2017 Aetna Pharmacy Drug Guide - Five Tier Open Value Small Group Formulary $\bf Abstral$

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 tablets 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 tablets 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 |

Accu-Chek Aviva Plus

Products Affected

• ACCU-CHEK AVIVA PLUS

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Accu-Chek Compact Plus Care

Products Affected

• ACCU-CHEK COMPACT PLUS CARE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Accu-Chek Multiclix Lancet Dev

Products Affected

• ACCU-CHEK MULTICLIX LANCET DEV

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Accu-Chek Nano SmartView

Products Affected

• ACCU-CHEK NANO SMARTVIEW

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 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 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 |

| Revision Date | Prior Authorization: November 21, 2016 Step Therapy: October 06, 2017 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/Act emra.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Act emra.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/Act emra.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Act emra.html |
| QL Criteria | 4 SYRINGES Per 28 DAYSs |
| 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/acti mmune.html |
| Exclusion Criteria | |
| Required Medical 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 |

Actoplus met XR

Products Affected

ACTOPLUS MET 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 |

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 |

Adagen

Products Affected

• ADAGEN

| 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 |

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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/pulmon aryhypertensionagents.html |
| QL Criteria | 2 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/pulmon aryhypertensionagents.html |
| QL Criteria | 3 TABS Per 1 DAYS |
| 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 diskus 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 |

Advocate Duo

Products Affected

• ADVOCATE DUO DEVICE

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Adynovate

Products Affected

adynovate

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

Adyphren

Products Affected

ADYPHREN

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

Adyphren Amp II

Products Affected

• ADYPHREN AMP II

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

Adyphren II

Products Affected

• ADYPHREN II

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

Adzenys XR-ODT

Products Affected

• ADZENYS XR-ODT

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Attention deficit hyperactivity disorder (ADHD) |
| Exclusion Criteria | |
| Required Medical Information | A documented diagnosis of Attention deficit hyperactivity disorder (ADHD) |
| Age Restrictions | For Quillivant Only- 17 years of age and older |
| Prescriber Restrictions | |
| Coverage Duration | 1 year |
| Other Criteria | |
| ST Criteria | A documented contraindication, intolerance, allergy, or failure of 14 days each of 3 of the following medications: amphetamine/dextroamphetamine/sr, 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 tablet 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 tablets 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 tablet 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 tabs Per 1 DAYS |
| 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 |

AgaMatrix Presto

Products Affected

• AGAMATRIX PRESTO

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 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 |

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/lys osomal_storage.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Alecensa

Products Affected

ALECENSA

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

Alendronate Sodium

Products Affected

• alendronate sodium oral tablet 10 mg, 40 mg, 5 mg

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

Alendronate Sodium

Products Affected

• alendronate sodium oral tablet 35 mg, 70 mg

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

Alfuzosin HCl ER

Products Affected

• alfuzosin hcl er

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

Alinia

Products Affected

• ALINIA ORAL SUSPENSION RECONSTITUTED

| QL Criteria | 60 ml 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 TABLET

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

Almotriptan Malate

Products Affected

• almotriptan malate

| QL Criteria | 6 tablets Per 30 dayss |
|----------------------|---|
| 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 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-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 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 |

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 one 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 |

ALPRAZolam ER

Products Affected

• alprazolam er

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

ALPRAZolam XR

Products Affected

• alprazolam xr

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

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 | 1 tablet 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 |

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 |

Amitiza

Products Affected

AMITIZA

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

Amlodipine Besylate-Valsartan

Products Affected

• amlodipine besylate-valsartan

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

Amlodipine-Olmesartan

Products Affected

• amlodipine-olmesartan

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

Amlodipine-Valsartan-HCTZ

Products Affected

• amlodipine-valsartan-hctz

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

Amnesteem

Products Affected

amnesteem

| 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 tablets 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 | |
| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62% |

| QL Criteria | 1 1.25 gm packet Per 1 day |
|----------------------|--|
| Notes/ References | Annual Review: 02/2017 |
| Revision Date | Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 | |
| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62% |

| QL Criteria | 5 grams-2 packets Per 1 day |
|----------------------|--|
| Notes/ References | Annual Review: 02/2017 |
| Revision Date | Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 | |
| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one month of Androgel 1.62% |

| QL Criteria | 4 pumps Per 1 day |
|----------------------|--|
| Notes/ References | Annual Review: 02/2017 |
| Revision Date | Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Anoro Ellipta

Products Affected

• ANORO ELLIPTA

| QL Criteria | 60 BLISTERS Per 30 DAYSs |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Antara

Products Affected

• ANTARA ORAL CAPSULE 30 MG, 90 MG

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

Anzemet

Products Affected

ANZEMET ORAL

| QL Criteria | 5 tablets Per 1 month |
|----------------------|---|
| 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 days |
|----------------------|---|
| 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 capsules 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, 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 TABS Per 1 DAYS |
| 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, 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 TABS Per 1 DAYS |
| 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/Alp ha-1 Antitrypsin Inhibitor Therapy.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Aranesp (Albumin Free)

Products Affected

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

• ARANESP (ALBUMIN FREE) INJECTION

| ARANESP (ALBUMIN FREE) INJECTION MCG/ML, 60 MCG/0.5ML | |
|---|---|
| PA Criteria | Criteria Details |
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Eryt hropoiesis_Stimulating_Agents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Arcalyst

Products Affected

ARCALYST

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Arca lyst.html |
| Exclusion Criteria | |
| Required Medical 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 capsule Per 1 day |
| Notes/ References | Annual Review: 07/2017 |
| Revision Date | Prior Authorization: November 29, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

ARIPiprazole

Products Affected

• aripiprazole oral solution

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

ARIPiprazole

Products Affected

• aripiprazole oral tablet 10 mg, 15 mg, 2 mg • aripiprazole oral tablet dispersible

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

Armodafinil

Products Affected

| • armodafinil oral tablet 150 mg | |
|----------------------------------|--|
| 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 |

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 |
|----------------------|---|
| 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 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 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 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 |

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 |

| 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 | 120 tablets Per 3 Days |
| Notes/ References | Annual Review: 06/2017 |
| Revision Date | Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Arzerra

Products Affected

ARZERRA

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Arzerra.html |
| Exclusion Criteria | |
| Required Medical 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 |

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 tablets 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 CP24 Per 1 DAYS |
|----------------------|---|
| 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 CP24 Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Atacand

Products Affected

• ATACAND ORAL TABLET 32 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 | 1 TABS Per 1 DAYS |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Atomoxetine HCl

Products Affected

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

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

Atomoxetine HCl

Products Affected

• atomoxetine hcl oral capsule 100 mg, 80 mg

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

Atorvastatin Calcium

Products Affected

• atorvastatin calcium oral

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

Austedo

Products Affected

• AUSTEDO

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/Aust edo.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Aust edo.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 |

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 |

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 | |

| 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 | |
| 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 |

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

Azilect

Products Affected

AZILECT

| 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 |

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 tablet Per 1 day |
| 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 capsules 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 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 |

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 Small Group Formulary Last Update 12/2017

Next Update

Bayer Contour Link Monitor

Products Affected

• BAYER CONTOUR LINK MONITOR

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Bayer Contour Monitor

Products Affected

• BAYER CONTOUR MONITOR KIT

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Bayer Contour Next EZ

Products Affected

• BAYER CONTOUR NEXT EZ

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Bayer Contour next Link

Products Affected

• BAYER CONTOUR NEXT LINK

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Bayer Contour Next Monitor

Products Affected

• BAYER CONTOUR NEXT MONITOR

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 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 tablet Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Benicar HCT

Products Affected

• BENICAR HCT

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

Benlysta

Products Affected

• BENLYSTA INTRAVENOUS

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benlysta.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/benl ysta.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/benl ysta.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/here ditary_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/here ditary_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 | 1 box Per 1 month |
| 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 | 56 ampules Per 30 DAYSs |
|----------------------|---|
| 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 |

