date	Prior Authorization: August 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Repatha

Products Affected

- REPATHA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
QL Criteria	2 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Repatha Pushtronex System

Products Affected

- REPATHA PUSHTRONEX SYSTEM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
QL Criteria	1 syringe Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Repatha SureClick

Products Affected

- REPATHA SURECLICK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html
QL Criteria	2 syringes Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Requip XL

Products Affected

- REQUIP XL ORAL TABLET EXTENDED
RELEASE 24 HOUR 12 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Requip XL

Products Affected

- REQUIP XL ORAL TABLET EXTENDED
RELEASE 24 HOUR 2 MG, 4 MG, 6 MG, 8
MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rescula

Products Affected

- RESCULA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of latanoprost and one week of Travatan Z
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Restoril

Products Affected

- RESTORIL ORAL CAPSULE 22.5 MG, 7.5 MG

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Retin-A

Products Affected

- RETIN-A

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Retin-A Micro

Products Affected

- RETIN-A MICRO

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Retin-A Micro Pump

Products Affected

- RETIN-A MICRO PUMP

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Revatio

Products Affected

- REVATIO INTRAVENOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Revatio

Products Affected

- REVATIO ORAL SUSPENSION
RECONSTITUTED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
QL Criteria	224 ml Per 1 fill
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Revatio

Products Affected

- REVATIO ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Reveal Blood Glucose Test

Products Affected

- REVEAL BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Revlimid

Products Affected

- REVLIMID

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rexall Blood Glucose Test

Products Affected

- REXALL BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rexulti

Products Affected

- REXULTI

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine).
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 08/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rhofade

Products Affected

- RHOFADE

QL Criteria	4 tubes Per 1 year
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RiaSTAP

Products Affected

- RIASTAP

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Rias tap.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: November 17, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Riax

Products Affected

- RIAx

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic benzoyl peroxide foam
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rightest GS100 Blood Glucose

Products Affected

- RIGHTEST GS100 BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rightest GS300 Blood Glucose

Products Affected

- RIGHTEST GS300 BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rightest GS550 Blood Glucose

Products Affected

- RIGHTEST GS550 BLOOD GLUCOSE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rilutek

Products Affected

- RILUTEK

PA Criteria	Criteria Details
Covered Uses	amyotrophic lateral sclerosis (ALS)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of amyotrophic lateral sclerosis (ALS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Riluzole

Products Affected

- *riluzole*

PA Criteria	Criteria Details
Covered Uses	amyotrophic lateral sclerosis (ALS)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of amyotrophic lateral sclerosis (ALS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Risedronate Sodium

Products Affected

- *risedronate sodium oral tablet 150 mg*

QL Criteria	1 tablet Per 30 Days
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Risedronate Sodium

Products Affected

- *risedronate sodium oral tablet 30 mg, 5 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Risedronate Sodium

Products Affected

- *risedronate sodium oral tablet 35 mg*

QL Criteria	4 tablets Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperDAL

Products Affected

- RISPERDAL ORAL SOLUTION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperDAL

Products Affected

- RISPERDAL ORAL TABLET 0.25 MG, 0.5 MG, 1 MG, 2 MG, 3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperDAL

Products Affected

- RISPERDAL ORAL TABLET 4 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperDAL M-TAB

Products Affected

- RISPERDAL M-TAB ORAL TABLET DISPERSIBLE 0.5 MG
- RISPERDAL M-TAB ORAL TABLET DISPERSIBLE 1 MG, 3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperDAL M-TAB

Products Affected

- RISPERDAL M-TAB ORAL TABLET
DISPERSIBLE 4 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperiDONE

Products Affected

- *risperidone oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg, 3 mg*
- *risperidone oral tablet dispersible 0.25 mg, 0.5 mg, 3 mg*
- *risperidone oral tablet dispersible 1 mg, 2 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperiDONE

Products Affected

- *risperidone oral tablet 4 mg*
- *risperidone oral tablet dispersible 4 mg*

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperiD ONE M-TAB

Products Affected

- RISPERIDONE M-TAB ORAL TABLET
DISPERSIBLE 0.5 MG, 1 MG, 2 MG, 3 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RisperiD ONE M-TAB

Products Affected

- RISPERIDONE M-TAB ORAL TABLET
DISPERSIBLE 4 MG

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ritalin

Products Affected

- RITALIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	6 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ritalin LA

Products Affected

- RITALIN LA ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 10 MG, 40 MG
- RITALIN LA ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 20 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 caps Per 1 Day
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ritalin LA

Products Affected

- RITALIN LA ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 30 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 caps Per 1 Day
Notes/ References	Annual Review: 09/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rixubis

Products Affected

- RIXUBIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rizatriptan Benzoate

Products Affected

- *rizatriptan benzoate*

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ROPINIRole HCl ER

Products Affected

- *ropinirole hcl er oral tablet extended release*
24 hour 12 mg

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ROPINIRole HCl ER

Products Affected

- *ropinirole hcl er oral tablet extended release*
24 hour 2 mg, 4 mg, 6 mg, 8 mg

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rosuvastatin Calcium

Products Affected

- *rosuvastatin calcium*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Roxicodone

Products Affected

- ROXICODONE ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rozerem

Products Affected

- ROZEREM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 7 days (one week) of the preferred generic alternative zolpidem or zolpidem er.
QL Criteria	1 tab Per 1 Day
Notes/ References	Annual Review: 08/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rubraca

Products Affected

- RUBRACA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Rubraca.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: January 09, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ruconest

Products Affected

- RUCONEST

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/hereditary_angioedema.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rydapt

Products Affected

- RYDAPT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Rydapt.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 capsules Per 1 day
Notes/References	
Revision Date	Prior Authorization: June 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

RyVent

Products Affected

- RYVENT

PA Criteria	Criteria Details
Covered Uses	Seasonal and perennial allergic rhinitis, vasomotor rhinitis, allergic conjunctivitis due to inhalant allergens and foods, mild uncomplicated allergic skin manifestations of urticaria and angioedema, dermatographism, ss therapy for anaphylactic reactions adjunctive to epinephrine, mmelioration of the severity of allergic reactions to blood or plasma.
Exclusion Criteria	
Required Medical Information	A documented diagnosis of seasonal and perennial allergic rhinitis, vasomotor rhinitis, allergic conjunctivitis due to inhalant allergens and foods, mild uncomplicated allergic skin manifestations of urticaria and angioedema, dermatographism, as therapy for anaphylactic reactions adjunctive to epinephrine, or amelioration of the severity of allergic reactions to blood or plasma.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of carbinoxamine
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Last Update 12/2017

Sabril

Products Affected

- SABRIL ORAL PACKET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/anticonvulsants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sabril

Products Affected

- SABRIL ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/anticonvulsants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	6 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Saizen

Products Affected

- SAIZEN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Saizen Click.Easy

Products Affected

- SAIZEN CLICK.EASY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Samsca

Products Affected

- SAMSCA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/samsca.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sancuso

Products Affected

- SANCUSO

QL Criteria	1 patch Per 21 Days
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SandoSTATIN

Products Affected

- SANDOSTATIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Sandostatin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SandoSTATIN LAR Depot

Products Affected

- SANDOSTATIN LAR DEPOT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Sandostatin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Santyl

Products Affected

- SANTYL

QL Criteria	60 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Saphris

Products Affected

- SAPHRIS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Saphris

Products Affected

- SAPHRIS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Savella

Products Affected

- SAVELLA

QL Criteria	2 tab Per 1 Day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Savella Titration Pack

Products Affected

- SAVELLA TITRATION PACK

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Seasonique

Products Affected

- SEASONIQUE

QL Criteria	90 days maximum Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Seebri Neohaler

