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) and 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

Abstral

Products Affected

- ABSTRAL

PA Criteria	Criteria Details
Covered Uses	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of fentanyl transmucosal lozenge and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 tablet Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Acamprosate Calcium

Products Affected

- *acamprosate calcium*

QL Criteria	6 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 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

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

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

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

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

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

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

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

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

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

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

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 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 02/2017

Revision Date	Prior Authorization: November 21, 2016 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015

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 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
QL Criteria	1 capsule Per 1 Day
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: November 21, 2016 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015

Acitretin

Products Affected

- *acitretin*

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

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 INTRAVENOUS

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

Actemra

Products Affected

- ACTEMRA SUBCUTANEOUS

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
QL Criteria	1 syringe Per 1 month
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

PA Criteria	Criteria Details
Covered Uses	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone
QL Criteria	120 lozenge Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Activella

Products Affected

- ACTIVELLA

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

Actonel

Products Affected

- ACTONEL ORAL TABLET 150 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	1 tablet Per 28 months
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	1 tab 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	4 tablets Per 28 months
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Actoplus Met

Products Affected

- ACTOPLUS MET

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

Actoplus met XR

Products Affected

- ACTOPLUS MET XR ORAL TABLET EXTENDED RELEASE 24 HOUR 15-1000 MG

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

Actoplus met XR

Products Affected

- ACTOPLUS MET XR ORAL TABLET EXTENDED RELEASE 24 HOUR 30-1000 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

Actos

Products Affected

- ACTOS

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

Aczone

Products Affected

- ACZONE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo and generic dapsone gel
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: November 06, 2017 Quantity Limits: August 25, 2015

Adalat CC

Products Affected

- ADALAT CC ORAL TABLET EXTENDED RELEASE 24 HOUR 30 MG, 90 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

Adalat CC

Products Affected

- ADALAT CC ORAL TABLET EXTENDED RELEASE 24 HOUR 60 MG

QL Criteria	2 tabs Per 1 day
Notes/ References	
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 tabs 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, dexmethylphenidate/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, dexmethylphenidate/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

Adefovir Dipivoxil

Products Affected

- *adefovir dipivoxil*

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

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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Advair Diskus

Products Affected

- ADVAIR DISKUS INHALATION AEROSOL POWDER BREATH ACTIVATED 100-50 MCG/DOSE, 250-50 MCG/DOSE

QL Criteria	1 diskus Per 1 month
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 INHALATION AEROSOL POWDER BREATH ACTIVATED 500-50 MCG/DOSE

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

Advair HFA

Products Affected

- ADVAIR 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

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

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 inections 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, dexmethylphenidate/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

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Asthma
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Asmanex and Qvar
QL Criteria	1 inhaler Per 1 month
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: November 30, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Afeditab CR

Products Affected

- AFEDITAB CR ORAL TABLET EXTENDED RELEASE 24 HOUR 30 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

Afeditab CR

Products Affected

- AFEDITAB CR ORAL TABLET EXTENDED RELEASE 24 HOUR 60 MG

QL Criteria	2 tabs Per 1 day
Notes/ References	
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/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 tab Per 1 day
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/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 tab Per 1 day
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

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/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

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 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 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 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

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*

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

Alendronate Sodium

Products Affected

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

QL Criteria	4 tabs Per 1 month
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 40 mg, 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

Alfuzosin HCl ER

Products Affected

- *alfuzosin hcl 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

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

Allegra Allergy

Products Affected

- ALLEGRA ALLERGY ORAL TABLET
  180 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

Allegra Allergy

Products Affected

- ALLEGRA ALLERGY ORAL TABLET 60 MG

QL Criteria	2 tabs Per 1 day
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 patches Per 28 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	severe diarrhea-predominant irritable bowel syndrome (IBS)
Exclusion Criteria	
Required Medical Information	Patient is female, and has a documented diagnosis of severe diarrhea-predominant irritable bowel syndrome (IBS) including one or more of the following: frequent and severe abdominal pain/discomfort, frequent urgency or fecal incontinence or disability or restriction of daily activities due to IBS, AND patient has chronic IBS symptoms generally lasting 6 months or longer, AND anatomic or biochemical abnormalities of the gastrointestinal tract have been excluded
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step Therapy
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: April 11, 2016 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 tabs 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 tabs 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

Altoprev

Products Affected

- ALTOPREV

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, or simvastatin)
QL Criteria	2 tabs 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

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Asthma
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Asmanex and Qvar
QL Criteria	1 inhaler Per 1 month
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: November 30, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ambien

Products Affected

- AMBIEN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
QL Criteria	2 tabs 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
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

Amcinonide

Products Affected

- *amcinonide external cream*
- *amcinonide external lotion*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of betamethasone dipropionate (cream/ointment/lotion)
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 caps 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*

QL Criteria	2 capsules 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

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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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	1.25 grams Per 1 day
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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	5 grams Per 1 day
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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	5 grams Per 1 fill
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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

Anoro Ellipta

Products Affected

- ANORO ELLIPTA

QL Criteria	1 kit Per 1 fill
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	10 tabs Per 1 fill
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

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

ApexiCon E

Products Affected

- APEXICON E

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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

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

Apriso

Products Affected

- APRISO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), 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

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, dexmethylphenidate/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

Aptiom

Products Affected

- APTIOM ORAL TABLET 200 MG

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	6 tablets Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aptiom

Products Affected

- APTIOM ORAL TABLET 400 MG

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	3 tablets Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aptiom

Products Affected

- APTIOM ORAL TABLET 600 MG

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: October 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Aptiom

Products Affected

- APTIOM ORAL TABLET 800 MG

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	1 tablet Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 19, 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
- ARANESP (ALBUMIN FREE) INJECTION 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

Arava

Products Affected

- ARAVA

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

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	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Serevent
QL Criteria	1 cap 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

Aricept

Products Affected

- ARICEPT

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 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

Arixtra

Products Affected

- ARIXTRA

QL Criteria	2 syringes 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 one month of Asmanex and Qvar
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 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 Breo, Dulera, Symbicort and propionate/salmeterol inhaler (generic Airduo)
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 Breo, Dulera, Symbicort and propionate/salmeterol inhaler (generic Airduo)
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

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

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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	3 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

Asacol HD

Products Affected

- ASACOL HD

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Delzicol, Lialda, or Pentasa
QL Criteria	6 tabs 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

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

Astagraf XL

Products Affected

- ASTAGRAF XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 0.5 MG

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

Astagraf XL

Products Affected

- ASTAGRAF XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 1 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

Atacand

Products Affected

- ATACAND

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
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

Atacand HCT

Products Affected

- ATACAND HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

Atelvia

Products Affected

- ATELVIA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	4 tabs 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

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

Atripla

Products Affected

- ATRIPLA

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

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 tab 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

Avalide

Products Affected

- AVALIDE ORAL TABLET 150-12.5 MG, 300-12.5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

Avandia

Products Affected

- AVANDIA ORAL TABLET 2 MG, 4 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

Avapro

Products Affected

- AVAPRO

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
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

Avita

Products Affected

- *avita external cream*

PA Criteria	Criteria Details
Covered Uses	Acne Vulgaris (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	50 grams Per 1 fill
Notes/ References	

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Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avita

Products Affected

- *avita external gel*

PA Criteria	Criteria Details
Covered Uses	Acne Vulgaris (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	

Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Avodart

Products Affected

- AVODART

ST Criteria	A documented contraindication, intolerance, allergy, or failure of dutasteride or finasteride
QL Criteria	1 cap 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
QL Criteria	1 kit Per 30 Dayss
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
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

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
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

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 amlodipine in combination with two of the following: Atacand, Avapro, Cozaar, Micardis
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

Azulfidine

Products Affected

- AZULFIDINE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Delzicol, Lialda, or Pentasa
QL Criteria	8 tabs 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), Delzicol, Lialda, or Pentasa
QL Criteria	8 tabs 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

PA Criteria	Criteria Details
Covered Uses	Seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy
Exclusion Criteria	
Required Medical Information	A documented diagnosis of seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy AND Concomitant use of an anticonvulsant drug
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	8 tabs Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Baraclude

Products Affected

- BARACLUDE 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

Basaglar KwikPen

Products Affected

- BASAGLAR KWIKPEN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Levemir and Tresiba
Notes/ References	
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: August 25, 2015

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Bebulin

Products Affected

- BEBULIN

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

Beconase AQ

Products Affected

- BECONASE AQ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone and either OTC Nasacort 24HR or Flonase OTC
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

PA Criteria	Criteria Details
Other Criteria	
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 zolpidem, zolpidem er, 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

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

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

Betamethasone Valerate

Products Affected

- *betamethasone valerate external cream*
- *betamethasone valerate external lotion*
- *betamethasone valerate external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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
QL Criteria	15 vials Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bethkis

Products Affected

- BETHKIS

QL Criteria	224 ml Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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 one month each of Anoro Ellipta and 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

Bicalutamide

Products Affected

- *bicalutamide*

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

Bimatoprost

Products Affected

- *bimatoprost ophthalmic*

PA Criteria	Criteria Details
Covered Uses	open-angle glaucoma, ocular hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of glaucoma or ocular hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
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: December 07, 2016 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

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

Bosulif

Products Affected

- BOSULIF 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/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	4 tablets Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bosulif

Products Affected

- BOSULIF 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/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	1 tab Per 1 day
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

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

Breo Ellipta

Products Affected

- BREO ELLIPTA

QL Criteria	2 blisters Per 1 days
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

Brilinta

Products Affected

- BRILINTA

QL Criteria	2 tabs 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 SOLUTION

PA Criteria	Criteria Details
Covered Uses	Partial-onset seizure
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	20 ML Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Briviact

Products Affected

- BRIVIACT ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Partial-onset seizure
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: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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*

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	For ages 5-8 documented inability to use metered dose inhalers
Age Restrictions	Less than 8 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	No prior authorization required for children 1-4 years of age. Medical Exception allowed for topical steroid treatment of eosinophilic esophagitis for which other treatments have been unsatisfactory and for Nasal Polyps when all criteria met: A diagnosis of chronic sinusitis with nasal polyposis, endoscopic sinus surgery has been performed, and standard nasal steroid sprays have been used as part of post-operative management and have failed.
QL Criteria	4 ml Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: January 20, 2017 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

PA Criteria	Criteria Details
Other Criteria	
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	3 tabs 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

BuPROPion HCl

Products Affected

- *bupropion hcl oral*

QL Criteria	6 tabs 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 tabs 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 tablet 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

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

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

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

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

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	2 bottles Per 1 fill
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

PA Criteria	Criteria Details
Other Criteria	
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
QL Criteria	1 pen Per 1 fill
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
QL Criteria	1 pen Per 1 fill
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

PA Criteria	Criteria Details
Covered Uses	Treatment of hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic beta-blockers
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Bystolic