Bexarotene

Products Affected

• bexarotene

| PA Criteria | Criteria Details |
|------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Targretin.html |
| Exclusion Criteria | |
| Required Medical 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 |

Bicalutamide

Products Affected

• bicalutamide

| 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 |

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 |

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 tablet 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/botu linum_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/botu linum_toxin.html |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Botox Cosmetic

Products Affected

• BOTOX COSMETIC INTRAMUSCULAR SOLUTION RECONSTITUTED 50 UNIT

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/botu linum_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 |

Bravelle

Products Affected

BRAVELLE

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

Breo Ellipta

Products Affected

 BREO ELLIPTA INHALATION AEROSOL POWDER BREATH ACTIVATED 100-25 MCG/INH

| QL Criteria | 2 blister Per 1 DAYS |
|----------------------|---|
| 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 INHALATION AEROSOL POWDER BREATH ACTIVATED 200-25 MCG/INH

| 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 |

Brilinta

Products Affected

• BRILINTA

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

Brisdelle

Products Affected

• BRISDELLE

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

Briviact

Products Affected

• BRIVIACT ORAL 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 | 4 milliliters 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 |

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, 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 | 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 |

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 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 |

BuPROPion HCl

Products Affected

• bupropion hcl oral

| 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 |

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 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 (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 month |
| 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 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 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 month |
| 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 tablet 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 tablets 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 month |
|----------------------|---|
| 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 suppository Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Candesartan Cilexetil

Products Affected

• candesartan cilexetil oral tablet 16 mg, 4 mg, 8 mg

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

Candesartan Cilexetil-HCTZ

Products Affected

• candesartan cilexetil-hctz

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

Capecitabine

Products Affected

• capecitabine

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 tablets 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 tablet 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/meta bolic_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

| QL Criteria | 1 TB24 Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Cardura XL

Products Affected

CARDURA XL

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

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 |

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 |

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 |

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/ga ucher_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 capsules 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/ga ucher_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 capsules Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Cetrotide

Products Affected

• CETROTIDE SUBCUTANEOUS KIT 0.25 MG

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

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 | |

| | 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/Chol bam.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Chorionic Gonadotropin

Products Affected

• chorionic gonadotropin intramuscular

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

Cialis

Products Affected

• CIALIS ORAL TABLET 2.5 MG

• CIALIS ORAL TABLET 5 MG

| PA Criteria | Criteria Details |
|------------------------------|--|
| 111 Ciliviiu | |
| 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 |

Ciclodan

Products Affected

• CICLODAN EXTERNAL SOLUTION

| 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 (paraaminosalicylic 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) |
| Notes/ References | Annual Review: 07/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Ciclopirox

Products Affected

• ciclopirox external solution

| 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 (paraaminosalicylic 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) |
| Notes/ References | Annual Review: 07/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Cimzia

Products Affected

• CIMZIA SUBCUTANEOUS KIT 2 X 200 MG

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Cimzia.html |
| 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 month |
| 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/Cinq air.html |
| Exclusion Criteria | |
| Required Medical 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/here ditary_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/here ditary_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 10 mg, 20 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 |

Citalopram Hydrobromide

Products Affected

• citalopram hydrobromide oral tablet 40 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 |

Claravis

Products Affected

• claravis

| 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 |

Clarinex-D 12 Hour

Products Affected

• CLARINEX-D 12 HOUR

| QL Criteria | 2 TB12 Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Clever Chek Auto-Code

Products Affected

• CLEVER CHEK AUTO-CODE

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Clever Choice Micro System

Products Affected

• CLEVER CHOICE MICRO SYSTEM

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Climara Pro

Products Affected

· CLIMARA PRO

| QL Criteria | 1 box (4 patches) Per 1 month |
|----------------------|---|
| 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 solution
- clindamycin phosphate 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 |

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 |

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 |

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 |

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 |

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 |

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 |

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 |

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

Products Affected

• clozapine oral tablet 100 mg

| QL Criteria | 9 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Products Affected

• clozapine oral tablet 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 |

Products Affected

• clozapine oral tablet 25 mg, 50 mg • clozapine oral tablet dispersible 25 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 |

Products Affected

• clozapine oral tablet dispersible 12.5 mg

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

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 |

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 |

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 |

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 |

CombiPatch

Products Affected

COMBIPATCH

| QL Criteria | 8 patches Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Combivent Respimat

Products Affected

• COMBIVENT RESPIMAT

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

Cometriq (100 mg Daily Dose)

Products Affected

• COMETRIQ (100 MG DAILY DOSE)

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 kits 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 caupsules 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 kits 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 tablet Per 1 day |
|----------------------|---|
| Notes/ References | |
| 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 month |
|----------------------|---|
| 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 capsule Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 one of the following: ACE Inhibitor or ACE Inhibitor/HCTZ combination or Angiotensin-Receptor Blocker or Angiotensin-Receptor Blocker/HCTZ combination |
| Notes/ References | Annual Review: 06/2017 |
| Revision Date | Prior Authorization: July 25, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 |

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 tablet 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/meta bolic_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 |

CVS Nicotine

Products Affected

• cvs nicotine transdermal patch 24 hour

| 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 |

CVS Nicotine Polacrilex

Products Affected

• cvs nicotine polacrilex mouth/throat lozenge 4 mg

| QL Criteria | 20 EA Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

CVS NTS Step 1

Products Affected

• cvs nts step 1

| 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 |

Cycloset

Products Affected

CYCLOSET

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

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/meta bolic_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/lys osomal_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/ophth almic_agents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 2 ML Per 1 DAYS |
| 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 EA Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Daklinza

Products Affected

DAKLINZA

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

Daliresp

Products Affected

DALIRESP

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

Dapsone

Products Affected

• dapsone external

| QL Criteria | 60 grams Per 30 dayss |
|----------------------|---|
| 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 | 1 capsule Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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/meta bolic_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/antivira l_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 |

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 |

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 TB24 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 |
|----------------------|---|
|----------------------|---|

Dexilant

Products Affected

DEXILANT

| 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 capsule 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 |
|----------------------|--|
|----------------------|--|

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 milliliters 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 tablets 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 | 4 capsules Per 1 Day |
|----------------------|---|
| 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 box Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 tablet Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: November 01, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Diclofenac Sodium

Products Affected

• diclofenac sodium transdermal gel 1 %

| QL Criteria | 200 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 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 tablets 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 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 step through one month each of generic Migranal and two of the following: naratriptan, rizatriptan, sumatriptan, zolmitriptan |
|----------------------|--|
| QL Criteria | 9 ML Per 30 Days |
| 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 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 12 hour 120 mg
- diltiazem hcl er 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 HCl ER

Products Affected

• diltiazem hcl er 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 Beads

Products Affected

- release 24 hour 120 mg, 180 mg, 300 mg, 360 mg
- diltiazem hcl er beads oral capsule extended diltiazem hcl er beads oral capsule extended release 24 hour 420 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 Beads

Products Affected

• diltiazem hcl er 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 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 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 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 |

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 |

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 capsules 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 ORAL TABLET 10 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 | 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 oral tablet 10 mg

| 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 | |
| QL Criteria | 1 tablet Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Donepezil HCl

Products Affected

• donepezil hcl oral tablet 23 mg, 5 mg • donepezil hcl oral tablet dispersible

| 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 |

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

Dronabinol

Products Affected

• dronabinol

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

Duavee

Products Affected

• DUAVEE

| QL Criteria | 1 TABS Per 1 DAYS |
|----------------------|---|
| 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 month |
|----------------------|---|
| 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 capsules Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

DULoxetine HCl

Products Affected

• duloxetine hcl oral capsule delayed release particles 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 |

DULoxetine HCl

Products Affected

• duloxetine hcl oral capsule delayed release particles 60 mg

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

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 |

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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical 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/botu linum_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 tablet Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Edarbyclor

Products Affected

EDARBYCLOR

| 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 tablet 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 tablet 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 tablet 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/lys osomal_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/ga ucher_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 |

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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Elmiron