Products Affected

- SEEBRI NEOHALER

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of either Spiriva or Incruse Ellipta
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Selzentry

Products Affected

- SELZENTRY ORAL SOLUTION

QL Criteria	8 bottles Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Selzentry

Products Affected

- SELZENTRY ORAL TABLET 25 MG

QL Criteria	8 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Selzentry

Products Affected

- SELZENTRY ORAL TABLET 75 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Semprex-D

Products Affected

- SEMPREX-D

QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sensipar

Products Affected

- SENSIPAR

PA Criteria	Criteria Details
Covered Uses	Secondary Hyperparathyroidism (HPT) in adult patients with chronic kidney disease (CKD) on dialysis, Hypercalcemia in adult patients with Parathyroid Carcinoma, or Hypercalcemia in adult patients with primary HPT for whom parathyroidectomy would be indicated on the basis of serum calcium levels, but who are unable to undergo parathyroidectomy.
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of Secondary Hyperparathyroidism (HPT) in an adult patient with chronic kidney disease (CKD) on dialysis, Hypercalcemia in an adult patient with parathyroid carcinoma (PC), or Hypercalcemia in an adult patient with primary hyperparathyroidism for whom parathyroidectomy would be indicated on the basis of serum calcium levels, but who are unable to undergo parathyroidectomy
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Serevent Diskus

Products Affected

- SEREVENT DISKUS

QL Criteria	1 box Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel

Products Affected

- SEROQUEL ORAL TABLET 100 MG, 50 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel

Products Affected

- SEROQUEL ORAL TABLET 200 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel

Products Affected

- SEROQUEL ORAL TABLET 25 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel

Products Affected

- SEROQUEL ORAL TABLET 300 MG
- SEROQUEL ORAL TABLET 400 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel XR

Products Affected

- SEROQUEL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 150 MG,
200 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel XR

Products Affected

- SEROQUEL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 300 MG,
400 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel XR

Products Affected

- SEROQUEL XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 50 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Serostim

Products Affected

- SEROSTIM SUBCUTANEOUS SOLUTION RECONSTITUTED 4 MG, 5 MG, 6 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sertraline HCl

Products Affected

- *sertraline hcl oral concentrate*

QL Criteria	10 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sertraline HCl

Products Affected

- *sertraline hcl oral tablet 100 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sertraline HCl

Products Affected

- *sertraline hcl oral tablet 25 mg*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sertraline HCl

Products Affected

- *sertraline hcl oral tablet 50 mg*

QL Criteria	45 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Signifor

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Signifor.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sildenafil Citrate

Products Affected

- *sildenafil citrate oral*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Silenor

Products Affected

- SILENOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 7 days each of generic doxepin and either zolpidem or zolpidem er
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Siliq

Products Affected

- SILIQ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Siliq.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Siliq.html
QL Criteria	2 injections Per 1 month
Notes/References	
Revision Date	Prior Authorization: July 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Simponi

Products Affected

- SIMPONI SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- SIMPONI SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Simponi.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Simponi.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Simponi Aria

Products Affected

- SIMPONI ARIA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Simponi_Aria.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Simponi_Aria.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Simvastatin

Products Affected

- *simvastatin oral*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Singulair

Products Affected

- SINGULAIR

QL Criteria	1 pack Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Singulair

Products Affected

- SINGULAIR

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sirturo

Products Affected

- SIRTURO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antimycobacterial_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	68 tablets Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sitavig

Products Affected

- SITAVIG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of oral acyclovir tablets
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sivextro

Products Affected

- SIVEXTRO ORAL

QL Criteria	6 tablets Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Smart Sense Premium Test

Products Affected

- SMART SENSE PREMIUM TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Smart Sense Value Test

Products Affected

- SMART SENSE VALUE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Smartest Blood Glucose Test

Products Affected

- SMARTEST BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sodium Phenylbutyrate

Products Affected

- *sodium phenylbutyrate oral powder 3 gm/tsp* • *sodium phenylbutyrate oral tablet*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Solaraze

Products Affected

- SOLARAZE

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Soliqua

Products Affected

- SOLIQUA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one of the following: Victoza, Byetta, Bydureon, Tanzeum, Trulicity, Adylixin, Lantus, Toujeo, Levemir, Tresiba, Basaglar
QL Criteria	5 pens Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Solus V2 Test

Products Affected

- SOLUS V2 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Somatuline Depot

Products Affected

- SOMATULINE DEPOT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/San_dostatin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Somavert

Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sonata

Products Affected

- SONATA ORAL CAPSULE 10 MG

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sonata

Products Affected

- SONATA ORAL CAPSULE 5 MG

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Soriatane

Products Affected

- SORIATANE ORAL CAPSULE 10 MG, 17.5 MG, 25 MG

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sorilux

Products Affected

- SORILUX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sovaldi

Products Affected

- SOVALDI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Spiriva HandiHaler

Products Affected

- SPIRIVA HANDIHALER

QL Criteria	1 box Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Spiriva Respimat

Products Affected

- SPIRIVA RESPIMAT INHALATION
AEROSOL SOLUTION 1.25 MCG/ACT

QL Criteria	1 inhaler Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Spiriva Respimat

Products Affected

- SPIRIVA RESPIMAT INHALATION
AEROSOL SOLUTION 2.5 MCG/ACT

QL Criteria	1 inhaler Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Spritam

Products Affected

- SPRITAM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of immediate release levitiracetam tablets
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sprix

Products Affected

- SPRIX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketorolac tablets
QL Criteria	5 days maximum Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sprycel

Products Affected

- SPRYCEL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stimate

Products Affected

- STIMATE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/misendocrine.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stiolto Respimat

Products Affected

- STIOLTO RESPIMAT

QL Criteria	1 inhaler Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stivarga

Products Affected

- STIVARGA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Strattera

Products Affected

- STRATTERA ORAL CAPSULE 10 MG, 18 MG, 25 MG, 40 MG, 60 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Strattera

Products Affected

- STRATTERA ORAL CAPSULE 100 MG,
80 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Strensiq

Products Affected

- STRENSIQ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Striant

Products Affected

- STRIANT

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, Gender Dysphoria, Gender reassignment surgery
Exclusion Criteria	Excluded in patients with carcinoma of the breast or suspected carcinoma of the prostate or if the patient will be using therapy for muscle building purposes
Required Medical Information	Documented diagnosis of gender dysphoria, documentation of undergoing gender reassignment surgery, or a diagnoses of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available) Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	2 buccals Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stribild

Products Affected

- STRIBILD

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviral_hiv.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Striverdi Respimat

Products Affected

- STRIVERDI RESPIMAT

PA Criteria	Criteria Details
Covered Uses	Chronic Obstructive Pulmonary Disorder (COPD)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Serevent
QL Criteria	1 inhaler Per 1 month
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Suboxone

Products Affected

- SUBOXONE SUBLINGUAL FILM 12-3
MG

QL Criteria	2 pack Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Suboxone

Products Affected

- SUBOXONE SUBLINGUAL FILM 2-0.5 MG, 4-1 MG, 8-2 MG

QL Criteria	90 pack Per 30 Days
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Subsys

Products Affected

- SUBSYS

PA Criteria	Criteria Details
Covered Uses	Breakthrough cancer pain, General anesthesia
Exclusion Criteria	
Required Medical Information	Documentation that member is terminally ill or has a documented diagnosis of cancer with concomitant use of around the clock long acting opioid therapy for cancer pain, requiring management of breakthrough pain and is intolerant of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step therapy may apply
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of the preferred generic alternative, fentanyl transmucosal lozenge, and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 sprays Per 30 Days
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 29, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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SulfaSALazine