Products Affected

- BYSTOLIC ORAL TABLET 20 MG

PA Criteria	Criteria Details
Covered Uses	Treatment of hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic beta-blockers
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: April 03, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Byvalson

Products Affected

- BYVALSON

PA Criteria	Criteria Details
Covered Uses	Treatment of hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 generic beta-blockers and 2 generic angiotensin receptor blockers (ARBs)
QL Criteria	1 tablet Per 1 day
Notes/ References	Annual Review: 08/2017
Revision Date	Prior Authorization: April 03, 2017 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/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

Calcipotriene

Products Affected

- *calcipotriene external cream*
- *calcipotriene external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of a medium to high potency topical steroid
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 fill
Notes/ References	Annual Review: 06/2016
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Calcitrene

Products Affected

- *calcitrene*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of a medium to high potency topical steroid
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 unit 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*

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

Candesartan Cilexetil-HCTZ

Products Affected

- *candesartan cilexetil-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

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/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

Caprelsa

Products Affected

- CAPRELSA 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/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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Caprelsa

Products Affected

- CAPRELSA ORAL TABLET 300 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	1 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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 LA

Products Affected

- CARDIZEM LA ORAL TABLET EXTENDED RELEASE 24 HOUR 120 MG, 180 MG, 300 MG, 360 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

Cardizem LA

Products Affected

- CARDIZEM LA ORAL TABLET EXTENDED RELEASE 24 HOUR 240 MG

QL Criteria	2 tabs Per 1 day
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 tab Per 1 day
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

Cartia XT

Products Affected

- CARTIA XT ORAL CAPSULE EXTENDED RELEASE 24 HOUR 120 MG, 300 MG
- CARTIA XT ORAL CAPSULE EXTENDED RELEASE 24 HOUR 180 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

Cartia XT

Products Affected

- CARTIA XT ORAL CAPSULE EXTENDED RELEASE 24 HOUR 240 MG

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

Casodex

Products Affected

- CASODEX

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

Cayston

Products Affected

- CAYSTON

QL Criteria	3 vials 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of two generic non steroidal anti-inflammatory drugs
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	
QL Criteria	2 caps Per 1 Day
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	2 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cevimeline HCl

Products Affected

- *cevimeline hcl*

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

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

Chenodal

Products Affected

- CHENODAL

PA Criteria	Criteria Details
Covered Uses	For treatment of cholesterol-type gallstones in patients over 18 years of age and have tried and failed 2 years of generic Actigall (ursodiol) therapy and are not able to undergo surgery due to systemic disease or age, and for treatment of diagnosed Cerebrotendinous Xanthomatosis (CTX) in patients over 18 years of age
Exclusion Criteria	Intrahepatic duct calculus, Chronic constipation in patients with cholesterol gallstones, Prophylaxis of recurrent gallstones, Hyperlipidemia, Rheumatoid Arthritis
Required Medical Information	Prior to initial coverage for gallstone disease, a cholecystogram or other appropriate imaging studies is required to determine presence of radiolucent gallstones, stones that are transparent to x-rays. Due to high risk of hepatotoxicity and adverse effects, for the first 3 months, authorization is required each month pending hepatic function tests (for both gallstones and CTX). After initial 3 months, authorization required every 3 months for length of treatment, pending hepatic function tests. At 6 months prior to authorization, the following results are required, serum cholesterol levels, hepatic function test, and cholecystogram (monitor dissolution of stones). Safety of use beyond a total of 24 months has not been established
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 month (initial authorization), 3 month (reauthorization)
Other Criteria	Max authorization up to 2 years
Notes/ References	

Revision Date	Prior Authorization: April 11, 2016 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

Cialis

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	diagnosis of benign prostatic hyperplasia
Exclusion Criteria	Erectile dysfunction (ED) diagnosis is not covered except for members with ED benefit rider or Fully Insured (FI) members in the state of NY.
Required Medical Information	A documented diagnosis of diagnosis of benign prostatic hyperplasia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year (30 tablets every 30 days)
Other Criteria	Member has failed two alpha blockers (e.g. Cardura (doxazosin), Hytrin (terazosin), Flomax (tamsulosin), Uroxatral (alfuzosin), Rapaflo (silodosin) and failed one 5-alpha reductase inhibitor (e.g. Avodart (dutasteride), Proscar (finasteride), Jalyn (dutasteride/tamsulosin).
QL Criteria	1 tablets Per 1 day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: April 22, 2016 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
QL Criteria	1 kit Per 1 month
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
QL Criteria	1 kit Per 1 month
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
QL Criteria	1 kit Per 1 year
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

Citalopram Hydrobromide

Products Affected

- *citalopram hydrobromide oral tablet*

QL Criteria	1 tabs 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 ORAL CAPSULE 10 MG, 30 MG
- *claravis oral capsule 20 mg, 40 mg*

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	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic oral antibiotic prescribed for treatment of acne (i.e., minocycline, doxycycline)
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

Clarinex

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

Clarinex-D 12 Hour

Products Affected

- CLARINEX-D 12 HOUR

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

Cleocin-T

Products Affected

- CLEOCIN-T EXTERNAL GEL
- CLEOCIN-T EXTERNAL LOTION
- CLEOCIN-T EXTERNAL SOLUTION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
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	4 patches Per 28 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	4 patches Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Clindagel

Products Affected

- CLINDAGEL

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

Clindamycin Phosphate

Products Affected

- *clindamycin phosphate external gel*
- *clindamycin phosphate external lotion*
- *clindamycin phosphate external solution*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
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 gel*
- *clobetasol propionate external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/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*

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 ml 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*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
QL Criteria	236 ml 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 shampoo*

QL Criteria	236 ml 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 solution*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
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 E

Products Affected

- *clobetasol propionate e*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/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 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 EXTERNAL LOTION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
QL Criteria	236 ml 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 EXTERNAL SHAMPOO

QL Criteria	236 ml 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 ml 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 ml Per 30 Days
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*

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

Clopidogrel Bisulfate

Products Affected

- *clopidogrel bisulfate oral*

QL Criteria	1 tab Per 1 fill
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*
- *clozapine oral tablet dispersible 100 mg*

QL Criteria	9 tabs 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*

QL Criteria	4 tabs 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*
- *clozapine oral tablet dispersible 25 mg*

QL Criteria	3 tabs 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 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

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

CloZAPine

Products Affected

- *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

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 tabs 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 tabs 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

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

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

Colazal

Products Affected

- COLAZAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Delzicol, Lialda, or Pentasa
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

Colchicine

Products Affected

- *colchicine oral*

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

Colyte with Flavor Packs

Products Affected

- COLYTE WITH FLAVOR PACKS ORAL SOLUTION RECONSTITUTED 240 GM

QL Criteria	1 bottle Per 1 fill
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 1 month
Notes/ References	
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/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

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	4 capsules Per 1 day
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	3 capsules Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Complera

Products Affected

- COMPLERA

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

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, dexmethylphenidate/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, dexmethylphenidate/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 40 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	
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 roll Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Coreg CR

Products Affected

- COREG CR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of carvedilol
QL Criteria	1 cap Per 1 day
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
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 than or equal to 70 beats per minute, and who are on maximally tolerated doses of beta-blockers (bisoprolol/bisoprolol-HCTZ, carvedilol, carvedilol CR, metoprolol succinate/metoprolol succinate-HCTZ, nevibolol) or have a documented contraindication to beta-blocker use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of a formualry ACE Inhibitor, ACE Inhibitor/HCTZ combination product, Angiotensin-Receptor Blocker, or Angiotensin-Receptor Blocker/HCTZ combination product
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: July 25, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Last update 12/2017
Next update

Cormax Scalp Application

Products Affected

- CORMAX SCALP APPLICATION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
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, dexmethylphenidate/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 100 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Cozaar

Products Affected

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
QL Criteria	2 EA 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 two generic statin medications (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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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 tabs 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	1 capsule 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

Cystagon

Products Affected

- CYSTAGON

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

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/ophthalmic_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 bottles Per 1 month
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, Anoro, Stiolto, Incruse, or Spiriva
QL Criteria	1 tab 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

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: August 25, 2015

Darifenacin Hydrobromide ER

Products Affected

- *darifenacin hydrobromide 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

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, dexmethylphenidate/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

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

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

Desonide

Products Affected

- *desonide external*

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

DesOwen

Products Affected

- DESOWEN EXTERNAL CREAM
- DESOWEN EXTERNAL LOTION

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

Desoximetasone

Products Affected

- *desoximetasone external*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of betamethasone dipropionate (cream/ointment/lotion)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Desoximetasone

Products Affected

- *desoximetasone external*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tabs 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.)
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

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.)
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

Detrol

Products Affected

- DETROL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare and 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 and Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin, oxybutynin XL)
QL Criteria	1 cap Per 1 day
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, dexmethylphenidate/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

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 tabs 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

DiazePAM

Products Affected

- *diazepam rectal*

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

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 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: November 01, 2017 Step Therapy: August 25, 2015 Quantity Limits: October 10, 2017

Diclofenac Sodium

Products Affected

- *diclofenac sodium transdermal gel 1 %*

QL Criteria	200 GM 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 %
- DIFFERIN EXTERNAL LOTION

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
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 tabs Per 1 fill
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diflorasone Diacetate

Products Affected

- *diflorasone diacetate external*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Diflorasone Diacetate

Products Affected

- *diflorasone diacetate external*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of betamethasone dipropionate (cream/ointment/lotion)
Notes/ References	
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 of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan
QL Criteria	8 ml Per 30 Days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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

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

Diltiazem CD

Products Affected

- *diltiazem cd oral capsule extended release 24 hour 120 mg, 180 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

Diltiazem CD

Products Affected

- *diltiazem cd oral capsule extended release 24 hour 240 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

DiltiaZEM CD

Products Affected

- *diltiazem cd oral capsule extended release 24 hour 300 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

Diltiazem HCl ER

Products Affected

- *diltiazem hcl er oral capsule extended release 24 hour 240 mg*

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

Diltiazem HCl ER Beads

Products Affected

- *diltiazem hcl er beads oral capsule extended release 24 hour 120 mg, 180 mg, 300 mg, 360 mg*

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

Diltiazem HCl ER Beads

Products Affected

- *diltiazem hcl er beads oral capsule extended release 24 hour 240 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

Diltiazem HCl ER Coated Beads

Products Affected

- *diltiazem hcl er coated beads oral capsule extended release 24 hour 120 mg, 180 mg*
- *diltiazem hcl er coated beads oral capsule extended release 24 hour 360 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

Diltiazem HCl ER Coated Beads

Products Affected

- *diltiazem hcl er coated beads oral tablet extended release 24 hour 180 mg, 300 mg, 360 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

Diltiazem HCl ER Coated Beads

Products Affected

- *diltiazem hcl er coated beads oral tablet extended release 24 hour 240 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