Products Affected

• ELMIRON

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

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 capsule 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 | 2 tablets Per 1 Day |
|----------------------|---|
| 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 tablet Per 1 day |
| 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 tablet Per 1 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 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 | 4 syringes Per 1 month |
| 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 | 8 syringes Per 1 month |
| 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 RECONSTITUTED

| 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 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/En brel.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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 | 8 syringes Per 1 month |
| 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
- endocet 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 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 |

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 EA Per 1 Day |
|----------------------|---|
| 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/Ent yvio.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ent yvio.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 | |
| 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 |

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/Eryt hropoiesis_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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Eprosartan Mesylate

Products Affected

• eprosartan mesylate

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

EQ Nicotine

Products Affected

• eq nicotine transdermal

| 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 |

Erivedge

Products Affected

• ERIVEDGE

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

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/Idio pathic_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 EA 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/Idio pathic_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/Idio pathic_Pulmonary_Fibrosis.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 3 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Escitalopram Oxalate

Products Affected

• escitalopram oxalate oral solution

| QL Criteria | 20 ml Per 1 DAYS |
|----------------------|---|
| 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 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 tablet 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 tablet 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 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Eszopiclone

Products Affected

eszopiclone

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

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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Evamist

Products Affected

EVAMIST

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

Evekeo

Products Affected

EVEKEO

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

Exalgo

Products Affected

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

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

Exforge HCT

Products Affected

• EXFORGE HCT

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of amlodipine with 2 of the following(brand or generic if available): Atacand HCT, Avalide, Hyzaar, Micardis HCT |
|----------------------|--|
| 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/Anit dotes.html |
| Exclusion Criteria | |
| Required Medical 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 | 1 box Per 1 month |
| 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/lys osomal_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 |

FaLessa

Products Affected

• FALESSA ORAL KIT 20-1-0.1 MCG-MG

| QL Criteria | 1.5 TABS Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Famciclovir

Products Affected

• famciclovir oral tablet 125 mg, 250 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 |

Famciclovir

Products Affected

• famciclovir oral tablet 500 mg

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

Fanapt

Products Affected

• FANAPT

| 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 |

Fanapt Titration Pack

Products Affected

• FANAPT TITRATION PACK

| 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 |
|----------------------|--|
| 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 EA 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 |

Felodipine ER

Products Affected

• felodipine 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 |

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

Fenofibrate Micronized

Products Affected

• fenofibrate micronized

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

Fenofibric Acid

Products Affected

• fenofibric acid oral tablet

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

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 one week each of two (2) immediate-release opioids including morphine, hydrocodone, oxycodone, or hydromorphone |
| QL Criteria | 120 Lozenges 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 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 week each of fentanyl transmucosal lozenge and two other short acting opioids (i.e., morphine, hydrocodone, oxycodone, hydromorphone) |
| QL Criteria | 120 tablets Per 30 Days |
| Notes/ References | Annual Review: 06/2017 |
| Revision Date | Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Ferriprox

Products Affected

FERRIPROX

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

Fetzima

Products Affected

• FETZIMA

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

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

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 CP24 Per 1 DAYS |
| 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 TABS Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Finasteride

Products Affected

• finasteride oral tablet 5 mg

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

Fioricet/Codeine

Products Affected

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

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | All FDA approved indications |
| Exclusion Criteria | |
| Required Medical Information | (1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for 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 capsules Per 30 Days |
| Notes/ References | |
| Revision Date | Prior Authorization: September 06, 2017 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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 alclometasone cream/ointment |
|----------------------|---|
| 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
 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 0.05 %
- 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 cream 0.1 %

• 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 capsule 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 capsules 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 capsules 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 capsules 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 20 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 |

FLUoxetine HCl

Products Affected

• fluoxetine hcl oral tablet 60 mg

| QL Criteria | 1 tabs Per 1 DAYS |
|----------------------|---|
| 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 capsules 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 tablets 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 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 ER

Products Affected

• fluvoxamine maleate er

| QL Criteria | 2 cap Per 1 DAYS |
|----------------------|---|
| 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 ORAL CAPSULE EXTENDED RELEASE 24 HOUR 20 MG EXTENDED RELEASE 24 HOUR 25 MG, $\,$

35 MG

• FOCALIN XR ORAL CAPSULE

| 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 |

Follistim AQ

Products Affected

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

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

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 |

FORA D10 2-in-1 Monitor

Products Affected

• FORA D10 2-IN-1 MONITOR

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

FORA D15g 2-in-1 Monitor

Products Affected

• FORA D15G 2-IN-1 MONITOR

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

FORA D20 2-in-1 Monitor

Products Affected

• FORA D20 2-IN-1 MONITOR

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| 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/bon e_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/bon e_disease_agents.html |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 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 tablets 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 Flash System

Products Affected

• FREESTYLE FLASH SYSTEM

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

FreeStyle Freedom Lite

Products Affected

• FREESTYLE FREEDOM LITE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

FreeStyle InsuLinx System

Products Affected

• FREESTYLE INSULINX SYSTEM

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 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 System

Products Affected

• FREESTYLE SYSTEM

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 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 SUSPENSION

| 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 | |
| Notes/ References | Annual Review: 03/2017 |
| Revision Date | Prior Authorization: April 27, 2017 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 TABS Per 1 DAYS |
| 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 capsules 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 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Gabitril

Products Affected

• GABITRIL ORAL TABLET 12 MG

| QL Criteria | 4 TABS Per 1 DAYS |
|----------------------|---|
| 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 DAYS |
|----------------------|---|
| 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 |

Gammaplex

Products Affected

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

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

Gamunex-C

Products Affected

• GAMUNEX-C

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

Ganirelix Acetate

Products Affected

• ganirelix acetate

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

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/Gatt ex.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 kit Per 1 month |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

GaviLyte-C

Products Affected

• gavilyte-c

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

GaviLyte-G

Products Affected

• gavilyte-g

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

Gelnique

Products Affected

- GELNIQUE TRANSDERMAL GEL 10 %

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of Vesicare 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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.html |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

GenVisc 850

Products Affected

• GENVISC 850

| PA Criteria | Criteria Details |
|------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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/antivira l_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 |

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

Gilotrif

Products Affected

• GILOTRIF

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Antineoplastics.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 1 tablet 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/Alp ha-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 Diagnostic

Products Affected

• GLUCAGEN DIAGNOSTIC

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

GlucaGen HypoKit

Products Affected

• GLUCAGEN HYPOKIT

| 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 |

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 |

Gonal-f

Products Affected

• GONAL-F

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

Gonal-f RFF

Products Affected

• GONAL-F RFF

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

Gonal-f RFF Rediject

Products Affected

• GONAL-F RFF REDIJECT

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

Gralise

Products Affected

• GRALISE ORAL TABLET 300 MG

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of gabapentin |
|----------------------|---|
| QL Criteria | 1 tablet 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 tablets 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 starter pack Per 1 month |
| 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 HCI 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/here ditary_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/here ditary_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 |

Heather

Products Affected

heather

| QL Criteria | 1.5 TABS Per 1 DAYS |
|----------------------|---|
| 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 INTRAVENOUS KIT 3000 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 |

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 |

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/sedati ve-hypnotics.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 1 capsules Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

HM Nicotine

Products Affected

• hm 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 |

HM Nicotine Polacrilex

Products Affected

• hm nicotine polacrilex mouth/throat lozenge 2 mg

| QL Criteria | 20 EA Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 | 2 tablets 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 |

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/act har.html |
| Exclusion Criteria | |
| Required Medical 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 |

HumuLIN 70/30

Products Affected

• HUMULIN 70/30

| 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 |

HumuLIN N

Products Affected

• HUMULIN N

| 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 |

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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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
- hydrocodone-ibuprofen oral tablet 5-200 mg, 7.5-200 mg

| 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 | |

| 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 |

HYDROmorphone HCl

Products Affected

| hydromorphone hcl oral liquid • hydromorphone hcl rectal | |
|--|--|
| 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 |