Products Affected

- *sulfasalazine oral*

QL Criteria	8 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sulfazine

Products Affected

- SULFAZINE

QL Criteria	8 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SUMatriptan Succinate

Products Affected

- *sumatriptan succinate oral*

QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sumavel DosePro

Products Affected

- SUMAVEL DOSEPRO SUBCUTANEOUS SOLUTION JET-INJECTOR

QL Criteria	6 syringes Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Supartz

Products Affected

- SUPARTZ INTRA-ARTICULAR SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Supprelin LA

Products Affected

- SUPPRELIN LA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Supreme Test

Products Affected

- SUPREME TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sure Edge Test

Products Affected

- SURE EDGE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SureChek Blood Glucose Test

Products Affected

- SURECHEK BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sure-Test EasyPlus Mini Test

Products Affected

- SURE-TEST EASYPLUS MINI TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sutent

Products Affected

- SUTENT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sylatron

Products Affected

- SYLATRON SUBCUTANEOUS KIT 200 MCG, 300 MCG, 600 MCG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Symbicort

Products Affected

- SYMBICORT

QL Criteria	1 inhaler Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Symbyax

Products Affected

- SYMBYAX

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Symbyax

Products Affected

- SYMBYAX

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SymLinPen 120

Products Affected

- SYMLINPEN 120 SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Covered Uses	All FDA Approved uses
Exclusion Criteria	Poor compliance with current insulin regimen, Poor compliance with prescribed self-blood glucose monitorings, An A1C greater than 9%, Recurrent severe hypoglycemia requiring assistance during the previous 6 months, Presence of hypoglycemia unawareness, Confirmed diagnosis of gastroparesis, Need for medications that stimulate GI motility , Patient is less than 18 years old, Concurrent use with other oral antidiabetic medications (except metformin and sulfonylureas) or drugs that alter gastrointestinal motility
Required Medical Information	A documented diagnosis of type 1 or type 2 diabetes mellitus and the patient concurrently using rapid or short-acting insulin (e.g., Humalog or regular insulin). For extended renewals: a documented diagnosis of type 1 or type 2 diabetes mellitus and the patient concurrently using rapid or short-acting insulin (e.g., Humalog or regular insulin), and the patient demonstrated an expected reduction in HbA1c since starting this therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	initial: 6 months - extended: 12 months
Other Criteria	
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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SymLinPen 60

Products Affected

- SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Covered Uses	All FDA Approved uses
Exclusion Criteria	Poor compliance with current insulin regimen, Poor compliance with prescribed self-blood glucose monitorings, An A1C greater than 9%, Recurrent severe hypoglycemia requiring assistance during the previous 6 months, Presence of hypoglycemia unawareness, Confirmed diagnosis of gastroparesis, Need for medications that stimulate GI motility , Patient is less than 18 years old, Concurrent use with other oral antidiabetic medications (except metformin and sulfonylureas) or drugs that alter gastrointestinal motility
Required Medical Information	A documented diagnosis of type 1 or type 2 diabetes mellitus and the patient concurrently using rapid or short-acting insulin (e.g., Humalog or regular insulin). For extended renewals: a documented diagnosis of type 1 or type 2 diabetes mellitus and the patient concurrently using rapid or short-acting insulin (e.g., Humalog or regular insulin), and the patient demonstrated an expected reduction in HbA1c since starting this therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	initial: 6 months - extended: 12 months
Other Criteria	
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: June 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Symproic

Products Affected

- SYMPROIC

PA Criteria	Criteria Details
Covered Uses	Treatment of opioid-induced constipation (OIC) in adult patients with chronic non-cancer pain
Exclusion Criteria	Patients with known or suspected gastrointestinal obstruction or at increased risk of recurrent obstruction or with a history of a hypersensitivity reaction to naldemedine
Required Medical Information	A documented diagnosis of opioid-induced constipation (OIC) in adult patients with chronic non-cancer pain and the patient has been taking opioids for 4 weeks or more
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Movantik or Relistor
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: November 06, 2017 Quantity Limits: November 15, 2017

Synagis

Products Affected

- SYNAGIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Synagis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synalgos-DC

Products Affected

- SYNALGOS-DC

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	<p>(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.</p>
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synarel

Products Affected

- SYNAREL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Syndros

Products Affected

- SYNDROS

PA Criteria	Criteria Details
Covered Uses	Anorexia associated with weight loss in patients with AIDS, Chemotherapy-induced nausea and vomiting
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Anorexia associated with weight loss in patients with AIDS, or Chemotherapy-induced nausea and vomiting
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
QL Criteria	4 bottles Per 1 month
Notes/References	
Revision Date	Prior Authorization: July 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synera

Products Affected

- SYNERA

QL Criteria	10 patches Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synjardy

Products Affected

- SYNJARDY

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synjardy XR

Products Affected

- SYNJARDY XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 10-1000
MG, 12.5-1000 MG, 5-1000 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synjardy XR

Products Affected

- SYNJARDY XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 25-1000
MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synvisc

Products Affected

- SYNVISC INTRA-ARTICULAR SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Synvisc One

Products Affected

- SYNVISC ONE INTRA-ARTICULAR SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/viscosupplements.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Syprine

Products Affected

- SYPRINE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Taclonex

Products Affected

- TACLONEX EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of a medium to high potency topical steroid
QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Taclonex

Products Affected

- TACLONEX EXTERNAL SUSPENSION

QL Criteria	60 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tacrolimus

Products Affected

- *tacrolimus external ointment 0.03 %*

PA Criteria	Criteria Details
Covered Uses	Atopic dermatitis, Vitiligo
Exclusion Criteria	
Required Medical Information	FOR PROTOPIC 0.1%: A documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or an adolescent 16 years of age or older with either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas. FOR PROTOPIC 0.03%: A documented diagnosis of mild to moderate atopic dermatitis (eczema) in patients less than 2 years of age for short-term use (up to 3 months)(Note: requirement of a trial of topical corticosteroid is not required) or a documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or child 2 years of age or older and either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks (14 days) of one preferred alternative topical corticosteroid (triamcinolone acetonide, fluocinonide cream, augmented betamethasone gel, betamethasone dipropionate, hydrocortisone valerate, or fluticasone propionate ointment)
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 26, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tacrolimus

Products Affected

- *tacrolimus external ointment 0.1 %*

PA Criteria	Criteria Details
Covered Uses	Atopic dermatitis, Vitiligo
Exclusion Criteria	
Required Medical Information	FOR PROTOPIC 0.1%: A documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or an adolescent 16 years of age or older with either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas. FOR PROTOPIC 0.03%: A documented diagnosis of mild to moderate atopic dermatitis (eczema) in patients less than 2 years of age for short-term use (up to 3 months)(Note: requirement of a trial of topical corticosteroid is not required) or a documented diagnosis of atopic dermatitis (eczema) or vitiligo in an adult or child 2 years of age or older and either a documented contraindication, intolerance or allergy to one preferred alternative topical corticosteroid indicated for the patient's condition, or a documented failure of an adequate trial of 2 weeks (14 days) of one preferred alternative topical corticosteroid indicated for the patient's condition, or the treatment is in an area at high risk for skin atrophy such as face, eyelids, or genital areas.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks (14 days) of one preferred alternative topical corticosteroid (triamcinolone acetonide, fluocinonide cream, augmented betamethasone gel, betamethasone dipropionate, hydrocortisone valerate, or fluticasone propionate ointment)
QL Criteria	60 GM Per 1 fill
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 26, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tafinlar