DilTIAZem HCl ER Coated Beads

Products Affected

- *diltiazem hcl er coated beads oral capsule extended release 24 hour 240 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

DilTIAZem HCl ER Coated Beads

Products Affected

- *diltiazem hcl er coated beads oral capsule extended release 24 hour 300 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

Dilt-XR

Products Affected

- *dilt-xr oral capsule extended release 24 hour 240 mg*

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

Diovan

Products Affected

- DIOVAN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, 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

Diovan HCT

Products Affected

- DIOVAN HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

Dipentum

Products Affected

- DIPENTUM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), 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 ORAL TABLET EXTENDED RELEASE 24 HOUR 10 MG, 15 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare and Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin, oxybutynin XL)
QL Criteria	2 tabs Per 1 Day
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 ORAL TABLET EXTENDED RELEASE 24 HOUR 5 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Vesicare and Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin, oxybutynin XL)
QL Criteria	1 tabs Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dolophine

Products Affected

- DOLOPHINE

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

Donepezil HCl

Products Affected

- *donepezil hcl*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Dovonex

Products Affected

- DOVONEX EXTERNAL CREAM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of a medium to high potency topical steroid
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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

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 caps 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

Duetact

Products Affected

- DUETACT

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

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 tabs 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*
- *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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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

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	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

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	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

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	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

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	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

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	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

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 04, 2017 Quantity Limits: August 25, 2015

Dyanavel XR

Products Affected

- DYANAVEL 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, dexmethylphenidate/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

- DYSPORT

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

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
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

Edarbyclor

Products Affected

- EDARBYCLOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

Edurant

Products Affected

- EDURANT

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

QL Criteria	1 cap 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

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

Elestrin

Products Affected

- ELESTRIN

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

Elocon

Products Affected

- ELOCON EXTERNAL CREAM
- ELOCON EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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

PA Criteria	Criteria Details
Other Criteria	
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

Emend

Products Affected

- EMEND ORAL CAPSULE 125 MG, 80 MG
- EMEND ORAL CAPSULE 40 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

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

Emtriva

Products Affected

- EMTRIVA ORAL CAPSULE

QL Criteria	1 cap Per 1 day
Notes/ References	
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 and Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin, oxybutynin 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

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 injections Per 1 month
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

Enoxaparin Sodium

Products Affected

- *enoxaparin sodium*

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

Enstilar

Products Affected

- ENSTILAR

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

Entecavir

Products Affected

- *entecavir*

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

Entresto

Products Affected

- ENTRESTO

PA Criteria	Criteria Details
Covered Uses	Heart Failure
Exclusion Criteria	Known or suspected pregnancy
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: April 11, 2016 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

QL Criteria	1 bottle 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

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/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

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 tab 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 cap 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 CAPSULE

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	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/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	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 tablet 10 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

Escitalopram Oxalate

Products Affected

- *escitalopram oxalate oral tablet 20 mg, 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

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

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

Estradiol-Norethindrone Acet

Products Affected

- *estradiol-norethindrone acet*

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

Estradiol-Norethindrone Acet

Products Affected

- *estradiol-norethindrone acet*

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

Estrogel

Products Affected

- ESTROGEL

QL Criteria	50 grams 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 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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 month
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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	120 tabs 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

Evoxac

Products Affected

- EVOXAC

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

Exalgo

Products Affected

- EXALGO 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	

PA Criteria	Criteria Details
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 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

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

Exelon

Products Affected

- EXELON TRANSDERMAL

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Exforge

Products Affected

- EXFORGE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of amlodipine in combination with two of the following: Atacand, Avapro, Cozaar, 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

Exforge HCT

Products Affected

- EXFORGE HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of amlodipine in combination with two of the following: Atacand HCT, Avalide, Hyzaar, Micardis 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
QL Criteria	15 vials Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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

Fabior

Products Affected

- FABIOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
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*

QL Criteria	21 tabs Per 1 fill
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 tabs Per 1 fill
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 one generic medication (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) plus Latuda
QL Criteria	2 tabs 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 one generic medication (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

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/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	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/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

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 tabs 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 tabs 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 tabs 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	3 tabs 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

Felodipine ER

Products Affected

- *felodipine 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

Femring

Products Affected

- FEMRING

QL Criteria	1 ring Per 90 dayss
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fenofibrate

Products Affected

- *fenofibrate oral capsule*

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 tablet 145 mg, 160 mg, 48 mg, 54 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

Fenofibrate Micronized

Products Affected

- *fenofibrate micronized*

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

Fenofibric Acid

Products Affected

- *fenofibric acid*

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

Fenofibric Acid

Products Affected

- *fenofibric acid*

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	(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	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	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone
QL Criteria	120 lozenge Per 30 Days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 10, 2016 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 fentanyl transmucosal lozenge and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	120 tablet 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/Antidotes.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 cap Per 1 Day
Notes/ References	Annual Review: 05/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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

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 cap 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

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

Fibricor

Products Affected

- FIBRICOR

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

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

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

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
QL Criteria	3 syringes Per 30 dayss
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

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

Flovent Diskus

Products Affected

- FLOVENT DISKUS

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

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

Fluocinolone Acetonide

Products Affected

- *fluocinolone acetonide external cream*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluocinolone Acetonide

Products Affected

- *fluocinolone acetonide external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Fluocinonide

Products Affected

- *fluocinonide external cream*
- *fluocinonide external gel*
- *fluocinonide external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of betamethasone dipropionate (cream/ointment/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

Fluocinonide

Products Affected

- *fluocinonide external solution*

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

FLUoxetine HCl

Products Affected

- *fluoxetine hcl oral capsule 10 mg*

QL Criteria	1 cap 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	4 caps Per 1 month
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*

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 tabs 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 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

Fluticasone Propionate

Products Affected

- *fluticasone propionate external cream*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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 tabs 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, dexmethylphenidate/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, dexmethylphenidate/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

Fondaparinux Sodium

Products Affected

- *fondaparinux sodium*

QL Criteria	2 syringes 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

Fosamax

Products Affected

- FOSAMAX ORAL TABLET 70 MG

QL Criteria	4 tablets Per 1 month
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	4 tabs 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

Fragmin

Products Affected

- FRAGMIN SUBCUTANEOUS SOLUTION 10000 UNIT/ML, 12500 UNIT/0.5ML, 15000 UNIT/0.6ML, 18000 UNT/0.72ML, 2500 UNIT/0.2ML, 5000 UNIT/0.2ML, 7500 UNIT/0.3ML, 95000 UNIT/3.8ML

QL Criteria	2 syringes Per 1 Day
Notes/ References	
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 Precision Neo Test

Products Affected

- FREESTYLE PRECISION NEO 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

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

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

PA Criteria	Criteria Details
Covered Uses	A diagnosis of partial-onset seizures OR generalized tonic-clonic seizures
Exclusion Criteria	
Required Medical Information	A documented diagnosis of partial-onset seizures or generalized tonic-clonic seizures, and documented use as adjunct therapy with one or more other FDA approved Anti-Epileptic Drug (AED).
Age Restrictions	12 years and greater
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: April 27, 2017 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 tablet*

QL Criteria	6 tabs 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 12 MG
- GABITRIL ORAL TABLET 4 MG

QL Criteria	4 tabs 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 16 MG

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

Galantamine Hydrobromide

Products Affected

- *galantamine hydrobromide*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Galantamine Hydrobromide ER

Products Affected

- *galantamine hydrobromide er*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 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

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 30 fillss
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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 and 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

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

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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and 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	Ulcerative colitis
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild to moderate ulcerative colitis in males. Note: Per Product Labeling, Giazo effectiveness was not demonstrated in female patients.
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 mesalamine DR (generic Asacol HD), Delzicol, Lialda, or Pentasa
QL Criteria	6 tabs Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: April 11, 2016 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 cap 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	1 tab Per 1 day
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

GlucaGen HypoKit

Products Affected

- GLUCAGEN HYPOKIT

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

Glyxambi

Products Affected

- GLYXAMBI

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Invokana/Invokamet and either Januvia/Janumet and either Tradjenta/Jentadueto
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

Gralise

Products Affected

- GRALISE ORAL TABLET 300 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of gabapentin
QL Criteria	1 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 tabs 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 1 fill
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*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
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	
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

Hepsera

Products Affected

- HEPSERA

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

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	
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 300 MG

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.
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: February 10, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Horizant

Products Affected

- HORIZANT ORAL TABLET EXTENDED RELEASE 600 MG

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.
QL Criteria	1 tablet Per 2 Days
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, 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

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 inections 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 inections 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 inections 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 inections 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 inections 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 inections 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/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

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	

PA Criteria	Criteria Details
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

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	

PA Criteria	Criteria Details
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

Hydrocodone-Ibuprofen

Products Affected

- *hydrocodone-ibuprofen oral tablet 10-200 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

Hydrocodone-Ibuprofen

Products Affected

- *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

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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

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

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 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

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PA Criteria	Criteria Details
Other Criteria	
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

Hyzaar

Products Affected

- HYZAAR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/hctz
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ibandronate Sodium

Products Affected

- *ibandronate sodium oral*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	1 tab Per 1 month
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/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	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 ORAL TABLET 10-200 MG
- *ibudone oral tablet 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

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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

Iclusig

Products Affected

- ICLUSIG ORAL TABLET 15 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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Iclusig

Products Affected

- ICLUSIG ORAL TABLET 45 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	1 tab Per 1 day
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/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

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/Idhifa.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/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	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/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 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	4 capsules Per 1 Day
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	8 vials Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Imitrex STATdose Refill

Products Affected

- IMITREX STATDOSE REFILL SUBCUTANEOUS SOLUTION CARTRIDGE

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

Imitrex STATdose System

Products Affected

- IMITREX STATDOSE SYSTEM SUBCUTANEOUS SOLUTION AUTO-INJECTOR 6 MG/0.5ML

QL Criteria	2 boxes Per 1 fill
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 Leishmania donovani, Cutaneous leishmaniasis due to Leishmania braziliensis, Leishmania guyanensis, and Leishmania panamensis, or Mucosal leishmaniasis due to Leishmania braziliensis
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	28 days
Other Criteria	
QL Criteria	3 capsules Per 1 day
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 XL

Products Affected

- INDERAL XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 80 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

Indomethacin

Products Affected

- *indomethacin oral*

QL Criteria	3 capsule Per 1 Day
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: August 25, 2015

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/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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

InnoPran XL

Products Affected

- INNOPRAN XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 120 MG

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

InnoPran XL

Products Affected

- INNOPRAN XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 80 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

Intelence

Products Affected

- INTELENCE ORAL TABLET 100 MG, 25 MG

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

Intelence

Products Affected

- INTELENCE ORAL TABLET 200 MG

QL Criteria	2 tabs 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

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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine, or Vyvanse
QL Criteria	1 tab 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	2 tabs 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and 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	2 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 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ipratropium Bromide