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 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 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 | 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 hel 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 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 | 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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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 |

| 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 |

Ibandronate Sodium

Products Affected

• ibandronate sodium intravenous solution 3 mg/3ml

| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bon e_disease_agents.html |
|----------------------|---|
| 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 tablet 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 EA 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 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 |

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 tablets 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 tablet 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/I dhifa.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 1 tablet Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 08, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Ilaris

Products Affected

• ILARIS

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Ilar is.html |
| Exclusion Criteria | |
| Required Medical 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/Ilar is.html |
| Exclusion Criteria | |
| Required Medical 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| QL Criteria | 4 EA 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 SOLUTION 20 MG/ACT

| QL Criteria | 0.27 ml Per 1 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 SOLUTION 5 MG/ACT

| QL Criteria | 0.21 ml Per 1 DAYS |
|----------------------|---|
| 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 | 84 capsules Per 28 days |
| Notes/ References | |
| Revision Date | Prior Authorization: August 16, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Increlex

Products Affected

INCRELEX

| PA Criteria | Criteria Details |
|------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Inc relex.html |
| Exclusion Criteria | |
| Required Medical 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 |

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/Infl ectra.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ingre zza.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/Ingre zza.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 tablets 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 CP24 Per 1 DAYS |
|----------------------|---|
| 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 tablets 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 tablets 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 tablets 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 |

Invokamet

Products Affected

INVOKAMET

| QL Criteria | 2 tablets Per 1 DAYS |
|----------------------|---|
| 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 | 2 tablets 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 month |
|----------------------|---|
| 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 tablet Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Irbesartan-Hydrochlorothiazide

Products Affected

• irbesartan-hydrochlorothiazide

| 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 |

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/I ressa.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 EA 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 tablets 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/Anit dotes.html |
| Exclusion Criteria | |
| Required Medical 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/Anit dotes.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Jakafi

Products Affected

JAKAFI

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 |

Janumet

Products Affected

JANUMET

| QL Criteria | 999 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Janumet XR

Products Affected

• JANUMET XR

| QL Criteria | 999 tablet Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Janumet XR

Products Affected

• JANUMET XR

| QL Criteria | 999 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Januvia

Products Affected

JANUVIA

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

Jardiance

Products Affected

JARDIANCE

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

Jentadueto

Products Affected

JENTADUETO

| QL Criteria | 2 tablets Per 1 day |
|----------------------|---|
| Notes/ References | Annual Review: 05/2016 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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/ophth almic_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 |

Jevtana

Products Affected

JEVTANA

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/jevta na.html |
| Exclusion Criteria | |
| Required Medical 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 |

Jinteli

Products Affected

• jinteli

| 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 |

Jolivette

Products Affected

• jolivette

| 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 |

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 |
|----------------------|---|
|----------------------|---|

Juxtapid

Products Affected

JUXTAPID

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Antilipi demic 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/Antilipi demic 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 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 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 capsules Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Kalbitor

Products Affected

KALBITOR

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/here ditary_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/here ditary_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 EA 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 tablets 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/lys osomal_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 |

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 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 |

Kepivance

Products Affected

KEPIVANCE

| 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 |

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 |
|----------------------|---|
|----------------------|---|

Ketoconazole

Products Affected

ketoconazole 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 |

Ketorolac Tromethamine

Products Affected

• ketorolac tromethamine ophthalmic

| QL Criteria | 1 vial 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 | 20 tablets Per 28 dayss |
|----------------------|---|
| 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/carb onic_anhydrase_inhibitor.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 4 tablets Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Kevzara

Products Affected

KEVZARA

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Ke vzara.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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 TB24 Per 1 DAYS |
| 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/Kin eret.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Kin eret.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 |

Kogenate FS

Products Affected

• KOGENATE FS INTRAVENOUS KIT 3000 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 |

Kogenate FS Bio-Set

Products Affected

• KOGENATE FS BIO-SET INTRAVENOUS KIT 3000 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 |

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 tablets Per 1 day |
| Notes/ References | Annual Review: 05/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Kombiglyze XR

Products Affected

 KOMBIGLYZE XR ORAL TABLET EXTENDED RELEASE 24 HOUR 5-1000 MG, 5-500 MG

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

Korlym

Products Affected

KORLYM

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

Kroger Blood Glucose

Products Affected

 KROGER BLOOD GLUCOSE KIT W/DEVICE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Kroger Premium Blood Glucose

Products Affected

• KROGER PREMIUM BLOOD GLUCOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 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/gou t.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/meta bolic_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/Antilipi demic 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/Antilipi demic Agents_HOFH.html |
| QL Criteria | 4 SOLN Per 30 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 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 |

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 TAB Per 1 DAILY |
| 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 TAB Per 1 DAILY |
| 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 TAB Per 1 DAILY |
| Notes/ References | Annual Review: 09/2017 |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Products Affected

• lamotrigine er oral tablet extended release 24 hour 100 mg, 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 | 1 tablet 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 |

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 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 |

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 |

Products Affected

• lamotrigine er oral tablet extended release 24 hour 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 TB24 Per 1 DAYS |
| Notes/ References | Annual Review: 09/2017 |
| Revision Date | Prior Authorization: April 11, 2016 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 |

Larin Fe 1.5/30

Products Affected

• LARIN FE 1.5/30

| 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 |

Latuda

Products Affected

• LATUDA ORAL TABLET 120 MG, 20 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 tablet 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 bottles 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 tablet 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 | 6 ML 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| QL Criteria | 30 day supply Per 1 prescription |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Lenvima 14 MG Daily Dose

Products Affected

• LENVIMA 14 MG DAILY DOSE

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

Lenvima 18 MG Daily Dose

Products Affected

• LENVIMA 18 MG DAILY DOSE

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

Lenvima 20 MG Daily Dose

Products Affected

• LENVIMA 20 MG DAILY DOSE

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 | 30 day supply Per 1 prescription |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Lenvima 24 MG Daily Dose

Products Affected

• LENVIMA 24 MG DAILY DOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 | 30 day supply Per 1 prescription |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Lenvima 8 MG Daily Dose

Products Affected

• LENVIMA 8 MG DAILY DOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 prescription |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

LevETIRAcetam ER

Products Affected

• levetiracetam er oral tablet extended release 24 hour 500 mg

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

LevETIRAcetam ER

Products Affected

• levetiracetam er oral tablet extended release 24 hour 750 mg

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

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 |

Lialda

Products Affected

• LIALDA

| 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 |

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 %

| 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: August 25, 2015 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 |

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 capsule 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 |

Lipofen

Products Affected

LIPOFEN

| 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 |

Livalo

Products Affected

• LIVALO

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of Crestor and one generic statin (atorvastatin, fluvastatin, lovastatin, pravastatin, 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 |

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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| 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 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 |

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 |

Lovastatin

Products Affected

• lovastatin

| 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 |

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/ophth almic_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/lys osomal_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 |

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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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 ORAL CAPSULE

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Antineoplastics.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| 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 |

Lynparza

Products Affected

• LYNPARZA ORAL TABLET

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

Lyza

Products Affected

• LYZA

| 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 |

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/ophth almic_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/hyd roxyprogesterone.html |
| Exclusion Criteria | |
| Required Medical 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

| 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 |

Matzim LA

Products Affected

• matzim la oral tablet extended release 24 hour 180 mg, 300 mg, 360 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 |

Matzim LA

Products Affected

• matzim la oral tablet extended release 24 hour 240 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 |

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 |

Meijer Blood Glucose

Products Affected

• MEIJER BLOOD GLUCOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Meijer Premium Blood Glucose

Products Affected

• MEIJER PREMIUM BLOOD GLUCOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 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 tablets 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 tablets 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 10 mg, 5 mg

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

Menopur

Products Affected

MENOPUR

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

Menostar

Products Affected

MENOSTAR

| QL Criteria | 1 box (4 patches) Per 1 month |
|----------------------|---|
| 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 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 |

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 tablet 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 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 |