Products Affected

- TAFINLAR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tagrisso

Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 TABLET Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Taltz

Products Affected

- TALTZ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Taltz.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Taltz.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tamiflu

Products Affected

- TAMIFLU ORAL CAPSULE

QL Criteria	20 capsules Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tamiflu

Products Affected

- TAMIFLU ORAL SUSPENSION
RECONSTITUTED 6 MG/ML

QL Criteria	480 pen Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tanzeum

Products Affected

- TANZEUM

QL Criteria	4 pens Per 1 month
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tarceva

Products Affected

- TARCEVA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tasigna

Products Affected

- TASIGNA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tazarotene

Products Affected

- *tazarotene external*

PA Criteria	Criteria Details
Covered Uses	acne vulgaris, plaque psoriasis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of acne vulgaris or plaque psoriasis
Age Restrictions	Prior authorization only required in patients older than 35 years of age.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tazorac

Products Affected

- TAZORAC

PA Criteria	Criteria Details
Covered Uses	acne vulgaris, plaque psoriasis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of acne vulgaris or plaque psoriasis
Age Restrictions	Prior authorization only required in patients older than 35 years of age.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tecfidera

Products Affected

- TECFIDERA ORAL RELEASE 240 MG
- TECFIDERA ORAL CAPSULE DELAYED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tecfidera

Products Affected

- TECFIDERA ORAL CAPSULE DELAYED
RELEASE 120 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
QL Criteria	14 capsules Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Technivie

Products Affected

- TECHNIVIE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tekturna

Products Affected

- TEKTURNA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 preferred ACE/ARB medications, such as lisinopril, quinapril, enalapril, benazepril, candesartan, irbesartan, losartan, or telmisartan
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tekturna HCT

Products Affected

- TEKTURNA HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 preferred ACE/ARB medications, such as lisinopril, quinapril, enalapril, benazepril, candesartan, irbesartan, losartan, or telmisartan
QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Telcare Blood Glucose Test

Products Affected

- TELCARE BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Telmisartan

Products Affected

- *telmisartan*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Telmisartan-HCTZ

Products Affected

- *telmisartan-hctz*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Temazepam

Products Affected

- *temazepam oral capsule 22.5 mg, 7.5 mg*

QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Temodar

Products Affected

- TEMODAR ORAL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Temovate

Products Affected

- TEMOVATE EXTERNAL CREAM
- TEMOVATE EXTERNAL OINTMENT
- TEMOVATE EXTERNAL GEL

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Temovate

Products Affected

- TEMOVATE EXTERNAL SOLUTION

QL Criteria	100 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Temozolomide

Products Affected

- *temozolomide*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Testim

Products Affected

- TESTIM

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	10 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Testosterone

Products Affected

- *testosterone transdermal gel 10 mg/act (2%)*

QL Criteria	4 grams Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Testosterone

Products Affected

- *testosterone transdermal gel 12.5 mg/act (1%)*
- *testosterone transdermal gel 50 mg/5gm (1%)*

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	10 grams Per 1 Day

Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Testosterone

Products Affected

- *testosterone transdermal gel 25 mg/2.5gm (1%)*

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	2.5 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Testosterone

Products Affected

- *testosterone transdermal solution*

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	6 milliliters Per 1 Day
Notes/References	

Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Tetrabenazine

Products Affected

- *tetrabenazine oral tablet 12.5 mg*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/xenazine.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 31, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tetrabenazine

Products Affected

- *tetrabenazine oral tablet 25 mg*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/xenazine.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 31, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TGT Blood Glucose Test

Products Affected

- *tgt blood glucose test*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Thalomid

Products Affected

- THALOMID

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Thiola

Products Affected

- THIOLA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TiaGABine HCl

Products Affected

- *tiagabine hcl oral tablet 2 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TiaGABine HCl

Products Affected

- *tiagabine hcl oral tablet 4 mg*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tilia Fe

Products Affected

- TILIA FE

QL Criteria	1.5 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tivorbex

Products Affected

- TIVORBEX

QL Criteria	3 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tobi Podhaler

Products Affected

- TOBI PODHALER

QL Criteria	8 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tolak

Products Affected

- TOLAK

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic fluorouracil 5% cream
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Topamax Sprinkle

Products Affected

- TOPAMAX SPRINKLE

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Topiramate

Products Affected

- *topiramate oral capsule sprinkle*

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Toviaz

Products Affected

- TOVIAZ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare or Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, Tolterodine ER, oxybutynin, oxybutynin XL)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tracleer

Products Affected

- TRACLEER

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tradjenta

Products Affected

- TRADJENTA

QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 05/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TraMADol HCl

Products Affected

- *tramadol hcl oral*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TraMADol HCl ER

Products Affected

- *tramadol hcl er oral capsule extended release 24 hour 100 mg, 200 mg, 300 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TraMADol HCl ER

Products Affected

- *tramadol hcl er oral tablet extended release 24 hour*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TraMADol HCl ER (Biphasic)

Products Affected

- *tramadol hcl er (biphasic) oral tablet
extended release 24 hour 100 mg, 200 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TraMADol HCl ER (Biphasic)

Products Affected

- *tramadol hcl er (biphasic) oral tablet extended release 24 hour 300 mg*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tramadol-Acetaminophen

Products Affected

- *tramadol-acetaminophen*

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tranexamic Acid

Products Affected

- *tranexamic acid oral*

QL Criteria	30 tablets Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trelegy Ellipta

Products Affected

- TRELEGY ELLIPTA

QL Criteria	1 inhaler Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: November 15, 2017

Trelstar Mixject

Products Affected

- TRELSTAR MIXJECT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tremfya

Products Affected

- TREMFYA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Tremfya.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Tremfya.html
QL Criteria	1 injection Per 56 Days
Notes/References	
Revision Date	Prior Authorization: August 02, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretinoin

Products Affected

- *tretinoin external*

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretinoin

Products Affected

- *tretinoin oral*

QL Criteria	30 days maximum Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretinoin Microsphere

Products Affected

- *tretinoin microsphere*

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretinoin Microsphere Pump

Products Affected

- *tretinoin microsphere pump*

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretin-X

Products Affected

- TRETIN-X EXTERNAL CREAM 0.075 %

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred alternatives indicated for the members condition, one of which has to be topical tretinoin.
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretin-X

Products Affected

- TRETIN-X EXTERNAL CREAM 0.075 %

PA Criteria	Criteria Details
Covered Uses	Acne vulgaris
Exclusion Criteria	
Required Medical Information	A documented diagnosis of any one of the following: (1) Acne vulgaris (includes comedonal, cystic, nodular & papular acne), (2) Actinic keratoses and lesions are on the face or lesions are not on the face and therapy includes the use of 5-fluorouracil in conjunction with tretinoin, (3) Hypertrophic scars or keloids and intralesional injection of corticosteroids is ineffective or not tolerated, (4) Keratosis follicularis (Darier's disease, Darier-White disease), (5) Facial flat warts, or (6) Multiple flat warts (includes common warts and plantar warts)
Age Restrictions	greater than 35
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretten

Products Affected

- TRETEN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Treximet

Products Affected

- TREXIMET

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan
QL Criteria	9 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Treximet

Products Affected

- TREXIMET

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan
QL Criteria	9 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trezix

Products Affected

- TREZIX ORAL CAPSULE 320.5-30-16 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 capsules Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tribenzor

Products Affected

- TRIBENZOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives, which include Exforge HCT, telmisartan/amlodipine used in combination with hctz, and any of the following medications used in combination with amlodipine: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, valsartan/hctz, or telmisartan/hctz
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tricor