Products Affected

- *ipratropium bromide nasal*

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

Iprivask

Products Affected

- IPRIVASK

QL Criteria	2 syringes 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-Hydrochlorothiazide

Products Affected

- *irbesartan-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

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/Iressa.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: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Isentress

Products Affected

- ISENTRESS ORAL TABLET

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

Isentress

Products Affected

- ISENTRESS ORAL TABLET CHEWABLE

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

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

Itraconazole

Products Affected

- *itraconazole oral*

QL Criteria	4 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

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 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/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

Jakafi

Products Affected

- JAKAFI ORAL TABLET 15 MG, 20 MG, 25 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/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 tabs 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 tabs 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 tab 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 tabs 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 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Jentadueto

Products Affected

- JENTADUETO

QL Criteria	2 tabs 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

Jentadueto 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

Jentadueto 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/ophthalmic_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

Jublia

Products Affected

- JUBLIA

PA Criteria	Criteria Details
Covered Uses	Onychomycosis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of onychomycosis confirmed by either a positive KOH stain (potassium hydroxide), positive PAS stain (para-aminosalicylic acid), a positive DTM (dermatophyte test medium) or positive fungal culture (NOTE: This positive test should be within the last 3 - 6 months and associated with the current infection)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Failure of an adequate trial of one systemic oral alternative is terbinafine (6 weeks for fingernail infections, 12 weeks for toenail infections), griseofulvin (6 months), itraconazole (60 days (PulsePak) for fingernail infections, 90 days for toenail), OR If member has hepatic dysfunction or increased risk for liver disease (for example, has a history of alcohol abuse or a history of hepatitis), or is female and is pregnant and/or breastfeeding. (No trial needed)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one systemic (oral) alternative such as terbinafine, itraconazole, or griseofulvin
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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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 tab Per 1 day
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

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	3 capsules Per 1 Day
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 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
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

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	

PA Criteria	Criteria Details
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 tablets 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

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 tabs 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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	4 tabs 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 Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
QL Criteria	2 EA 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 tabs 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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Kerydin

Products Affected

- KERYDIN

PA Criteria	Criteria Details
Covered Uses	Onychomycosis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of onychomycosis confirmed by either a positive KOH stain (potassium hydroxide), positive PAS stain (para-aminosalicylic acid), a positive DTM (dermatophyte test medium) or positive fungal culture (NOTE: This positive test should be within the last 3 - 6 months and associated with the current infection)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Failure of an adequate trial of one systemic oral alternative is terbinafine (6 weeks for fingernail infections, 12 weeks for toenail infections), griseofulvin (6 months), itraconazole (60 days (PulsePak) for fingernail infections, 90 days for toenail), OR If member has hepatic dysfunction or increased risk for liver disease (for example, has a history of alcohol abuse or a history of hepatitis), or is female and is pregnant and/or breastfeeding. (No trial needed)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one systemic (oral) alternative such as terbinafine, itraconazole, or griseofulvin
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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Ketoconazole

Products Affected

- *ketoconazole oral*

QL Criteria	2 tabs Per 1 day
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	4 tablets Per 1 Day
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/rxnonmedicare/data/2017/MISC/carbonic_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 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

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
QL Criteria	1 syringe Per 1 Day
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
QL Criteria	2 tabs 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
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

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/antidiabetic 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 tabs 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

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 syringe Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET DISPERSIBLE 100 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 lamotrigine or lamotrigine er
QL Criteria	2 tabs Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET DISPERSIBLE 200 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 lamotrigine or lamotrigine er
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET DISPERSIBLE 25 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 lamotrigine or lamotrigine er
QL Criteria	6 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal ODT

Products Affected

- LAMICTAL ODT ORAL TABLET DISPERSIBLE 50 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 lamotrigine or lamotrigine er
QL Criteria	3 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL KIT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of immediate release lamotrigine
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

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET EXTENDED RELEASE 24 HOUR 200 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	3 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET EXTENDED RELEASE 24 HOUR 250 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LaMICtal XR

Products Affected

- LAMICTAL XR ORAL TABLET EXTENDED RELEASE 24 HOUR 300 MG

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIgine

Products Affected

- *lamotrigine oral tablet dispersible 100 mg, 200 mg*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIgine

Products Affected

- *lamotrigine oral tablet dispersible 25 mg*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	6 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIgine

Products Affected

- *lamotrigine oral tablet dispersible 50 mg*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	3 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 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*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

LamoTRIgine ER

Products Affected

- *lamotrigine er oral tablet extended release 24 hour 200 mg*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	3 tabs Per 1 day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 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*

PA Criteria	Criteria Details
Covered Uses	Epilepsy, Bipolar I Disorder (only for ODT formulation)
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Epilepsy or Bipolar I disorder (Only ODT formulation will be reviewed for Bipolar I).
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 immediate release lamotrigine
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lansoprazole

Products Affected

- *lansoprazole oral capsule delayed release*

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

Lantus

Products Affected

- LANTUS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Levemir and Tresiba
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lantus SoloStar

Products Affected

- LANTUS SOLOSTAR SUBCUTANEOUS SOLUTION PEN-INJECTOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Levemir and Tresiba
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, 40 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, and clozapine
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 20 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, and clozapine
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 60 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, and clozapine
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, and clozapine
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

Lazanda

Products Affected

- LAZANDA NASAL SOLUTION 100 MCG/ACT, 400 MCG/ACT

PA Criteria	Criteria Details
Covered Uses	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of fentanyl transmucosal lozenge and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	15 bottles Per 1 fill
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lazanda

Products Affected

- LAZANDA NASAL SOLUTION 300 MCG/ACT

PA Criteria	Criteria Details
Covered Uses	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of fentanyl transmucosal lozenge and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone)
QL Criteria	4 botttles Per 30 days
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: October 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Leflunomide

Products Affected

- *leflunomide 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

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	5 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 day supply Per 1 fill
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 day supply Per 1 fill
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 day supply Per 1 Day
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/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 day supply Per 1 fill
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/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 day supply Per 1 fill
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/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 day supply Per 1 Day
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, or simvastatin)
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

Lescol XL

Products Affected

- LESCOL XL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (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

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 tabs 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 tabs 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	(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

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 10 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

Lexapro

Products Affected

- LEXAPRO ORAL TABLET 20 MG
- LEXAPRO ORAL TABLET 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

Lialda

Products Affected

- LIALDA

QL Criteria	4 tabs Per 1 Day
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), of 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	*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
QL Criteria	50 GM 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	
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

Lidocaine PAK

Products Affected

- *lidocaine pak*

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

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 GM 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

Linezolid

Products Affected

- *linezolid oral suspension reconstituted*

QL Criteria	150 ml Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Linezolid

Products Affected

- *linezolid oral tablet*

QL Criteria	28 tablets Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Linzess

Products Affected

- LINZESS

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

Linzess

Products Affected

- LINZESS

QL Criteria	1 capsules 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (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

Lipofen

Products Affected

- LIPOFEN

QL Criteria	1 cap 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 two generic statin medications (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

Lofibra

Products Affected

- LOFIBRA ORAL CAPSULE 134 MG, 67 MG

QL Criteria	1 cap 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 tab 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/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	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

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

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

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

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

Losartan Potassium

Products Affected

- *losartan potassium oral tablet 25 mg, 50 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

Lotronex

Products Affected

- LOTRONEX

PA Criteria	Criteria Details
Covered Uses	severe diarrhea-predominant irritable bowel syndrome (IBS)
Exclusion Criteria	
Required Medical Information	Patient is female, and has a documented diagnosis of severe diarrhea-predominant irritable bowel syndrome (IBS) including one or more of the following: frequent and severe abdominal pain/discomfort, frequent urgency or fecal incontinence or disability or restriction of daily activities due to IBS, AND patient has chronic IBS symptoms generally lasting 6 months or longer, AND anatomic or biochemical abnormalities of the gastrointestinal tract have been excluded
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step Therapy
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: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Lovastatin

Products Affected

- *lovastatin*

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

Lovenox

Products Affected

- LOVENOX

QL Criteria	2 syringes 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/ophthalmic_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

Lumigan

Products Affected

- LUMIGAN OPHTHALMIC SOLUTION
  0.01 %

PA Criteria	Criteria Details
Covered Uses	open-angle glaucoma, ocular hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of glaucoma or ocular hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
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: December 07, 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
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

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

Lynparza

Products Affected

- LYNPARZA

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

QL Criteria	30 tabs Per 1 fill
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/ophthalmic_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	
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Maprotiline HCl

Products Affected

- *maprotiline hcl oral tablet 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

Maprotiline HCl

Products Affected

- *maprotiline hcl oral tablet 50 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

Maprotiline HCl

Products Affected

- *maprotiline hcl oral tablet 75 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

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 caps 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

Matzim LA

Products Affected

- MATZIM LA ORAL TABLET EXTENDED RELEASE 24 HOUR 180 MG, 300 MG, 360 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

Matzim LA

Products Affected

- MATZIM LA ORAL TABLET EXTENDED RELEASE 24 HOUR 240 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

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	12 tabs 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	12 tablets 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 ORAL TABLET 0.5 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	3 tabs Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Mekinist

Products Affected

- MEKINIST ORAL TABLET 2 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	1 tab Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Memantine HCl

Products Affected

- *memantine hcl oral tablet*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Menostar

Products Affected

- MENOSTAR

QL Criteria	4 patches Per 28 fills
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

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

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 tabs 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

Methadone HCl

Products Affected

- *methadone hcl oral concentrate*

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. (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).

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	(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).

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 solution 5 mg/5ml*

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. (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).

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 tablet*
- *methadone hcl oral tablet soluble*

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. (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).

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	(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).