Methadone HCl

Products Affected

• methadone hcl oral concentrate

| PA Criteria | Criteria Details |
|-----------------------|------------------------------|
| Covered Uses | All FDA approved indications |
| Exclusion Criteria | |

| PA Criteria | Criteria Details |
|---------------------------------|---|
| 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). |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | See required medical information |

| PA Criteria | Criteria Details |
|----------------------|---|
| 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 | |

| PA Criteria | Criteria Details |
|---------------------------------|---|
| 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). |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | See required medical information |

| PA Criteria | Criteria Details |
|----------------------|---|
| Other Criteria | |
| QL Criteria | 30 mg 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 | |

| PA Criteria | Criteria Details |
|---------------------------------|--|
| 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 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). |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | See required medical information |

| PA Criteria | Criteria Details |
|----------------------|---|
| Other Criteria | |
| QL Criteria | 60 mg 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

| PA Criteria | Criteria Details |
|-----------------------|------------------------------|
| Covered Uses | All FDA approved indications |
| Exclusion Criteria | |

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Required Medical Information | (I) 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). |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | See required medical information |

| PA Criteria | Criteria Details |
|----------------------|---|
| 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 | |

| PA Criteria | Criteria Details |
|---------------------------------|--|
| 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 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). |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | See required medical information |

| PA Criteria | Criteria Details |
|----------------------|---|
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: November 27, 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 tablets 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 | 4 tablets Per 1 Day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Products Affected

• methylphenidate hcl oral solution 10 mg/5ml

| QL Criteria | 30 milliliters 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 |

Products Affected

• methylphenidate hcl oral solution 5 mg/5ml

| QL Criteria | 60 milliliters 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 |

Products Affected

• methylphenidate hcl oral tablet

| QL Criteria | 6 tablet 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 |

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 |

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 |

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 |

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 |

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 |

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 |

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 capsule Per 1 day |
|----------------------|---|
| Notes/ References | Annual Review: 09/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Methylphenidate HCl ER (LA)

Products Affected

- methylphenidate hcl er (la) oral capsule extended release 24 hour 20 mg
- methylphenidate hcl er (la) oral capsule extended release 24 hour 40 mg

| QL Criteria | 1 capsule Per 1 day |
|----------------------|---|
| Notes/ References | Annual Review: 09/2016 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Methylphenidate HCl ER (LA)

Products Affected

• methylphenidate hcl er (la) oral capsule extended release 24 hour 30 mg

| QL Criteria | 2 capsules Per 1 day |
|----------------------|---|
| Notes/ References | Annual Review: 09/2016 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Methylphenidate HCl ER (LA)

Products Affected

• methylphenidate hcl er (la) oral capsule extended release 24 hour 60 mg

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

Metoprolol Succinate ER

Products Affected

• metoprolol succinate er oral tablet extended release 24 hour 200 mg

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

Metoprolol Succinate ER

Products Affected

• metoprolol succinate er oral tablet extended release 24 hour 25 mg

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

Miacalcin

Products Affected

• MIACALCIN INJECTION

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bon e_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/bon e_disease_agents.html |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Mimvey

Products Affected

• mimvey

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

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/Eryt hropoiesis_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 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 tabs Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Modafinil

Products Affected

• modafinil oral tablet 100 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 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 | 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 |

Modafinil

Products Affected

• modafinil oral tablet 200 mg

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Excessive daytime sleepiness associated with narcolepsy, Excessive daytime sleepiness associated with obstructive sleep apnea/hypopnea syndrome (OSAHS), Shift Work Sleep Disorder |
| Exclusion Criteria | |
| Required Medical Information | FOR NARCOLEPSY: Documentation of diagnostic testing and clinical notations supporting diagnosis of Narcolepsy, such as MSLT, clinical progress notes, etc. (Failure to adequately support the diagnosis of narcolepsy may result in denial of coverage). FOR OSAHS: The prescribing physician is a sleep specialist, ear, nose and throat, neurologist or pulmonologist or has obtained a consult from a sleep specialist, and a Standard Diagnostic Nocturnal Polysomnography (NPSG) has confirmed the diagnosis of OSAHS, and the patient has received nasal continuous positive airway pressure (CPAP) or bilevel positive airway pressure (BIPAP) for at least 1 month, and CPAP or BIPAP therapy must be continued on a routine basis in combination with 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 tablets 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 |

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 |

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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical 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 tablet 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
 morphine sulfate rectal

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

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

Morphine Sulfate

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 (Concentrate)

Products Affected

• morphine sulfate (concentrate) oral solution 100 mg/5ml, 20 mg/ml

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

| 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 |

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 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 | 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 100 mg, 30 mg, 60 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 | 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 ER

Products Affected

• morphine sulfate er oral tablet extended release 15 mg, 200 mg

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

| PA Criteria | Criteria Details |
|----------------------|--|
| Coverage Duration | Length of Therapy; see required medical information |
| 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 capsules Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Mozobil

Products Affected

MOZOBIL

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Mozobil.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: October 27, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Multaq

Products Affected

MULTAQ

| 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 |

Mupirocin

Products Affected

• mupirocin external

| 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 |

Mupirocin Calcium

Products Affected

mupirocin calcium

| 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 |

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/my alept.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 | 0.5 VIAL Per 1 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 |

Myobloc

Products Affected

 MYOBLOC INTRAMUSCULAR SOLUTION 2500 UNIT/0.5ML, 5000 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/botu linum_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 |

Myorisan

Products Affected

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

| 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 |

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 tablet 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 tablets Per 1 day |
| Notes/ References | Annual Review: 03/2017 |
| Revision Date | Prior Authorization: September 12, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Myzilra

Products Affected

• myzilra

| 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 |

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/lys osomal_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 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 |

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/bon e_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 ctg 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/ANEOPL/ Nerlynx.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 6 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: August 02, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Nesina

Products Affected

NESINA

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of 1 month each of Januvia, Janumet, or Janumet XR; Tradjenta or Jentadueto; and generic alogliptin, alogliptin/pioglitazone, or alogliptin/metformin |
|----------------------|---|
| QL Criteria | 1 tablet Per 1 day |
| Notes/ References | Annual Review: 05/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Neulasta

Products Affected

• NEULASTA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

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

Neupogen

Products Affected

 NEUPOGEN INJECTION SOLUTION 300 MCG/ML, 480 MCG/1.6ML
 NEUPOGEN INJECTION SOLUTION PREFILLED SYRINGE

| WEG/WE, 400 WEG/1.0WE | |
|---------------------------------|--|
| 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 |

Neutek 2Tek Glucose/Pressure

Products Affected

• NEUTEK 2TEK GLUCOSE/PRESSURE

| QL Criteria | 1 meter Per 1 year |
|----------------------|---|
| 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 tablet 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 | |
| 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 |

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 pack 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 |

Nexplanon

Products Affected

NEXPLANON

| QL Criteria | 1 implant Per 1 year |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Next Choice One Dose

Products Affected

• next choice one dose

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

Nicoderm CQ

Products Affected

NICODERM CQ

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

Nicorelief

Products Affected

• nicorelief mouth/throat gum

| QL Criteria | 24 EA Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Nicorette

Products Affected

• NICORETTE MOUTH/THROAT GUM

| QL Criteria | 24 EA Per 1 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Nicotine

Products Affected

• nicotine transdermal patch 24 hour

| 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 Step 1

Products Affected

• nicotine step 1

| 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 Step 2

Products Affected

• nicotine step 2

| 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 Step 3

Products Affected

• nicotine step 3

| 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 |

Nicotrol

Products Affected

NICOTROL

| QL Criteria | 3 boxes-504 crtrg Per 1 month |
|----------------------|---|
| 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 | 4 bottles Per 1 day |
|----------------------|---|
| 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 tablets 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 tablet 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 tablets 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 tablet 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 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Nikki

Products Affected

NIKKI

| 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 |

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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| 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, 25.5 mg, 34 mg, 40 mg, 8.5 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 |