Products Affected

- TRICOR

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Triglide

Products Affected

- TRIGLIDE ORAL TABLET 160 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tri-Legest Fe

Products Affected

- TRI-LEGEST FE

QL Criteria	1.5 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trintellix

Products Affected

- TRINTELLIX

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 tablet Per 1 Day
Notes/References	

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Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Triptodur

Products Affected

- TRIPTODUR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trokendi XR

Products Affected

- TROKENDI XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 100 MG,
25 MG
- TROKENDI XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 50 MG

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trokendi XR

Products Affected

- TROKENDI XR ORAL CAPSULE
EXTENDED RELEASE 24 HOUR 200 MG

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TRUEtest Test

Products Affected

- TRUETEST TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

TrueTrack Test

Products Affected

- TRUETRACK TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trulance

Products Affected

- TRULANCE

PA Criteria	Criteria Details
Covered Uses	Chronic idiopathic constipation (CIC)
Exclusion Criteria	
Required Medical Information	A documented diagnosis of chronic idiopathic constipation
Age Restrictions	18 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of lactulose and either Linzess or Amitiza
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: April 05, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trulicity

Products Affected

- TRULICITY

QL Criteria	4 pens (2 ml) Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Truvada

Products Affected

- TRUVADA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviral_hiv.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tudorza Pressair

Products Affected

- TUDORZA PRESSAIR INHALATION
AEROSOL POWDER BREATH
ACTIVATED

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of either Spiriva or Incruse Ellipta
QL Criteria	1 pack Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tybost

Products Affected

- TYBOST

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tykerb

Products Affected

- TYKERB

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tylenol with Codeine #3

Products Affected

- TYLENOL WITH CODEINE #3

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tylenol with Codeine #4

Products Affected

- TYLENOL WITH CODEINE #4

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tymlos

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
QL Criteria	1 pen Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tysabri

Products Affected

- TYSABRI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tyvaso

Products Affected

- TYVASO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tyvaso Refill

Products Affected

- TYVASO REFILL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tyvaso Starter

Products Affected

- TYVASO STARTER

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uceris

Products Affected

- UCERIS ORAL

PA Criteria	Criteria Details
Covered Uses	ulcerative colitis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of active, mild to moderate ulcerative colitis and a documented contraindication or intolerance or allergy or failure of an adequate trial of one month each of two preferred 5-ASA therapies (i.e., balsalazide, Canasa, Delzicol) and one preferred generic corticosteroid therapy (i.e., budesonide sr, prednisone, prednisolone)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tab Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 21, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uceris

Products Affected

- UCERIS RECTAL

PA Criteria	Criteria Details
Covered Uses	ulcerative colitis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of ACTIVE mild to moderate distal ulcerative colitis extending up to 40 cm from the anal verge, requiring induction of remission.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	4 canisters Per 1 month
Notes/References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uloric

Products Affected

- ULORIC

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ultima Test

Products Affected

- ULTIMA TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ultracet

Products Affected

- ULTRACET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ultram

Products Affected

- ULTRAM

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

UltraTRAK PRO Test

Products Affected

- ULTRATRAK PRO TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

UltraTRAK Ultimate Test

Products Affected

- ULTRATRAK ULTIMATE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ultravate

Products Affected

- ULTRAVATE EXTERNAL CREAM
- ULTRAVATE EXTERNAL OINTMENT

QL Criteria	50 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ultravate

Products Affected

- ULTRAVATE EXTERNAL LOTION

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Unistrip1 Generic

Products Affected

- UNISTRIP1 GENERIC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uptravi

Products Affected

- UPTRAVI ORAL TABLET 1000 MCG, 1200 MCG, 1400 MCG, 1600 MCG, 400 MCG, 600 MCG, 800 MCG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uptravi

Products Affected

- UPTRAVI ORAL TABLET 200 MCG PACK
- UPTRAVI ORAL TABLET THERAPY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uroxatral

Products Affected

- UROXATRAL

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Utibron Neohaler

Products Affected

- UTIBRON NEOHALER

PA Criteria	Criteria Details
Covered Uses	Chronic Ostructive Pulmonary Disease (COPD)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Chronic obstructive pulmonary disease (COPD)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month of either Anoro Ellipta or Stiolto
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valchlor

Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valcyte

Products Affected

- VALCYTE ORAL SOLUTION
RECONSTITUTED

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviraltopical.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valcyte

Products Affected

- VALCYTE ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviraltopical.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	102 tablets Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ValGANciclovir HCl

Products Affected

- *valganciclovir hcl oral solution reconstituted*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviraltopical.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1000 milliliters Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ValGANciclovir HCl

Products Affected

- *valganciclovir hcl oral tablet*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ID/antiviraltopical.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	102 tablets Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valsartan-Hydrochlorothiazide

Products Affected

- *valsartan-hydrochlorothiazide*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valsartan-Hydrochlorothiazide

Products Affected

- *valsartan-hydrochlorothiazide*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valtrex

Products Affected

- VALTRESX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic valacyclovir
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vanatol LQ

Products Affected

- VANATOL LQ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two days of one generic alternative tablet/capsule combination of acetaminophen/butalbital/caffeine
QL Criteria	90 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vanos

Products Affected

- VANOS

QL Criteria	120 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vantas

Products Affected

- VANTAS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Varubi

Products Affected

- VARUBI ORAL

QL Criteria	4 tablets Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vascepa

Products Affected

- VASCEPA ORAL CAPSULE 0.5 GM

QL Criteria	8 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vascepa

Products Affected

- VASCEPA ORAL CAPSULE 1 GM

QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vasotec

Products Affected

- VASOTEC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of enalapril and two other Angiotensin Converting Enzyme (ACE) Inhibitors
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vecamyl

Products Affected

- VECAMYL

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html
QL Criteria	10 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Veletri

Products Affected

- VELETRI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Veltassa

Products Affected

- VELTASSA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Veltassa.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 packet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vemlidy

Products Affected

- VEMLIDY

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/Vemlidy.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/Vemlidy.html
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: December 13, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venclexta

Products Affected

- VENCLEXTA ORAL TABLET 10 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Venclexta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	40 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venclexta

Products Affected

- VENCLEXTA ORAL TABLET 100 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Venclexta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venclexta

Products Affected

- VENCLEXTA ORAL TABLET 50 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Venclexta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venclexta Starting Pack

Products Affected

- VENCLEXTA STARTING PACK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Venclexta.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 pack Per 28 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl

Products Affected

- *venlafaxine hcl oral tablet 100 mg, 25 mg*

QL Criteria	3 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl

Products Affected

- *venlafaxine hcl oral tablet 37.5 mg*

QL Criteria	4 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl

Products Affected

- *venlafaxine hcl oral tablet 50 mg*

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl

Products Affected

- *venlafaxine hcl oral tablet 75 mg*

QL Criteria	5 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl ER

Products Affected

- *venlafaxine hcl er oral capsule extended release 24 hour 150 mg*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl ER

Products Affected

- *venlafaxine hcl er oral capsule extended release 24 hour 37.5 mg, 75 mg*

QL Criteria	1 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl ER

Products Affected

- *venlafaxine hcl er oral tablet extended release 24 hour 150 mg*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Venlafaxine HCl ER

Products Affected

- *venlafaxine hcl er oral tablet extended release 24 hour 225 mg, 75 mg*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ventavis

Products Affected

- VENTAVIS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmonaryhypertensionagents.html
Notes/References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Verdeso

Products Affected

- VERDESO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic desonide
QL Criteria	100 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Verdrocet

Products Affected

- VERDROCET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Versacloz

Products Affected

- VERSACLOZ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of clozapine
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Verzenio