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
- METHADOSE ORAL TABLET SOLUBLE

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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Last update 12/2017
Next update

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

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

Methamphetamine HCl

Products Affected

- *methamphetamine hcl*

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 tabs 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

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, dexmethylphenidate/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, dexmethylphenidate/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 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 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 cap 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)*

QL Criteria	1 cap 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)*

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 Succinate ER

Products Affected

- *metoprolol succinate er oral tablet extended release 24 hour 100 mg, 50 mg*

QL Criteria	1.5 tabs 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 tabs 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 tab 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 tabs 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 NASAL

ST Criteria	http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bone_disease_agents.html
QL Criteria	1 bottle Per 1 fill
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred generic alternatives from the following agents: candesartan, eprosartan, irbesartan, losartan, valsartan, olmesartan, or telmisartan
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

Micardis HCT

Products Affected

- MICARDIS HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any two preferred alternatives from the following: candesartan/hctz, eprosartan/hctz, irbesartan/hctz, losartan/hctz, telmisartan/hctz, olmesartan/hctz, or valsartan/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

Mimvey

Products Affected

- MIMVEY

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

Minivelle

Products Affected

- MINIVELLE

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

Mirapex ER

Products Affected

- MIRAPEX 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

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*

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

Mirtazapine

Products Affected

- *mirtazapine 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

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 tabs 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

Mometasone Furoate

Products Affected

- *mometasone furoate external cream*
- *mometasone furoate external ointment*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Monoclate-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

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

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

Morphine Sulfate

Products Affected

- *morphine sulfate oral solution 10 mg/5ml*

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

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

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

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	
QL Criteria	4 tablet 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	
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

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

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

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

Multaq

Products Affected

- MULTAQ

QL Criteria	2 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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, dexmethylphenidate/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

Myorisan

Products Affected

- *myorisan oral capsule 10 mg, 20 mg, 40 mg*
- MYORISAN ORAL CAPSULE 30 MG

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	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic oral antibiotic prescribed for treatment of acne (i.e., minocycline, doxycycline)
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

Myrbetriq

Products Affected

- MYRBETRIQ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one preferred generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin)
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

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 tablet 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

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

Namenda

Products Affected

- NAMENDA ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Namenda Titration Pak

Products Affected

- NAMENDA TITRATION PAK

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Namenda XR

Products Affected

- NAMENDA XR

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Namenda XR Titration Pack

Products Affected

- NAMENDA XR TITRATION PACK

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Namzaric

Products Affected

- NAMZARIC

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 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

Nasonex

Products Affected

- NASONEX

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone and either OTC Nasacort 24HR or Flonase OTC
Notes/ References	
Revision Date	Prior Authorization: August 25, 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

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/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	
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

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 Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
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

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

PA Criteria	Criteria Details
Covered Uses	Moderate to severe restless leg syndrome, Parkinson's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of moderate to severe restless leg syndrome or Parkinson's Disease
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	1 patch Per 1 day
Notes/ References	
Revision Date	Prior Authorization: April 22, 2016 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 tabs Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nevirapine ER

Products Affected

- *nevirapine er oral tablet extended release 24 hour 100 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

Nevirapine ER

Products Affected

- *nevirapine er oral tablet extended release 24 hour 400 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

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/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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

NexIUM

Products Affected

- NEXIUM ORAL CAPSULE DELAYED RELEASE 40 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
QL Criteria	1 cap Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015

NexIUM

Products Affected

- NEXIUM ORAL PACKET

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
QL Criteria	1 packet Per 1 Day
Notes/ References	Annual Review: 02/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015

NexIUM 24HR

Products Affected

- NEXIUM 24HR

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

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 gum*

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

Nicotine Polacrilex

Products Affected

- *nicotine polacrilex mouth/throat lozenge*

QL Criteria	20 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

Nifediac CC

Products Affected

- NIFEDIAC CC ORAL TABLET EXTENDED RELEASE 24 HOUR 30 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

Nifedical XL

Products Affected

- NIFEDICAL XL ORAL TABLET EXTENDED RELEASE 24 HOUR 60 MG

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

NIFEdipine ER

Products Affected

- *nifedipine er oral tablet extended release 24 hour 30 mg, 90 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

NIFEdipine ER

Products Affected

- *nifedipine er oral tablet extended release 24 hour 60 mg*

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

NIFEdipine ER Osmotic Release

Products Affected

- *nifedipine er osmotic release oral tablet extended release 24 hour 30 mg, 90 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

NIFEdipine ER Osmotic Release

Products Affected

- *nifedipine er osmotic release oral tablet extended release 24 hour 60 mg*

QL Criteria	2 tabs Per 1 day
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/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	3 capsules Per 28 days
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Nisoldipine ER

Products Affected

- *nisoldipine er oral tablet extended release 24 hour 17 mg, 20 mg, 34 mg, 40 mg, 8.5 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

Nisoldipine ER

Products Affected

- *nisoldipine er oral tablet extended release 24 hour 30 mg*

QL Criteria	2 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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

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

Norditropin FlexPro

Products Affected

- NORDITROPIN FLEXPRO

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

Northera

Products Affected

- NORTHERA ORAL CAPSULE 100 MG, 200 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 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

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

Noxafil

Products Affected

- NOXAFIL ORAL TABLET DELAYED RELEASE

QL Criteria	93 tablets 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

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/Neumega.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

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

PA Criteria	Criteria Details
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of two of the following: morphine, 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

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

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

PA Criteria	Criteria Details
Other Criteria	
ST Criteria	FOR A DIAGNOSIS OF PAIN: A documented contraindication, intolerance, allergy, or failure of two of Butrans, Hysingla ER, or Oxycontin. FOR A DIAGNOSIS OF DIABETIC PERIPHERAL NEUROPATHY: A documented contraindication, intolerance, allergy, or failure of Cymbalta and Lyrica.
QL Criteria	2 tabs 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

Nuedexta

Products Affected

- NUEDEXTA

QL Criteria	2 caps 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, 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
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 tablets Per 1 Day

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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/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

Nymalize

Products Affected

- NYMALIZE ORAL SOLUTION 60 MG/20ML

QL Criteria	135.2 ml Per 1 Day
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_Cholagitis.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_Cholagitis.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

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/Odomzo.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: August 25, 2015 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 15 mg, 20 mg, 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

OLANZapine

Products Affected

- *olanzapine oral tablet 2.5 mg*

QL Criteria	2 tabs 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*

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

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

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 tabs Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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 and either OTC Nasacort 24HR or Flonase OTC
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Omnitrope

Products Affected

- OMNITROPE

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

OneTouch Ultra Blue

Products Affected

- ONETOUCH ULTRA BLUE

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

OneTouch Verio

Products Affected

- ONETOUCH VERIO IN VITRO STRIP

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

Onfi

Products Affected

- ONFI ORAL SUSPENSION

PA Criteria	Criteria Details
Covered Uses	Seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy
Exclusion Criteria	
Required Medical Information	A documented diagnosis of seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy AND Concomitant use of an anticonvulsant drug
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Onfi

Products Affected

- ONFI ORAL TABLET 10 MG, 20 MG

PA Criteria	Criteria Details
Covered Uses	Seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy
Exclusion Criteria	
Required Medical Information	A documented diagnosis of seizures associated with Lennox-Gastaut syndrome or refractory(therapy resistant) epilepsy AND Concomitant use of an anticonvulsant drug
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Onglyza

Products Affected

- ONGLYZA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
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

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

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

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	(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	
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

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	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oravig

Products Affected

- ORAVIG

QL Criteria	14 tabs Per 1 fill
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

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
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 125 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/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 28 Days
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/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

Orenitram

Products Affected

- ORENITRAM

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 Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin
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

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/rxnonmedicare/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

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

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

Oxtellar XR

Products Affected

- OXTELLAR XR ORAL TABLET EXTENDED RELEASE 24 HOUR 150 MG, 300 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of oxcarbazepine
QL Criteria	2 tabs 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of oxcarbazepine
QL Criteria	4 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Oxybutynin Chloride

Products Affected

- *oxybutynin chloride oral tablet*

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

Oxybutynin Chloride ER

Products Affected

- *oxybutynin chloride er oral tablet extended release 24 hour 10 mg, 15 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

Oxybutynin Chloride ER

Products Affected

- *oxybutynin chloride er oral tablet extended release 24 hour 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

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

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

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

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 HCl ER

Products Affected

- *oxycodone hcl er 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	
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	4 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

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

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

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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

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

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-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

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

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	4 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

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

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

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

PA Criteria	Criteria Details
Other Criteria	
QL Criteria	4 tabs Per 1 day
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 tablets 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 tablets 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 tabs 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 tabs 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 ML 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 tabs 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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

PEG 3350/Electrolytes

Products Affected

- *peg 3350/electrolytes*

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

PEG-3350/Electrolytes

Products Affected

- *peg-3350/electrolytes*

QL Criteria	1 bottle Per 1 fill
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

Penlac

Products Affected

- PENLAC

PA Criteria	Criteria Details
Covered Uses	Onychomycosis
Exclusion Criteria	
Required Medical Information	A documented diagnosis of onychomycosis confirmed by either a positive KOH stain (potassium hydroxide), positive PAS stain (para-aminosalicylic acid), a positive DTM (dermatophyte test medium) or positive fungal culture (NOTE: This positive test should be within the last 3 - 6 months and associated with the current infection)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Failure of an adequate trial of one systemic oral alternative is terbinafine (6 weeks for fingernail infections, 12 weeks for toenail infections), griseofulvin (6 months), itraconazole (60 days (PulsePak) for fingernail infections, 90 days for toenail), OR If member has hepatic dysfunction or increased risk for liver disease (for example, has a history of alcohol abuse or a history of hepatitis), or is female and is pregnant and/or breastfeeding. (No trial needed)
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one systemic (oral) alternative such as terbinafine, itraconazole, or griseofulvin
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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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

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

Percocet

Products Affected

- PERCOCET ORAL TABLET 10-325 MG, 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

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

Percocet

Products Affected

- PERCOCET ORAL TABLET 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	
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

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

Pioglitazone HCl

Products Affected

- *pioglitazone hcl*

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

Pioglitazone HCl-Glimepiride

Products Affected

- *pioglitazone hcl-glimepiride*

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

Pioglitazone HCl-Metformin HCl

Products Affected

- *pioglitazone hcl-metformin hcl*

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

Plavix

Products Affected

- PLAVIX

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

Plavix

Products Affected

- PLAVIX

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

Plegridy

Products Affected

- PLEGRIDY 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/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 syringes Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Plegridy

Products Affected

- PLEGRIDY 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	
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 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/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 kit Per 365 Days
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
  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	
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

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/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 cap Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pradaxa

Products Affected

- PRADAXA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Eliquis and Xarelto
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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/PCSK9.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/PCSK9.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*

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, or 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

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 strips 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 strips 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 strips 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 strips 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 strips 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 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prenate Mini

Products Affected

- PRENATE MINI ORAL CAPSULE 18-0.6-0.4-350 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

Prevacid

Products Affected

- PREVACID ORAL CAPSULE DELAYED RELEASE 30 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

Prevacid 24HR

Products Affected

- PREVACID 24HR

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

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 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 02/2017

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Last update 12/2017
Next update

Revision Date	Prior Authorization: November 21, 2016 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015

Prezista

Products Affected

- PREZISTA ORAL SUSPENSION

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

Prezista

Products Affected

- PREZISTA ORAL TABLET 150 MG, 600 MG, 75 MG

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

Prezista

Products Affected

- PREZISTA ORAL TABLET 800 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

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

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Last update 12/2017
Next update

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

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

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Last update 12/2017
Next update

Revision Date	Prior Authorization: August 25, 2015 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 tablet Per 1 Day
Notes/ References	Annual Review: 05/2017