Nisoldipine ER

Products Affected

• nisoldipine er oral tablet extended release 24 hour 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 |

Nitroglycerin

Products Affected

• nitroglycerin translingual solution

| QL Criteria | 12 grams Per 1 fill |
|----------------------|---|
| 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/meta bolic_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 |

Nora-BE

Products Affected

• nora-be

| 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 |

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 |

Norlyroc

Products Affected

NORLYROC

| 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 |

Northera

Products Affected

• NORTHERA ORAL CAPSULE 100 MG

| PA Criteria | Criteria Details |
|------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northe ra.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 EA Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Northera

Products Affected

• NORTHERA ORAL CAPSULE 200 MG

| PA Criteria | Criteria Details |
|------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/Northe ra.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 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/Northe ra.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 EA Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Novarel

Products Affected

• novarel intramuscular solution reconstituted 10000 unit

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

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 |

Noxafil

Products Affected

• NOXAFIL ORAL TABLET DELAYED RELEASE

| QL Criteria | 93 TBEC Per 30 DAYSs |
|----------------------|---|
| Notes/ References | Annual Review: 09/2017 |
| 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/Inter leukin 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 |

| 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 | 6 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 |

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 |

| 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 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 |

Nuedexta

Products Affected

NUEDEXTA

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

Nuplazid

Products Affected

NUPLAZID

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/Nupl azid.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 |

NuvaRing

Products Affected

NUVARING

| QL Criteria | 1 ring Per 28 dayss |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 |
|----------------------|---|
| Notes/ References | Annual Review: 05/2017 |
| Revision Date | Prior Authorization: June 19, 2017 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/San dostatin.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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/Idio pathic_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 EA 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, • olanzapine oral tablet dispersible 5 mg, 7.5 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 |

OLANZapine

Products Affected

• olanzapine oral tablet 2.5 mg

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

OLANZapine-FLUoxetine HCl

Products Affected

• olanzapine-fluoxetine hcl

| 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 |

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 |

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

Omega-3-acid Ethyl Esters

Products Affected

• omega-3-acid ethyl esters

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

Omnaris

Products Affected

OMNARIS

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of 2 weeks of two of the following: 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 2

Products Affected

• ONETOUCH ULTRA 2

| QL Criteria | 1 KIT Per 365 DAYSs |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

OneTouch Ultra Blue

Products Affected

• ONETOUCH ULTRA BLUE

| QL Criteria | 300 strips Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

OneTouch Ultra Mini

Products Affected

• ONETOUCH ULTRA MINI

| QL Criteria | 1 KIT Per 365 DAYSs |
|----------------------|---|
| 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 |

OneTouch Verio IQ System

Products Affected

• ONETOUCH VERIO IQ SYSTEM

| QL Criteria | 1 KIT Per 365 DAYSs |
|----------------------|---|
| 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 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 |

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

Onzetra Xsail

Products Affected

ONZETRA XSAIL

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of sumatriptan nasal spray |
|----------------------|---|
| QL Criteria | 1 kit Per 30 Days |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Opana 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 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 | 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 |

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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 1 tablet Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Oravig

Products Affected

ORAVIG

| QL Criteria | 14 tablets 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/Ore ncia.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ore ncia.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/Ore ncia.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ore ncia.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/Ore ncia.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ore ncia.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/Ore ncia.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ore ncia.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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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/meta bolic_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 EA Per 1 Day |
| 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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical 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 tablet Per 1 day |
| Notes/ References | Annual Review: 05/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Osphena

Products Affected

OSPHENA

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

Otezla

Products Affected

• OTEZLA ORAL TABLET

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Ote zla.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ote zla.html |
| QL Criteria | 2 TABS Per 1 DAYS |
| 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/Ote zla.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Ote zla.html |
| QL Criteria | 1 KIT Per 365 DAYSs |
| 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 |

Ovidrel

Products Affected

OVIDREL

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

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

Oxtellar XR

Products Affected

• OXTELLAR XR ORAL TABLET EXTENDED RELEASE 24 HOUR 600 MG

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one month of oxcarbazepine |
|----------------------|---|
| 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

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 concentrate 100 mg/5ml • oxycodone hcl 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 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 HCI ER

Products Affected

• oxycodone hcl er oral tablet er 12 hour abuse-deterrent 10 mg, 20 mg, 40 mg, 80 mg

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

Products Affected

• oxycodone hcl er oral tablet er 12 hour abuse-deterrent 15 mg, 30 mg, 60 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 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 |

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 | |

| 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 |

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 | 28 tablets Per 1 fill |
| 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 | |

| PA Criteria | Criteria Details |
|----------------------|--|
| Coverage Duration | Length of Therapy; see required medical information |
| 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 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 |

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 tablet 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 tablets 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 oral tablet extended release 24 hour 12.5 mg, 37.5 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 |

PARoxetine HCl ER

Products Affected

• paroxetine hcl er oral tablet extended release 24 hour 25 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 |

PARoxetine Mesylate

Products Affected

• paroxetine mesylate

| QL Criteria | 1 capsule 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 |

Paxil

Products Affected

• PAXIL ORAL SUSPENSION

| QL Criteria | 30 ml Per 1 DAYS |
|----------------------|---|
| 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 | 4 liters 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 | 4 liters 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 |

Pentasa

Products Affected

• PENTASA ORAL CAPSULE EXTENDED RELEASE 250 MG

| QL Criteria | 16 capsules 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 tablets 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 |

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 | 4 milliliters 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 |

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 EXTERNAL GEL 0.015 %

| QL Criteria | 3 unit dose tubes Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Picato

Products Affected

• PICATO EXTERNAL GEL 0.05 %

| QL Criteria | 2 unit dose tubes 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 tablet 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 tablet 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 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Plegridy

Products Affected

• PLEGRIDY 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 | 28 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.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 |

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/PCS K9.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| QL Criteria | 2 syringes Per 1 Day |
| 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 TAB Per 1 DAILY |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 |

Pravastatin Sodium

Products Affected

• pravastatin sodium

| 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 |

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

Pregnyl

Products Affected

pregnyl

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

Premarin

Products Affected

• PREMARIN ORAL

| 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 |

Premphase

Products Affected

PREMPHASE

| 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 |

Prempro

Products Affected

PREMPRO

| 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 |

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 | 2 tablets 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 |
|---------------|--|
|---------------|--|

Prezista

Products Affected

• PREZISTA ORAL SUSPENSION

| QL Criteria | 2 bottles 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 150 MG, 600 MG, 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 |

Prezista

Products Affected

• PREZISTA ORAL TABLET 800 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 |

PriLOSEC OTC

Products Affected

· PRILOSEC 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 |

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 |

| 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 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 |
|---------------|---|
|---------------|---|

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/Eryt hropoiesis_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/lys osomal_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 | 8 CAP Per 1 DAYS |
| 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/lys osomal_storage.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/lysosomal_storage.html |
| QL Criteria | 25 CAP Per 1 DAYS |
| Notes/ References | |
| Revision Date | Prior Authorization: January 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Prodigy AutoCode Blood Glucose

Products Affected

 PRODIGY AUTOCODE BLOOD GLUCOSE KIT

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Progesterone Micronized

Products Affected

• progesterone micronized 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 |

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/Alp ha-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/bon e_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/bon e_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 EA Per 1 Day |
| 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 tablet Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 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 |

Protonix

Products Affected

• PROTONIX 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) |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: October 06, 2017 Quantity Limits: August 25, 2015 |

Proventil HFA

Products Affected

• PROVENTIL HFA

| ST Criteria | A documented contraindication, intolerance, allergy, or failure 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 |

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 betamethasone dipropionate (cream/ointment/lotion) |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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 | 2 ampules Per 1 day |
| 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 | 3.5 ML Per 1 DAYS |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Qbrelis