Products Affected

- VERZENIO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Verzenio.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: November 16, 2017

Viberzi

Products Affected

- VIBERZI

QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vicodin

Products Affected

- VICODIN ORAL TABLET 5-300 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vicodin ES

Products Affected

- VICODIN ES ORAL TABLET 7.5-300 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vicodin HP

Products Affected

- VICODIN HP ORAL TABLET 10-300 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Victory AGM-4000 Test

Products Affected

- VICTORY AGM-4000 TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Victoza

Products Affected

- VICTOZA SUBCUTANEOUS SOLUTION
PEN-INJECTOR

QL Criteria	3 pens Per 30 Days
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Viekira Pak

Products Affected

- VIEKIRA PAK

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Viekira XR

Products Affected

- VIEKIRA XR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	84 tablets Per 1 month
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vigabatrin

Products Affected

- *vigabatrin*

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/anticonvulsants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	6 packets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Viibryd

Products Affected

- VIIBRYD ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
QL Criteria	1 tab Per 1 Day
Notes/References	Annual Review: 05/2017

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Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Viibryd Starter Pack

Products Affected

- VIIBRYD STARTER PACK

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For quantities over the allowed amount for the prescribed medication, a member must meet one of the following: (1) Member requires a dose including half tablets, (2) member's dose is being titrated by physician (3-month limit), (3) member has had intolerance to drug administered as a single daily dose, or (4) member's dose cannot be achieved with proposed quantity limits for a given strength (ex. needs 375mg per day and would require 5 capsules of venlafaxine sr cap or Effexor XR 75mg to achieve dose.)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 3 different antidepressants from at least two different therapeutic subclasses. Examples include SSRIs (fluoxetine, citalopram), SNRIs (duloxetine, venlafaxine), TCAs (amitriptyline, nortriptyline), and heterocyclic antidepressants (mirtazapine, trazodone)
Notes/References	Annual Review: 05/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Vimizim

Products Affected

- VIMIZIM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vimovo

Products Affected

- VIMOVO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two weeks of one preferred generic non steroidal anti-inflammatory agent
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vimpat

Products Affected

- VIMPAT ORAL SOLUTION

QL Criteria	40 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vimpat

Products Affected

- VIMPAT ORAL TABLET 100 MG, 150 MG, 200 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vimpat

Products Affected

- VIMPAT ORAL TABLET 50 MG

QL Criteria	6 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Viokace

Products Affected

- VIOKACE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Creon and Zenpep
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vistogard

Products Affected

- VISTOGARD

QL Criteria	20 packs Per 1 prescription
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Visudyne

Products Affected

- VISUDYNE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/EYE/opthalmic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vivelle-Dot

Products Affected

- VIVELLE-DOT

QL Criteria	8 patch Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vivlodex

Products Affected

- VIVLODEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two generic non steroidal anti-inflammatory drugs
QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vocal Point Blood Glucose Test

Products Affected

- VOCAL POINT BLOOD GLUCOSE TEST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vogelxo

Products Affected

- VOGELXO TRANSDERMAL GEL 50 MG/5GM (1%)

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	10 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vogelxo Pump

Products Affected

- VOGELXO PUMP

PA Criteria	Criteria Details
Covered Uses	Primary hypogonadism or hypogonadotropic hypogonadism, gender dysphoria, gender reassignment
Exclusion Criteria	Patients with carcinoma of the breast or suspected carcinoma of the prostate, patient will be using therapy for muscle building purposes
Required Medical Information	A documented diagnosis of primary hypogonadism or hypogonadotropic hypogonadism as defined by either one of the following: (1) Member has undergone bilateral orchiectomy (no total fasting serum testosterone levels required), or (2) Having two consecutive low total fasting serum testosterone levels (below the testing laboratory's reference range or below 300ng/dl if reference ranges are not available), or for persons with low normal total testosterone levels (above 300 ng/dL but below 400 ng/dL), two consecutive low free or bioavailable fasting serum testosterone levels (below the testing laboratory's reference range or less than 225 picomoles per liter (pmol/L) (6 ng/dL) if reference ranges are not available)(Note: Two morning samples drawn between 7:00 a.m. and 10:00 a.m. obtained on two different days is required for NEW starts only), or (3) Member has a documented diagnosis of gender dysphoria or documentation of undergoing gender reassignment surgery.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62%
QL Criteria	10 grams Per 1 Day

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Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Voltaren

Products Affected

- VOLTAREN TRANSDERMAL

QL Criteria	200 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vonvendi

Products Affected

- VONVENDI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vosevi

Products Affected

- VOSEVI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Votrient

Products Affected

- VOTRIENT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vpriv

Products Affected

- VPRIV

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: ?http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gaucher_disease.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vraylar

Products Affected

- VRAYLAR ORAL CAPSULE 1.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	4 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vraylar

Products Affected

- VRAYLAR ORAL CAPSULE 3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vraylar

Products Affected

- VRAYLAR ORAL CAPSULE 4.5 MG, 6 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vraylar

Products Affected

- VRAYLAR ORAL CAPSULE THERAPY PACK

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vytorin

Products Affected

- VYTORIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic simvastatin in combination with generic ezetimibe, or generic ezetimibe-simvastatin
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vyvanse

Products Affected

- VYVANSE

QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Vyvanse

Products Affected

- VYVANSE

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

WaveSense Presto

Products Affected

- WAVESENSE PRESTO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of our preferred formulary Lifescan and Abbott products such as Onetouch and Freestyle
QL Criteria	300 EA Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Wellbutrin SR

Products Affected

- WELLBUTRIN SR

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Wellbutrin XL

Products Affected

- WELLBUTRIN XL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of bupropion XL and two selective serotonin reuptake inhibitors (SSRIs)
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Wilate

Products Affected

- WILATE INTRAVENOUS KIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xadago

Products Affected

- XADAGO

PA Criteria	Criteria Details
Covered Uses	Adjunctive treatment to levodopa/carbidopa in patients with Parkinson's disease (PD) experiencing "off" episodes
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Parkinson's disease and concurrent use of levodopa/carbidopa
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of rasagaline or selegiline
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: June 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Xalatan

Products Affected

- XALATAN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of latanoprost and one week of Travatan Z
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xalkori

Products Affected

- XALKORI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xanax

Products Affected

- XANAX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic alprazolam
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xanax XR

Products Affected

- XANAX XR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic alprazolam
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xatmep

Products Affected

- XATMEP

PA Criteria	Criteria Details
Covered Uses	Treatment of acute lymphoblastic leukemia (ALL) or polyarticular juvenile idiopathic arthritis (pJIA) in pediatric patients
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Acute Lymphoblastic Leukemia (ALL) in a pediatric patient (18 years and younger) as part of a multi-phase, combination chemotherapy maintenance regimen or a diagnosis of Polyarticular Juvenile Idiopathic Arthritis (PJIA) in pediatric patients (18 years and younger) who have had an insufficient therapeutic response to, or are intolerant of, an adequate trial of first-line therapy including full dose non-steroidal anti-inflammatory agents (NSAIDs). Regardless of diagnosis, the patient must have a documented inability to swallow tablets/capsules.
Age Restrictions	Approved for those 18 years of age or younger
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: July 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xeljanz

Products Affected

- XELJANZ

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Xeljanz_XeljanzXR.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Xeljanz_XeljanzXR.html
QL Criteria	2 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xeljanz XR

Products Affected

- XELJANZ XR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Xeljanz_XeljanzXR.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Xeljanz_XeljanzXR.html
QL Criteria	1 tablet Per 1 day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xeloda