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Last update 12/2017
Next update

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

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

Procardia XL

Products Affected

- PROCARDIA XL ORAL TABLET EXTENDED RELEASE 24 HOUR 30 MG, 90 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

Procardia XL

Products Affected

- PROCARDIA XL ORAL TABLET EXTENDED RELEASE 24 HOUR 60 MG

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

ProCentra

Products Affected

- PROCENTRA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, dexmethylphenidate/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: August 25, 2015 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

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	240 caps Per 30 monthss
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	750 caps Per 30 monthss
Notes/ References	
Revision Date	Prior Authorization: January 11, 2017 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	
QL Criteria	1 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Prometrium

Products Affected

- PROMETRIUM

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

Propafenone HCl ER

Products Affected

- *propafenone hcl er*

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

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	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of dutasteride or finasteride
Notes/ References	
Revision Date	Prior Authorization: October 25, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Protonix

Products Affected

- PROTONIX ORAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 generic RX or OTC proton pump inhibitors (i.e. esomeprazole mag, lansoprazole, omeprazole, pantoprazole, rabeprazole) (not required for Nexium requests for members under one year of age)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: October 06, 2017 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	

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: October 11, 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 one 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 tabs Per 1 day

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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 cap 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

Psorcon

Products Affected

- *psorcon*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pulmicort

Products Affected

- PULMICORT

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	For ages 5-8 documented inability to use metered dose inhalers
Age Restrictions	Less than 8 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	No prior authorization required for children 1-4 years of age. Medical Exception allowed for topical steroid treatment of eosinophilic esophagitis for which other treatments have been unsatisfactory and for Nasal Polyps when all criteria met: A diagnosis of chronic sinusitis with nasal polyposis, endoscopic sinus surgery has been performed, and standard nasal steroid sprays have been used as part of post-operative management and have failed.
QL Criteria	4 ml Per 1 Day
Notes/ References	Annual Review: 07/2017
Revision Date	Prior Authorization: January 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Pulmicort Flexhaler

Products Affected

- PULMICORT FLEXHALER

PA Criteria	Criteria Details
Covered Uses	Asthma
Exclusion Criteria	
Required Medical Information	Documented diagnosis of Asthma
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of Asmanex and Qvar
QL Criteria	1 inhaler Per 1 month
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: November 30, 2016 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	
QL Criteria	60 units Per 1 fill
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/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	1 bottle Per 1 month
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

Qnasl

Products Affected

- QNASL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone and either OTC Nasacort 24HR or Flonase OTC
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Qnasl Childrens

Products Affected

- QNASL CHILDRENS

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 weeks of flunisolide or mometasone and either OTC Nasacort 24HR or Flonase OTC
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 ORAL CAPSULE ER 24 HOUR SPRINKLE 100 MG, 25 MG, 50 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic topiramate
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

Qudexy XR

Products Affected

- QUDEXY XR ORAL CAPSULE ER 24 HOUR SPRINKLE 150 MG, 200 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic topiramate
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

QUEtiapine Fumarate

Products Affected

- *quetiapine fumarate oral tablet 100 mg, 50 mg*

QL Criteria	3 tabs 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 tabs 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 tabs 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 tabs 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*

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

QUEtiapine Fumarate ER

Products Affected

- *quetiapine fumarate er oral tablet extended release 24 hour 50 mg*

QL Criteria	6 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

QuilliChew ER

Products Affected

- QUILLICHEW 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, dexmethylphenidate/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

QuilliChew ER

Products Affected

- QUILLICHEW 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, dexmethylphenidate/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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 bottle Per 1 fill
Notes/ References	Annual Review: 09/2017
Revision Date	Prior Authorization: May 16, 2017 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

QL Criteria	2 tabs 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	http://www.aetna.com/products/rxnonmedicare/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

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
QL Criteria	20 bottles Per 30 Days
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

Razadyne

Products Affected

- RAZADYNE ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 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	
QL Criteria	12 injections Per 28 Days
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	
QL Criteria	12 injections Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rebif Rebidose 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	
QL Criteria	12 injections Per 28 Days
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	
QL Criteria	12 injections Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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

Rectiv

Products Affected

- RECTIV

QL Criteria	30 grams Per 1 fill
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	20 inhalations Per 1 fill
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

PA Criteria	Criteria Details
Covered Uses	Opioid-induced constipation (OIC) in adults with chronic non-cancer pain, OIC in adults with advanced illness
Exclusion Criteria	
Required Medical Information	A documented diagnosis of opioid induced constipation due to non-cancer pain, OR a documented diagnosis of an advanced illness (i.e., incurable cancer, end-stage COPD/emphysema, cardiovascular disease/heart failure,Alzheimer's disease/dementia, HIV/AIDS), receiving palliative care, and response to laxative therapy has not been sufficient and documented concommitant use of opioid therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	0.6 ML Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: September 09, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Relistor

Products Affected

- RELISTOR SUBCUTANEOUS SOLUTION 8 MG/0.4ML

PA Criteria	Criteria Details
Covered Uses	Opioid-induced constipation (OIC) in adults with chronic non-cancer pain, OIC in adults with advanced illness
Exclusion Criteria	
Required Medical Information	A documented diagnosis of opioid induced constipation due to non-cancer pain, OR a documented diagnosis of an advanced illness (i.e., incurable cancer, end-stage COPD/emphysema, cardiovascular disease/heart failure,Alzheimer's disease/dementia, HIV/AIDS), receiving palliative care, and response to laxative therapy has not been sufficient and documented concommitant use of opioid therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	0.4 ML Per 1 Day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: September 09, 2016 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 EA 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	
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

Repaglinide-Metformin HCl

Products Affected

- *repaglinide-metformin hcl*

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

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/PCSK9.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/PCSK9.html
QL Criteria	2 injections 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/PCSK9.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/PCSK9.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/PCSK9.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/PCSK9.html
QL Criteria	2 injections Per 28 Days
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Requip XL

Products Affected

- REQUIP XL ORAL TABLET EXTENDED RELEASE 24 HOUR 12 MG

QL Criteria	2 tabs 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 tab 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

PA Criteria	Criteria Details
Covered Uses	open-angle glaucoma, ocular hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of glaucoma or ocular hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
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: December 07, 2016 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 cap 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Revatio

Products Affected

- REVATIO INTRAVENOUS RECONSTITUTED
- REVATIO ORAL SUSPENSION

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 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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 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	
QL Criteria	1 cap Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rexulti

Products Affected

- REXULTI

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder, Schizophrenia
Exclusion Criteria	
Required Medical Information	Documented diagnosis of major depressive disorder or Schizophrenia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Step Therapy
ST Criteria	FOR MAJOR DEPRESSIVE DISORDER: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine). FOR SCHIZOPHRENIA: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine) and Latuda.
QL Criteria	1 tablet Per 1 Day
Notes/ References	Annual Review: 08/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: October 27, 2017 Quantity Limits: August 25, 2015

Reyataz

Products Affected

- REYATAZ ORAL CAPSULE 150 MG
- REYATAZ ORAL CAPSULE 300 MG

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

Reyataz

Products Affected

- REYATAZ ORAL CAPSULE 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

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/Riastap.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

Rilutek

Products Affected

- RILUTEK

PA Criteria	Criteria Details
Covered Uses	amyotrophic lateral sclerosis (ALS)
Exclusion Criteria	
Required Medical Information	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: April 11, 2016 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	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: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Risedronate Sodium

Products Affected

- *risedronate sodium oral tablet 150 mg*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
QL Criteria	1 tablet Per 28 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*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
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*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alendronate 70mg
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

Risedronate Sodium

Products Affected

- *risedronate sodium oral tablet delayed release*

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 aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or 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

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	2 tabs 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 3 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 tabs 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	4 tabs 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	2 tabs 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 3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
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

RisperDAL M-TAB

Products Affected

- RISPERDAL M-TAB ORAL TABLET DISPERSIBLE 4 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, clozapine, or Latuda
QL Criteria	4 tabs 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*
- *risperidone oral tablet dispersible 0.5 mg*
- *risperidone oral tablet dispersible 1 mg, 2 mg*

QL Criteria	2 tabs 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 3 mg*
- *risperidone oral tablet dispersible 3 mg*

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

RisperiDONE M-TAB

Products Affected

- RISPERIDONE M-TAB ORAL TABLET DISPERSIBLE 0.5 MG, 1 MG, 2 MG

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

RisperiDONE M-TAB

Products Affected

- RISPERIDONE M-TAB ORAL TABLET DISPERSIBLE 3 MG

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

RisperiDONE M-TAB

Products Affected

- RISPERIDONE M-TAB ORAL TABLET DISPERSIBLE 4 MG

QL Criteria	4 tabs 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, dexmethylphenidate/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, dexmethylphenidate/sr, dextroamphetamine, methamphetamine, methylphenidate/er/sr, atomoxetine or Vyvanse
QL Criteria	1 cap 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, dexmethylphenidate/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

Rivastigmine

Products Affected

- *rivastigmine*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rivastigmine Tartrate

Products Affected

- *rivastigmine tartrate*

PA Criteria	Criteria Details
Covered Uses	Alzheimer's Disease
Exclusion Criteria	
Required Medical Information	Documented diagnosis of mild, moderate, severe Alzheimer's Disease
Age Restrictions	less than 40 years old
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 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/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

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 tabs 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 tab 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 tablets 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

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

Rozerem

Products Affected

- ROZEREM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
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 ORAL TABLET 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/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	2 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: January 09, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Rubraca

Products Affected

- RUBRACA 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/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

Rythmol SR

Products Affected

- RYTHMOL SR

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

Sabril

Products Affected

- SABRIL

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 packs 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 ORAL TABLET 15 MG

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	
QL Criteria	1 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Samsca

Products Affected

- SAMSCA ORAL TABLET 30 MG

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	
QL Criteria	2 tabs Per 1 day
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 1 fill
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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and 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

Savaysa

Products Affected

- SAVAYSA

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

Savella

Products Affected

- SAVELLA

PA Criteria	Criteria Details
Covered Uses	Fibromyalgia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of fibromyalgia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of duloxetine and Lyrica
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: July 18, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Savella Titration Pack

Products Affected

- SAVELLA TITRATION PACK

PA Criteria	Criteria Details
Covered Uses	Fibromyalgia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of fibromyalgia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of duloxetine and Lyrica
QL Criteria	2 tablets Per 1 Day
Notes/ References	Annual Review: 03/2017
Revision Date	Prior Authorization: July 18, 2017 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 one month each of Spiriva and 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 150 MG

QL Criteria	2 tabs 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 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

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

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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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	3 tabs 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	4 tabs 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 one generic medication such as aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine and Latuda
QL Criteria	6 EA 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

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	2 EA 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 400 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	2 tabs 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