Products Affected

QBRELIS

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Hypertension, Heart Failure, Myocardial Infarction |
| Exclusion Criteria | |
| Required Medical Information | A documented diagnosis of hypertension (Approved only for ages 6 and older), Heart failure, or Myocardial Infarction AND must have a documented inability to swallow tablets/capsules |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 1 year |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 09, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 the preferred generic alternative, topiramate |
|----------------------|--|
| QL Criteria | 1 CAPS Per 1 DAYS |
| 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 the preferred generic alternative, topiramate |
|----------------------|--|
| 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 |

Products Affected

• quetiapine fumarate oral tablet 100 mg, 50 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 |

Products Affected

• quetiapine fumarate oral tablet 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 |

Products Affected

• quetiapine fumarate oral tablet 25 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 |

Products Affected

• quetiapine fumarate oral tablet 300 mg, 400 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 |

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 |

Products Affected

• quetiapine fumarate er oral tablet extended release 24 hour 300 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 |

Products Affected

• quetiapine fumarate er oral tablet extended release 24 hour 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 |

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 | 12 milliliters 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 |

RA Nicotine

Products Affected

• ra nicotine transdermal

| 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 |

RABEprazole Sodium

Products Affected

• rabeprazole sodium

| 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 | |
| QL Criteria | 1 tablet Per 1 Day |
| Notes/ References | Annual Review: 02/2017 |
| Revision Date | Prior Authorization: November 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Ranexa

Products Affected

RANEXA

| 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 |

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/meta bolic_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/meta bolic_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 capsules Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: December 13, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Rayos

Products Affected

RAYOS

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

Rebetol

Products Affected

REBETOL ORAL SOLUTION

| QL Criteria | 5 bottles Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Rebif

Products Affected

• REBIF SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CNS/multiple_sclerosis.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 12 syringes 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 syringes 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 syringes 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 syringes Per 28 Days |
| 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 | 1 tube 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 | 40 disks Per 1 year |
|----------------------|---|
| 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 dayss |
| 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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 |

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/PCS K9.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| QL Criteria | 2 units Per 28 Days |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Repatha Pushtronex System

Products Affected

• REPATHA PUSHTRONEX SYSTEM

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| QL Criteria | 1 syringe Per 30 days |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Repatha SureClick

Products Affected

• REPATHA SURECLICK

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/PCS K9.html |
| QL Criteria | 2 units Per 28 Days |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 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 |

Retin-A Micro Pump

Products Affected

• RETIN-A MICRO PUMP EXTERNAL GEL 0.08 %

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

Revatio

Products Affected

• REVATIO INTRAVENOUS

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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 |

Revatio

Products Affected

 REVATIO ORAL SUSPENSION RECONSTITUTED

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/pulmon aryhypertensionagents.html |
| 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 capsule 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

| • F | EYATAZ ORAL CAPSULE 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 |

Reyataz

Products Affected

• REYATAZ ORAL CAPSULE 200 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 |

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 |

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 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 |

Risedronate Sodium

Products Affected

• risedronate sodium oral tablet 30 mg, 5 mg

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

Risedronate Sodium

Products Affected

• risedronate sodium oral tablet 35 mg

release

• risedronate sodium oral tablet delayed

| 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 |

RisperiDONE

Products Affected

- risperidone oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg
- risperidone oral tablet dispersible 1 mg, 2 mg
- risperidone oral tablet dispersible 0.5 mg

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

RisperiDONE

Products Affected

risperidone oral tablet 3 mg
 risperidone oral tablet dispersible 3 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 |

RisperiDONE

Products Affected

risperidone oral tablet 4 mg
 risperidone oral tablet dispersible 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 |

RisperiDONE M-TAB

Products Affected

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

RisperiDONE M-TAB

Products Affected

 RISPERIDONE M-TAB ORAL TABLET DISPERSIBLE 3 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 |

RisperiDONE M-TAB

Products Affected

• RISPERIDONE M-TAB ORAL TABLET DISPERSIBLE 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 |

Rituxan

Products Affected

• RITUXAN INTRAVENOUS SOLUTION

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/Rit uxan.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Rit uxan.html |
| Notes/ References | |
| 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 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

ROPINIRole HCl ER

Products Affected

• ropinirole hcl er oral tablet extended release 24 hour 2 mg, 4 mg, 6 mg, 8 mg

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

Rosuvastatin Calcium

Products Affected

• rosuvastatin calcium

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

Rozerem

Products Affected

ROZEREM

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of zolpidem, zaleplon, or eszopiclone |
|----------------------|---|
| QL Criteria | 1 tablet Per 1 day |
| Notes/ References | Annual Review: 08/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Rubraca

Products Affected

• RUBRACA

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

Ruconest

Products Affected

RUCONEST

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/MISC/here ditary_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 |

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

Saizen

Products Affected

• SAIZEN

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

Saizen Click. Easy

Products Affected

• SAIZEN CLICK.EASY

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

Samsca

Products Affected

• SAMSCA 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 tablet 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 tablets 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 month |
|----------------------|---|
| Notes/ References | Annual Review: 10/2017 |
| Revision Date | Prior Authorization: August 25, 2015 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/San dostatin.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 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 |

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 • 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 |

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 |

Sensipar

Products Affected

• SENSIPAR

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/my alept.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 |

Serevent Diskus

Products Affected

SEREVENT 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 |

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 tablet 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, 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 | 2 tablets 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 |
|---------------|---|
|---------------|---|

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 tablets 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 tablet 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 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Sharobel

Products Affected

SHAROBEL

| 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 |

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/Sig nifor.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 SOLN Per 1 DAYS |
| 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/pulmon aryhypertensionagents.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: 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/Sili q.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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

| | TREFIELED STRINGE |
|------------------------------|--|
| 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 syringe Per 1 month |
| 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/Sim poni_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 vial 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 tablet 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/antimyc obacterial_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 EA Per 365 Days |
| 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 tablets Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Skyla

Products Affected

SKYLA

| QL Criteria | 1 IUD Per 365 DAYSs |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

SM Nicotine

Products Affected

• sm nicotine transdermal

| 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 |

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/meta bolic_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 |

Solia

Products Affected

• solia

| 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 |

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

Somatuline Depot

Products Affected

• SOMATULINE DEPOT

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

Somavert

Products Affected

SOMAVERT

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/growthhormone.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| 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 |

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 |

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 TABS Per 1 DAYS |
| 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 |

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 tablet 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 tablets 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/Stel ara.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Stel ara.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/Stel ara.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Stel ara.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 | |
| 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 |

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/lys osomal_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/antivira l_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 |

Striverdi Respimat

Products Affected

• STRIVERDI RESPIMAT

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

Suboxone

Products Affected

• SUBOXONE SUBLINGUAL FILM 12-3 MG

| QL Criteria | 2 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 |

Suboxone

Products Affected

• SUBOXONE SUBLINGUAL FILM 2-0.5 MG, 4-1 MG, 8-2 MG

| 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 |

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 |

SulfaSALAzine

Products Affected

• sulfasalazine oral

| 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 |

Sulfazine

Products Affected

• sulfazine

| 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 |

SUMAtriptan

Products Affected

• sumatriptan nasal

| QL Criteria | 3 nasal sprays Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

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 |

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 |

Products Affected

• sumatriptan succinate subcutaneous solution auto-injector 4 mg/0.5ml

| QL Criteria | 2 boxes (4 doses) Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Products Affected

• sumatriptan succinate 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 |

SUMAtriptan Succinate Refill

Products Affected

• sumatriptan succinate refill subcutaneous solution cartridge

| QL Criteria | 2 boxes (4 doses) Per 1 month |
|----------------------|---|
| 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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.html |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Supartz FX

Products Affected

SUPARTZ FX

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

Sutent

Products Affected

• SUTENT ORAL CAPSULE 37.5 MG, 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 | | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.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 | |

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 month |
|----------------------|---|
| 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 | |
| Poor compliance with current insulin regimen, Poor compliance prescribed self-blood glucose monitorings, An A1C greater than 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, Concurruse with other oral antidiabetic medications (except metformin 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 Small Group Formulary Last Update 12/2017