Products Affected

- XELODA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xenazine

Products Affected

- XENAZINE ORAL TABLET 12.5 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/xenazine.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 31, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xenazine

Products Affected

- XENAZINE ORAL TABLET 25 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/xenazine.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 31, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xeomin

Products Affected

- XEOMIN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/botulinum_toxin.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xermelo

Products Affected

- XERMELO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Xermelo.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	3 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: April 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xgeva

Products Affected

- XGEVA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xhance

Products Affected

- XHANCE

PA Criteria	Criteria Details
Covered Uses	Treatment of nasal polyps in patients 18 years of age or older
Exclusion Criteria	Hypersensitivity to any ingredient in Xhance
Required Medical Information	A documented diagnosis of nasal polyps
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic mometasone nasal
QL Criteria	1 spray bottle Per 30 Days
Notes/References	
Revision Date	Prior Authorization: November 06, 2017 Step Therapy: November 06, 2017 Quantity Limits: November 16, 2017

Xifaxan

Products Affected

- XIFAXAN ORAL TABLET 200 MG

QL Criteria	9 tablets Per 1 fill
Notes/ References	Annual Review: 04/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xifaxan

Products Affected

- XIFAXAN ORAL TABLET 550 MG

PA Criteria	Criteria Details
Covered Uses	Hepatic Encephalopathy, Irritable Bowel Syndrome (IBS) with Diarrhea.
Exclusion Criteria	
Required Medical Information	FOR HEPATIC ENCEPHALOPATHY: Member must have a documented diagnosis and be 18 years and older. FOR IBS WITH DIARRHEA: Member must have a documented diagnosis and must have been prescribed a 14-day course of therapy with three times a day dosing. For reauthorization of 2nd or 3rd course of therapy, there must be at least a 10-week treatment free period from the previous course of therapy.
Age Restrictions	18 years or older
Prescriber Restrictions	
Coverage Duration	HEPATIC ENCEPHALOPATHY: 1 year. IBS: 14 days.
Other Criteria	
QL Criteria	3 tablets Per 1 Day
Notes/References	Annual Review: 04/2017
Revision Date	Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xigduo XR

Products Affected

- XIGDUO XR ORAL TABLET EXTENDED
RELEASE 24 HOUR 10-1000 MG, 10-500
MG, 5-500 MG

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xigduo XR

Products Affected

- XIGDUO XR ORAL TABLET EXTENDED
RELEASE 24 HOUR 5-1000 MG

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ximino

Products Affected

- XIMINO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one of the following: minocycline cap 50mg, 75mg, 100mg; doxycycline monohydrate cap 50mg, 100mg; doxycycline hyclate cap 50mg, 100mg; or doxycycline hyclate tab 100mg
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: November 02, 2017 Quantity Limits: August 25, 2015

Xodol

Products Affected

- XODOL

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

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QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xolair

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/RESP/Xolair.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/RESP/Xolair.html
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xolegel

Products Affected

- XOLEGEL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic ketoconazole cream
QL Criteria	50 gm Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xopenex HFA

Products Affected

- XOPENEX HFA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 week each of Ventolin HFA AND Proair
QL Criteria	2 inhalers Per 1 fill
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xtampza ER

Products Affected

- XTAMPZA ER

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred alternatives which include Hysingla ER, Embeda and Oxycontin
QL Criteria	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xtandi

Products Affected

- XTANDI

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xulane

Products Affected

- XULANE

QL Criteria	3 patches Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xultophy

Products Affected

- XULTOPHY

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one of the following: Victoza, Byetta, Bydureon, Tanzeum, Trulicity, Adylixin, Lantus, Toujeo, Levemir, Tresiba, Basaglar
QL Criteria	5 pens Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xuriden

Products Affected

- XURIDEN

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/metabolic_agents.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 packets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xylon

Products Affected

- XYLON

QL Criteria	120 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: November 16, 2017

Xyntha

Products Affected

- XYNTHA INTRAVENOUS KIT 1000 UNIT, 2000 UNIT, 250 UNIT, 500 UNIT

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xyntha Solofuse

Products Affected

- XYNTHA SOLOFUSE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/bloodproducts_coagulants.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Xyrem

Products Affected

- XYREM

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/cataplaxy-xyrem.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	18 ml Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Yosprala

Products Affected

- YOSPRALA

PA Criteria	Criteria Details
Covered Uses	Documented history of cardiovascular or cerebrovascular events
Exclusion Criteria	
Required Medical Information	A documented history of cardiovascular or cerebrovascular events in a patient greater than 55 years of age or a patient who has a documented history of gastric ulcers
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 40 mg dosage of omeprazole in combination with aspirin
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zafirlukast

Products Affected

- *zafirlukast*

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zaleplon

Products Affected

- *zaleplon oral capsule 10 mg*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zaleplon

Products Affected

- *zaleplon oral capsule 5 mg*

QL Criteria	4 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zamicet

Products Affected

- ZAMICET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information
Other Criteria	

Notes/ References	
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zarxio

Products Affected

- ZARXIO

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/G-CSF.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zavesca

Products Affected

- ZAVESCA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: ?http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/gaucher_disease.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zegerid

Products Affected

- ZEGERID ORAL CAPSULE 40-1100 MG

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 caps Per 1 Day
Notes/References	
Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zegerid

Products Affected

- ZEGERID ORAL PACKET

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented diagnosis of one of the following: Gastroesophageal reflux disease, Complications related to GERD (e.g. esophageal strictures, Barrett's Esophagus), Peptic ulcer disease, Treatment and prevention of gastroduodenal ulcers associated with NSAIDs, Zollinger-Ellison Syndrome, or Helicobacter pylori eradication (Additional documentation of two concurrent antibiotics (i.e. amoxicillin or clarithromycin or metronidazole or tetracycline) that will be used in the treatment regimen combined with the requested PPI as part of the therapy are required). In addition for approval the following criteria must also be met: Documentation of an inability to swallow tablets/capsules.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of 2 generic alternatives: omeprazole, pantoprazole, esomeprazole, or lansoprazole
QL Criteria	1 pack Per 1 Day
Notes/References	

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Revision Date	Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015
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Zejula

Products Affected

- ZEJULA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Zejula.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	3 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: May 09, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zelapar

Products Affected

- ZELAPAR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic selegiline
QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zelboraf

Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	8 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zemaira

Products Affected

- ZEMAIRA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Alpha-1 Antitrypsin Inhibitor Therapy.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zembrace SymTouch

Products Affected

- ZEMBRACE SYMTOUCH

ST Criteria	A documented contraindication, intolerance, allergy, or failure of generic Imitrex injection
QL Criteria	8 syringes Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zemplar

Products Affected

- ZEMPLAR ORAL CAPSULE 1 MCG, 2 MCG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of paricalcitol and calcitriol
QL Criteria	1 capsule Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zenatane

Products Affected

- ZENATANE

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zenatane

Products Affected

- ZENATANE

PA Criteria	Criteria Details
Covered Uses	Severe recalcitrant nodular or cystic acne
Exclusion Criteria	
Required Medical Information	Member is enrolled in the FDA iPLEDGE program and, because of significant adverse reactions associated with its use, should be reserved for patients with multiple severe nodular acne who are unresponsive to conventional therapy, including topical acne products and systemic antibiotics
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 capsules Per 1 Day
Notes/References	Annual Review: 02/2016
Revision Date	Prior Authorization: August 22, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zenzedi

Products Affected

- ZENZEDI ORAL TABLET 10 MG, 5 MG

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zenzedi

Products Affected

- ZENZEDI ORAL TABLET 15 MG, 2.5 MG, 20 MG, 30 MG, 7.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexamethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zepatier