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder, Bipolar disorder or schizophrenia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder, Bipolar Disorder or Schizophrenia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	FOR MAJOR DEPRESSIVE DISORDER: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine). FOR A DIAGNOSIS OF BIPOLAR DISORDER OR SCHIZOPHRENIA: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine) and Latuda.
QL Criteria	1 tab Per 1 day
Notes/ References	Annual Review: 06/2017

Revision Date	Prior Authorization: December 20, 2016 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

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder, Bipolar disorder or schizophrenia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder, Bipolar Disorder or Schizophrenia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	FOR MAJOR DEPRESSIVE DISORDER: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine). FOR A DIAGNOSIS OF BIPOLAR DISORDER OR SCHIZOPHRENIA: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine) and Latuda.
QL Criteria	2 tabs Per 1 day
Notes/ References	Annual Review: 06/2017

Revision Date	Prior Authorization: December 20, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SEROquel XR

Products Affected

- SEROQUEL XR ORAL TABLET EXTENDED RELEASE 24 HOUR 50 MG

PA Criteria	Criteria Details
Covered Uses	Major depressive disorder, Bipolar disorder or schizophrenia
Exclusion Criteria	
Required Medical Information	A documented diagnosis of Major Depressive Disorder, Bipolar Disorder or Schizophrenia
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	FOR MAJOR DEPRESSIVE DISORDER: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine). FOR A DIAGNOSIS OF BIPOLAR DISORDER OR SCHIZOPHRENIA: A documented contraindication, intolerance, allergy, or failure of one atypical generic antipsychotic (i.e. aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone, paliperidone er, or clozapine) and Latuda.
QL Criteria	6 tabs Per 1 day
Notes/ References	Annual Review: 06/2017
Revision Date	Prior Authorization: December 20, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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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 tablet 100 mg*

QL Criteria	2 tabs 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	1.5 tag Per 1 Day
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	
QL Criteria	2 amps Per 1 day
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	3 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 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
QL Criteria	1 pen Per 1 fill
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
QL Criteria	1 syringe Per 1 month
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	188 tabs Per 365 dayss
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sivextro

Products Affected

- SIVEXTRO ORAL

QL Criteria	6 tabs Per 1 fill
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

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

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/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

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
QL Criteria	1 cap Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Soolantra

Products Affected

- SOOLANTRA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of any of the preferred topical generic alternatives, metronidazole or sulfacetamide sodium with sulfur
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

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

Soriatane

Products Affected

- SORIATANE ORAL CAPSULE 17.5 MG, 25 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

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 tab 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 capsule Per 1 Day
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

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

Sporanox

Products Affected

- SPORANOX ORAL CAPSULE

QL Criteria	1 cap 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

Sporanox Pulsepak

Products Affected

- SPORANOX PULSEPAK

QL Criteria	1 cap 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

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

Sprycel

Products Affected

- SPRYCEL ORAL TABLET 100 MG, 140 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	1 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sprycel

Products Affected

- SPRYCEL ORAL TABLET 20 MG, 50 MG, 70 MG, 80 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 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stelara

Products Affected

- STELARA INTRAVENOUS

PA Criteria	Criteria Details
Covered Uses	Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Stelara.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/Stelara.html
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stelara

Products Affected

- STELARA 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/Stelara.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/Stelara.html
QL Criteria	1 syringe Per 1 month
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Stiolto Respimat

Products Affected

- STIOLTO RESPIMAT

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

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/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 tabs 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, dexmethylphenidate/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, dexmethylphenidate/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

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	
QL Criteria	1 tab Per 1 day
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 30 Days
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

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

Subsys

Products Affected

- SUBSYS

PA Criteria	Criteria Details
Covered Uses	For pain due to malignant diagnosis only
Exclusion Criteria	Use in non-malignant pain
Required Medical Information	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 meet step therapy requirements, or the patient is terminally ill.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months

PA Criteria	Criteria Details
Other Criteria	For additional quantities, the member must have a documented diagnosis of cancer and prescription is written by an oncologist or pain specialist, or the member is enrolled in a hospice program or meets hospice criteria, or the member is terminally ill, or the patient has signed an opioid agreement in support of clinical guidelines by the American Pain Society and the American Academy of Pain Medicine. In addition, there must be documentation of one of the following: (1) A Healthcare Provider verbal confirmation that an agreement has been signed by the patient meets the criteria requirement (exceptions to requiring the signed opioid agreement for additional quantities are only for those patients that have a diagnosis of cancer or that are enrolled in a hospice program), or (2) the member has current diagnosis of cancer(see exception to opioid agreement above) as the primary cause of the pain and is currently on long-acting opioid and is being titrated on the long-acting opioid by physician, and the member has tried and failed an adequate trial of two weeks of a single entity or combination pain medication containing an immediate release acting opioid (ex. oxycodone, morphine sulfate oral(Roxanol), oxymorphone(Opana), hydromorphone(Dilaudid), oxycodone/apap(Percocet))
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one week each of 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: October 10, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sular

Products Affected

- SULAR ORAL TABLET EXTENDED RELEASE 24 HOUR 17 MG, 34 MG, 8.5 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

SulfaSALAzine

Products Affected

- *sulfasalazine oral*

QL Criteria	8 tabs 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 tabs Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

SUMAtriptan

Products Affected

- *sumatriptan 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

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

SUMAtriptan Succinate

Products Affected

- *sumatriptan succinate subcutaneous solution 6 mg/0.5ml*

QL Criteria	8 vials Per 1 fill
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 subcutaneous solution auto-injector 4 mg/0.5ml, 6 mg/0.5ml*

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

SUMAtriptan Succinate Refill

Products Affected

- *sumatriptan succinate refill subcutaneous solution cartridge*

QL Criteria	2 boxes Per 1 fill
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

Sutent

Products Affected

- SUTENT ORAL CAPSULE 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/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 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sutent

Products Affected

- SUTENT ORAL CAPSULE 25 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	2 capsules Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sutent

Products Affected

- SUTENT ORAL CAPSULE 37.5 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	1 cap Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Sutent

Products Affected

- SUTENT ORAL CAPSULE 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/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 cap Per 1 day
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/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

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 ORAL CAPSULE 12-25 MG, 12-50 MG, 6-25 MG, 6-50 MG

QL Criteria	1 cap 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

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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

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

Synalar

Products Affected

- SYNALAR EXTERNAL CREAM
- SYNALAR EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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	(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

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 SUSPENSION

QL Criteria	60 gram 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*

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: October 11, 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	4 caps Per 1 Day
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/Tagrisso.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: August 25, 2015 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	3 bottles Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tanzeum

Products Affected

- TANZEUM

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Victoza and Trulicity
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/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 tab Per 1 day
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/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	4 caps Per 1 day
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	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 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	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
Notes/ References	
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Taztia XT

Products Affected

- TAZTIA XT ORAL CAPSULE EXTENDED RELEASE 24 HOUR 120 MG, 180 MG, 300 MG, 360 MG

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

Taztia XT

Products Affected

- TAZTIA XT ORAL CAPSULE EXTENDED RELEASE 24 HOUR 240 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

Tecfidera

Products Affected

- TECFIDERA

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 caps Per 1 day
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 2 preferred ACE-I or ARB . Formulary Angiotensin Converting Enzyme Inhibitors (ACEI) & ACEI combinations include: Prinivil, Zestril (lisinopril), Lotensin/Lotensin HCT/Lotrel (benazepril), Vasotec (enalapril), Accupril (quinapril), Mavik (trandolapril),Univasc (moexipril). Formulary Angiotensin Receptor Blocker (ARB) & ARB combinations include: Cozaar/Hyzaar (losartan), Benicar/Benicar HCT (olmesartan), Micardis/Micardis HCT (telmisartan) , Diovan/Diovan HCT (valsartan), Avapro/Avalide (irbesartan), Atacand/Atacand HCT (candesartan), Teveten /Teveten HCT (eprosartan), Edarbi/Edarbyclor (azilsartan)
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

Tekturna HCT

Products Affected

- TEKTURNA HCT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of 2 preferred ACE-I or ARB . Formulary Angiotensin Converting Enzyme Inhibitors (ACEI) & ACEI combinations include: Prinivil, Zestril (lisinopril), Lotensin/Lotensin HCT/Lotrel (benazepril), Vasotec (enalapril), Accupril (quinapril), Mavik (trandolapril),Univasc (moexipril). Formulary Angiotensin Receptor Blocker (ARB) & ARB combinations include: Cozaar/Hyzaar (losartan), Benicar/Benicar HCT (olmesartan), Micardis/Micardis HCT (telmisartan) , Diovan/Diovan HCT (valsartan), Avapro/Avalide (irbesartan), Atacand/Atacand HCT (candesartan), Teveten /Teveten HCT (eprosartan), Edarbi/Edarbyclor (azilsartan)
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

Telmisartan

Products Affected

- *telmisartan*

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

Telmisartan-Amlodipine

Products Affected

- *telmisartan-amlodipine*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of amlodipine in combination with two of the following: Atacand, Avapro, Cozaar, Micardis
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

Telmisartan-HCTZ

Products Affected

- *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

Temazepam

Products Affected

- *temazepam oral capsule 22.5 mg, 7.5 mg*

QL Criteria	1 cap 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	
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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
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/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

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	
QL Criteria	10 grams Per 1 Day
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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	
QL Criteria	2.5 grams Per 1 Day
Notes/ References	Annual Review: 02/2017

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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 ml Per 1 day
Notes/ References	

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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

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

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/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

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

Tiazac

Products Affected

- TIAZAC ORAL CAPSULE EXTENDED RELEASE 24 HOUR 120 MG, 180 MG, 300 MG, 360 MG, 420 MG

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

Tiazac

Products Affected

- TIAZAC ORAL CAPSULE EXTENDED RELEASE 24 HOUR 240 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

Tivicay

Products Affected

- TIVICAY

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

Tivicay

Products Affected

- TIVICAY

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

Tobi

Products Affected

- TOBI

QL Criteria	56 vials Per 1 fill
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	1 box Per 28 dayss
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tobramycin

Products Affected

- *tobramycin inhalation*

QL Criteria	56 vials Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tolterodine Tartrate ER

Products Affected

- *tolterodine tartrate er*

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

Topicort

Products Affected

- TOPICORT EXTERNAL CREAM
- TOPICORT EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of betamethasone dipropionate (cream/ointment/lotion)
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Topicort

Products Affected

- TOPICORT EXTERNAL CREAM
- TOPICORT EXTERNAL GEL
- TOPICORT EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of triamcinolone (cream/ointment/lotion)
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 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Toprol XL

Products Affected

- TOPROL XL ORAL TABLET EXTENDED RELEASE 24 HOUR 100 MG, 50 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of metoprolol succinate
QL Criteria	1.5 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Toprol XL

Products Affected

- TOPROL XL ORAL TABLET EXTENDED RELEASE 24 HOUR 200 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of metoprolol succinate
QL Criteria	2 tabs Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Toprol XL

Products Affected

- TOPROL XL ORAL TABLET EXTENDED RELEASE 24 HOUR 25 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of metoprolol succinate
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

Toujeo SoloStar

Products Affected

- TOUJEO SOLOSTAR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month each of Levemir and Tresiba
Notes/ References	Annual Review: 03/2016
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 and Myrbetriq and one generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin, oxybutynin 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

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 tab 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

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

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	1 tabletts 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)*

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-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

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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

Tranexamic Acid

Products Affected

- *tranexamic acid oral*

QL Criteria	30 tablet 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	2 blisters Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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 cream*
- *tretinoin external gel 0.01 %, 0.025 %*

PA Criteria	Criteria Details
Covered Uses	Acne Vulgaris (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	

Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretinoin Microsphere

Products Affected

- *tretinoin microsphere*

PA Criteria	Criteria Details
Covered Uses	Acne Vulgaris (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
Notes/ References	

Revision Date	Prior Authorization: April 11, 2016 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 (including comedonal, cystic, nodular and papular acne), actinic keratoses with lesions, hypertrophic scars or keloids, keratosis follicularis (e.g., Darier's disease, Darier-White disease), facial flat warts, multiple flat warts (e.g.,common warts, plantar warts)
Exclusion Criteria	
Required Medical Information	For members greater than 35 years old, the following criteria must be met: Documented diagnosis of acne vulgaris (includes comedonal, cystic, nodular & papular acne), or Documented diagnosis of 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, or • Documented diagnosis of hypertrophic scars or keloids AND intralesional injection of corticosteroids was ineffective or not tolerated, or Documented diagnosis of keratosis follicularis (Darier's disease, Darier-White disease), or Documented diagnosis of facial flat warts, or Documented diagnosis of multiple flat warts (includes common warts and plantar warts)
Age Restrictions	Prior authorization only applies for members greater than 35 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of tretinoin and Epiduo
Notes/ References	

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Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Tretten

Products Affected

- TRETTEN

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

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

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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

Tribenzor

Products Affected

- TRIBENZOR

ST Criteria	A documented contraindication, intolerance, allergy, or failure of amlodipine in combination with two of the following: Atacand HCT, Avalide, Hyzaar, Micardis 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

Tricor

Products Affected

- TRICOR

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

Tridesilon

Products Affected

- TRIDESILON

ST Criteria	A documented contraindication, intolerance, allergy, or failure of alclometasone cream/ointment
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 tab Per 1 day
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Trilipix

Products Affected

- TRILIPIX

QL Criteria	1 cap 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	

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

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

Triumeq

Products Affected

- TRIUMEQ

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

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic topiramate
QL Criteria	1 cap 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of generic topiramate
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

Trospium Chloride

Products Affected

- *trospium chloride*

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

Trospium Chloride ER

Products Affected

- *trospium chloride er*

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

Trulicity

Products Affected

- TRULICITY

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

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 one month each of Spiriva and Incruse Ellipta
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

TussiCaps

Products Affected

- TUSSICAPS

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

Twynsta

Products Affected

- TWYNSTA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of amlodipine in combination with two of the following: Atacand, Avapro, Cozaar, Micardis
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

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/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

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

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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

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

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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

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
QL Criteria	1 amp Per 1 day
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
QL Criteria	1 amp Per 1 day
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
QL Criteria	1 amp Per 1 day
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Uceris

Products Affected

- UCERIS ORAL

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one month of mesalamine DR (generic Asacol HD), Delzicol, Lialda, or Pentasa
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

Uceris

Products Affected

- UCERIS RECTAL

PA Criteria	Criteria Details
Covered Uses	Active mild to moderate 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 42 months
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ulesfia

Products Affected

- ULESFIA

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

Uloric

Products Affected

- ULORIC

ST Criteria	A documented contraindication, intolerance, allergy, or failure of allopurinol
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

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

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

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

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

Ultravate

Products Affected

- ULTRAVATE EXTERNAL CREAM
- ULTRAVATE EXTERNAL OINTMENT

ST Criteria	A documented contraindication, intolerance, allergy, or failure of augmented betamethasone (cream/ointment/lotion/gel)
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

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 tab 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 Obstructive Pulmonary Disorder (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 one month each of Anoro Ellipta and 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/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 tube Per 1 month
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 ML 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 30s
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Valsartan

Products Affected

- *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

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

Valtrex

Products Affected

- VALTREX

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

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 caps Per 1 Day
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	2 tabs 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

Veltin

Products Affected

- VELTIN

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 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 tabs 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 tabs 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 tabs 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 tabs 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 cap 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 cap 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 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 tablet extended release 24 hour 225 mg, 37.5 mg, 75 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

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	
Notes/ References	
Revision Date	Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Verapamil HCl ER

Products Affected

- *verapamil hcl er oral capsule extended release 24 hour 100 mg, 300 mg*

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

Verapamil HCl ER

Products Affected

- *verapamil hcl er 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

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

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

Verelan PM

Products Affected

- VERELAN PM ORAL CAPSULE EXTENDED RELEASE 24 HOUR 100 MG, 300 MG

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

Verelan PM

Products Affected

- VERELAN PM 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

Versacloz

Products Affected

- VERSACLOZ

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Clozaril tablets
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: August 25, 2015

VESIcare

Products Affected

- VESICARE

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one preferred generic (i.e. trospium, trospium ER, tolterodine, tolterodine ER, oxybutynin)
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

Viberzi

Products Affected

- VIBERZI

PA Criteria	Criteria Details
Covered Uses	Diarrhea-predominant irritable bowel syndrome (IBS)
Exclusion Criteria	No known or suspected history of any of the following: does not have a gallbladder, diagnosis of pancreatitis, diagnosis of alcoholism, member drinks more than 3 alcoholic beverages/day, severe (Child-Pugh C) hepatic impairment, or anatomic or biochemical abnormalities of the gastrointestinal tract (e.g., biliary duct obstruction, sphincter of Oddi dysfunction, or severe constipation)
Required Medical Information	A documented diagnosis of diarrhea-predominant irritable bowel syndrome (IBS)
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
QL Criteria	2 tablets Per 1 day
Notes/ References	Annual Review: 10/2017
Revision Date	Prior Authorization: April 27, 2017 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

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

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

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Next update

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

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

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

Victoza

Products Affected

- VICTOZA SUBCUTANEOUS SOLUTION PEN-INJECTOR

QL Criteria	9 ML 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

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	3 tablets Per 1 Day
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

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

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

QL Criteria	2 tabs 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

Viramune XR

Products Affected

- VIRAMUNE XR ORAL TABLET EXTENDED RELEASE 24 HOUR 100 MG

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

Viramune XR

Products Affected

- VIRAMUNE XR ORAL TABLET EXTENDED RELEASE 24 HOUR 400 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

Viread

Products Affected

- VIREAD 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

Vistogard

Products Affected

- VISTOGARD

QL Criteria	20 packs Per 1 fill
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/ophthalmic_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 patches Per 28 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

Voltaren

Products Affected

- VOLTAREN TRANSDERMAL

QL Criteria	200 GM 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	4 tabs Per 1 day
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 one generic medication (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) plus Latuda
QL Criteria	4 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 3 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication (aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone) plus Latuda
QL Criteria	2 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 4.5 MG, 6 MG

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic medication (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 one generic medication (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

Wellbutrin SR

Products Affected

- WELLBUTRIN SR

QL Criteria	2 tabs 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/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

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

Xalatan

Products Affected

- XALATAN

PA Criteria	Criteria Details
Covered Uses	open-angle glaucoma, ocular hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of glaucoma or ocular hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
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: December 07, 2016 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 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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 tabs 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/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
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 tabs 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 tabs 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

Xifaxan

Products Affected

- XIFAXAN ORAL TABLET 200 MG

QL Criteria	9 tabs 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 ENCHEPHALOPATHY: 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

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

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

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

Xopenex HFA

Products Affected

- XOPENEX HFA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one 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

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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	
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/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	4 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 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: August 25, 2015

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/cataplexy-xyrem.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

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

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

2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Formulary
Last update 12/2017
Next update

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

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	
QL Criteria	3 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zegerid OTC

Products Affected

- ZEGERID OTC

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

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

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 tabs 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 cap 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 oral capsule 10 mg, 20 mg, 40 mg*
- ZENATANE ORAL CAPSULE 30 MG

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	
ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic oral antibiotic prescribed for treatment of acne (i.e., minocycline, doxycycline)
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 tabs 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 tablet 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 and either OTC Nasacort 24HR or Flonase OTC
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Ziana

Products Affected

- ZIANA

ST Criteria	A documented contraindication, intolerance, allergy, or failure of Epiduo
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

Zioptan

Products Affected

- ZIOPTAN

PA Criteria	Criteria Details
Covered Uses	open-angle glaucoma, ocular hypertension
Exclusion Criteria	
Required Medical Information	A documented diagnosis of glaucoma or ocular hypertension
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	
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: December 07, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of two generic statin medications (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

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

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 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

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

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	4 caps Per 1 day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZOLMitriptan

Products Affected

- *zolmitriptan oral tablet 2.5 mg*
- *zolmitriptan oral tablet dispersible 2.5 mg*

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

ZOLMitriptan

Products Affected

- *zolmitriptan oral tablet 5 mg*
- *zolmitriptan oral tablet dispersible 5 mg*

QL Criteria	3 tablets Per 1 fill
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 tabs 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	1.5 tag 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*

QL Criteria	2 tabs 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*

ST Criteria	A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone
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

Zomig

Products Affected

- ZOMIG NASAL

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 sprays Per 30 fills
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zomig

Products Affected

- ZOMIG ORAL TABLET 2.5 MG

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

Zomig

Products Affected

- ZOMIG ORAL TABLET 5 MG

QL Criteria	3 tablets Per 1 fill
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 tabs Per 30 days
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 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 tab 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

ST Criteria	A documented contraindication, intolerance, allergy, or failure of one generic non steroidal anti-inflammatory drug (NSAID)
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

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	3 tablets Per 1 day
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	3 tabs 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 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

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 04, 2017 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/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 tablets Per 1 Day
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	
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

ZyPREXA

Products Affected

- ZYPREXA ORAL

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	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 Zydis

Products Affected

- ZYPREXA ZYDIS

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	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/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 tabs 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/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 tablets Per 1 Day
Notes/ References	
Revision Date	Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyvox

Products Affected

- ZYVOX ORAL SUSPENSION RECONSTITUTED

QL Criteria	150 ml Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

Zyvox

Products Affected

- ZYVOX ORAL TABLET

QL Criteria	28 tabs Per 1 fill
Notes/ References	
Revision Date	Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015

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