Products Affected

- ZEPATIER

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/GI/hepatitis_c.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zetia

Products Affected

- ZETIA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of ezetimibe
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zetonna

Products Affected

- ZETONNA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zileuton ER

Products Affected

- *zileuton er*

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zinbryta

Products Affected

- ZINBRYTA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html
QL Criteria	1 injection Per 30 Days
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Ziprasidone HCl

Products Affected

- *ziprasidone hcl*

QL Criteria	2 caps Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zocor

Products Affected

- ZOCOR

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zodex 12-Day

Products Affected

- ZODEX 12-DAY

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented inability to use single tablets or reach appropriate dosing regimen with using single tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of dexamethasone tabs
Notes/References	
Revision Date	Prior Authorization: October 04, 2017 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

Zohydro ER

Products Affected

- ZOHYDRO ER ORAL CAPSULE ER 12 HOUR ABUSE-DETERRENT

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	(1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Length of Therapy; see required medical information

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PA Criteria	Criteria Details
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two preferred alternatives which include Hysingla ER, Embeda and Oxycontin
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoladex

Products Affected

- ZOLADEX

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Gonadotropins.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoledronic Acid

Products Affected

- *zoledronic acid intravenous concentrate*

QL Criteria	1 vial Per 21 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoledronic Acid

Products Affected

- *zoledronic acid intravenous solution*

QL Criteria	1 100 ml bottle Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolinza

Products Affected

- ZOLINZA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days maximum Per 1 fill
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZOLMitriptan

Products Affected

- *zolmitriptan oral*

QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoloft

Products Affected

- ZOLOFT ORAL CONCENTRATE

QL Criteria	10 ml Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoloft

Products Affected

- ZOLOFT ORAL TABLET 100 MG

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoloft

Products Affected

- ZOLOFT ORAL TABLET 25 MG

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zoloft

Products Affected

- ZOLOFT ORAL TABLET 50 MG

QL Criteria	45 tab Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolpidem Tartrate

Products Affected

- *zolpidem tartrate oral tablet 10 mg*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolpidem Tartrate

Products Affected

- *zolpidem tartrate oral tablet 5 mg*

QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolpidem Tartrate

Products Affected

- *zolpidem tartrate sublingual*

QL Criteria	1 tablet Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolpidem Tartrate ER

Products Affected

- *zolpidem tartrate er*

QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zolpimist

Products Affected

- ZOLPIMIST

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 7 days (one week) of the preferred generic alternative zolpidem or zolpidem er.
QL Criteria	1 bottle Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomacton

Products Affected

- ZOMACTON

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zometa

Products Affected

- ZOMETA

QL Criteria	1 bottle Per 7 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomig

Products Affected

- ZOMIG NASAL SOLUTION 2.5 MG

QL Criteria	6 ml Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomig

Products Affected

- ZOMIG NASAL SOLUTION 5 MG

QL Criteria	1 bottle Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomig

Products Affected

- ZOMIG ORAL

QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomig ZMT

Products Affected

- ZOMIG ZMT

QL Criteria	6 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZonaCort 11 Day

Products Affected

- ZONACORT 11 DAY

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented inability to use single tablets or reach appropriate dosing regimen with using single tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of dexamethasone tabs
Notes/References	
Revision Date	Prior Authorization: October 04, 2017 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

ZonaCort 7 Day

Products Affected

- ZONACORT 7 DAY

PA Criteria	Criteria Details
Covered Uses	All FDA approved indications
Exclusion Criteria	
Required Medical Information	A documented inability to use single tablets or reach appropriate dosing regimen with using single tablets
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of dexamethasone tabs
Notes/References	
Revision Date	Prior Authorization: October 04, 2017 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

Zonalon

Products Affected

- ZONALON

QL Criteria	45 grams Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zontivity

Products Affected

- ZONTIVITY

PA Criteria	Criteria Details
Covered Uses	Reduction of the reduction of thrombotic cardiovascular events in patients with a history of myocardial infarction (MI) or with peripheral arterial disease (PAD)
Exclusion Criteria	Do not use in patients with history of stroke, history of transient ischemic attack (TIA), or history of intracranial hemorrhage (ICH), or active pathological bleeding
Required Medical Information	Documented diagnosis or history of myocardial infarction (MI) or peripheral arterial disease (PAD) and concurrent use of aspirin or clopidogrel.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tablet Per 1 Day
Notes/References	Annual Review: 06/2017
Revision Date	Prior Authorization: July 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zorbtive

Products Affected

- ZORBTIVE

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zorvolex

Products Affected

- ZORVOLEX

QL Criteria	3 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zovirax

Products Affected

- ZOVIRAX EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week of generic acyclovir ointment
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zubsolv

Products Affected

- ZUBSOLV SUBLINGUAL TABLET
SUBLINGUAL 0.7-0.18 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	90 tablets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zubsolv

Products Affected

- ZUBSOLV SUBLINGUAL TABLET
SUBLINGUAL 1.4-0.36 MG, 5.7-1.4 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	90 tab Per 30 Days
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zubsolv

Products Affected

- ZUBSOLV SUBLINGUAL TABLET
SUBLINGUAL 11.4-2.9 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zubsolv

Products Affected

- ZUBSOLV SUBLINGUAL TABLET
SUBLINGUAL 2.9-0.71 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zubsolv

Products Affected

- ZUBSOLV SUBLINGUAL TABLET
SUBLINGUAL 8.6-2.1 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of the preferred alternatives, buprenorphine-naloxone sublingual tablet and Suboxone SL film
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 04/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zuplenz

Products Affected

- ZUPLENZ

QL Criteria	12 pack Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zurampic

Products Affected

- ZURAMPIC

PA Criteria	Criteria Details
Covered Uses	Treatment of hyperuricemia associated with gout
Exclusion Criteria	
Required Medical Information	A documented diagnosis of gout, and will be used in combination with a xanthine oxidase inhibitor (allopurinol OR febuxostat)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of allopurinol or febuxostat
QL Criteria	1 tablet Per 1 Day
Notes/References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: October 05, 2017 Quantity Limits: August 25, 2015

Zyban

Products Affected

- ZYBAN

QL Criteria	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyclara

Products Affected

- ZYCLARA

QL Criteria	56 EA Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyclara Pump

Products Affected

- ZYCLARA PUMP EXTERNAL CREAM
2.5 %

QL Criteria	2 bottle Per 365 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyclara Pump

Products Affected

- ZYCLARA PUMP EXTERNAL CREAM
3.75 %

QL Criteria	56 packets Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zydelig

Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyflo

Products Affected

- ZYFLO

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyflo CR

Products Affected

- ZYFLO CR

QL Criteria	4 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zykadia

Products Affected

- ZYKADIA

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	30 days supply Per 1 prescription
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZyPREXA

Products Affected

- ZYPREXA ORAL TABLET 10 MG, 15 MG, 20 MG, 7.5 MG
- ZYPREXA ORAL TABLET 5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZyPREXA

Products Affected

- ZYPREXA ORAL TABLET 2.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	2 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZyPREXA Zydis

Products Affected

- ZYPREXA ZYDIS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of 1 generic (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) Plus Latuda
QL Criteria	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zytiga

Products Affected

- ZYTIGA ORAL TABLET 250 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	4 tablets Per 1 Day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zytiga

Products Affected

- ZYTIGA ORAL TABLET 500 MG

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPPL/Antineoplastics.html
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Refer to the clinical policy bulletin above for details.
Other Criteria	
QL Criteria	2 tablets Per 1 day
Notes/References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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