Next Update

SymlinPen 60

Products Affected

• SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR

| PA Criteria | Criteria Details | |
|--|---|--|
| Covered Uses | All FDA Approved uses | |
| Poor compliance with current insulin regimen, Poor compliance prescribed self-blood glucose monitorings, An A1C greater than 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, Concurruse with other oral antidiabetic medications (except metformin 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 Small Group 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/Syna gis.html | |
| Exclusion Criteria | | |
| Required Medical 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

| • | SYNAL | AR EXTERNAL | L 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 | |
| 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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 |

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 |

Synribo

Products Affected

SYNRIBO

| 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 |

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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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/visc osupplements.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/visc osupplements.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/meta bolic_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 grams 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| QL Criteria | 4 EA 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 |

Take Action

Products Affected

• take action

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

Targretin

Products Affected

• TARGRETIN EXTERNAL

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Targretin.html |
| Exclusion Criteria | |
| Required Medical 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 |

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 capsules 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 capsule 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 capsules 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 CPDR Per 1 DAYS |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 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 EA 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 EA 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 two 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 tablet Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Tekturna HCT

Products Affected

TEKTURNA HCT

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

Telmisartan-HCTZ

Products Affected

• telmisartan-hctz

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

Temazepam

Products Affected

• temazepam oral capsule 22.5 mg, 7.5 mg

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

Temovate

Products Affected

• TEMOVATE EXTERNAL CREAM

| 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 |

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%)

| (170) | (170) |
|---------------------------------|---|
| 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 |
|----------------------|--|
| Revision Date | Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Testosterone

Products Affected

• testosterone transdermal gel 25 mg/2.5gm (1%)

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

| QL Criteria | 2.5 grams Per 1 Day |
|----------------------|--|
| Notes/ References | Annual Review: 02/2017 |
| Revision Date | Prior Authorization: May 11, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Testosterone

Products Affected

• testosterone transdermal solution

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

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

Testosterone Cypionate

Products Affected

• testosterone cypionate intramuscular solution 100 mg/ml

| QL Criteria | 10 vials Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Testosterone Cypionate

Products Affected

• testosterone cypionate intramuscular solution 200 mg/ml

| QL Criteria | 10 ml Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 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/xena zine.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/xena zine.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| QL Criteria | 4 tablets Per 1 Day |
| Notes/ References | |
| Revision Date | Prior Authorization: May 31, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TGT Blood Glucose Monitoring

Products Affected

• TGT BLOOD GLUCOSE MONITORING

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TGT Nicotine Step One

Products Affected

• tgt nicotine step one

| 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 |

TGT Nicotine Step Three

Products Affected

• tgt nicotine step three

| 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 |

TGT Nicotine Step Two

Products Affected

• tgt nicotine step two

| 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 |

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/meta bolic_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 |

Thrive

Products Affected

• thrive mouth/throat gum 2 mg

| QL Criteria | 24 EA Per 1 DAYS |
|----------------------|---|
| 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 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 |

Tirosint

Products Affected

TIROSINT

| 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 |

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

Tivorbex

Products Affected

TIVORBEX

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

Tobi Podhaler

Products Affected

TOBI PODHALER

| QL Criteria | 1 CAPS 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 | 10 ml Per 1 day |
|----------------------|---|
| 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 | 999 capsule 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 tablet 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: December 22, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Tradjenta

Products Affected

TRADJENTA

| QL Criteria | 1 tablet Per 1 day |
|----------------------|---|
| Notes/ References | Annual Review: 05/2016 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TraMADol HCl

Products Affected

• tramadol hcl oral

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | All FDA approved indications |
| Exclusion Criteria | |
| Required Medical Information | (1) A MEMBER WILL RECEIVE LIFETIME APPROVAL OF THE REQUESTED MEDICATION WHEN: Member is in hospice care, if terminally ill or has end-of-life care (other than hospice), or has an active oncology diagnosis (pain medication being used for pain for cancer patients) or palliative care. (2) FOR A DOCUMENTED DIAGNOSIS OF MODERATE TO SEVERE ACUTE PAIN (INCLUDING POST-SURGICAL PAIN), TRAUMATIC INJURY, OR FOR A CHILD ON AN OPIOID WEAN AT TIME OF HOSPITAL DISCHARGE: additional medication after initial coverage will be approved for up to 1 month. (3) FOR A DOCUMENTED DIAGNOSIS OF CHRONIC PAIN: which includes conditions of chronic pain not mentioned above, the prescriber must certify there is an active treatment plan that includes but is not limited to a specific treatment objective, the use of other pharmacological and non-pharmacological agents for pain relief as appropriate, that there has been an informed consent document signed and an addiction risk assessment performed, and that a written/signed agreement between prescriber and patient addressing issues of prescription management, diversion, and the use of other substances exists. When these criteria are met, the medication will be approved for an initial 3 months. Continuation requests may be approved for up to an additional 6 months. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Length of Therapy; see required medical information |
| 2017 A atma Dla arma | ary David Child Eine Tion On on Walne Small Chang Farmulany |

| 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 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 | 1 tablet 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 | |
| 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 |

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 |

| PA Criteria | Criteria Details |
|----------------------|--|
| Other Criteria | |
| QL Criteria | 8 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Tranexamic Acid

Products Affected

• tranexamic acid oral

| QL Criteria | 30 tablets Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Trelegy Ellipta

Products Affected

• TRELEGY ELLIPTA

| QL Criteria | 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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/Tre mfya.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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 | |
|----------------------|--|
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Trezix

Products Affected

• TREZIX ORAL CAPSULE 320.5-30-16 MG

| Criteria Details |
|--|
| All FDA approved indications |
| |
| (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. |
| |
| |
| 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 |

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 tablet 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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 EA 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

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of Topamax |
|----------------------|---|
| QL Criteria | 1 CP24 Per 1 DAYS |
| 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 tablets 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 capsule Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TRUEresult Blood Glucose

Products Affected

• TRUERESULT BLOOD GLUCOSE

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TrueTrack Blood Glucose

Products Affected

• TRUETRACK BLOOD GLUCOSE KIT

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

TrueTrack Smart System

Products Affected

• TRUETRACK SMART SYSTEM

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Type 1 Diabetes Mellitus, Type 2 Diabetes Mellitus |
| Exclusion Criteria | |
| Required Medical Information | Documentation of a physical limitation that makes utilization of a Lifescan product unsafe, inaccurate or otherwise not feasible. Such limitations may include, but are not limited to, manual dexterity or visual impairment issues not accommodated by the features and capabilities of the Lifescan product line, or the member has hematocrit levels which are chronically less than 30% or greater than 55%. |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years, 1 meter per year |
| Other Criteria | |
| QL Criteria | 1 KIT Per 365 DAYSs |
| Notes/ References | |
| Revision Date | Prior Authorization: April 11, 2016 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 30 fills |
| 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 capsules Per 1 fill |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Tybost

Products Affected

TYBOST

| QL Criteria | 1 EA 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 | |
| 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 |

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/bon e_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/bon e_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/pulmon aryhypertensionagents.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 SOLN Per 1 DAYS |
| 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/pulmon aryhypertensionagents.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 ML 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/pulmon aryhypertensionagents.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 ML 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 Asacol HD, Delzicol, Lialda or Pentasa |
|----------------------|---|
| 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 |

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 | 2 canisters Per 1 month |
| 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 tablet Per 1 day |
| 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/pulmon aryhypertensionagents.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: 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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 |

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 GM Per 1 Day |
| 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/antivira ltopical.html |
| Exclusion Criteria | |
| Required Medical 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 |

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/antivira ltopical.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: November 13, 2017 |

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/antivira ltopical.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 TABS Per 30 DAYSs |
| 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 tablet Per 1 day |
|----------------------|---|
| 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical 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 DAYS |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Vecamyl

Products Affected

VECAMYL

| PA Criteria | Criteria Details |
|---------------------------------|--|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/CV/antihypertensive_misc.html |
| QL Criteria | 10 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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/Vemlid y.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Vemlid y.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 |

Products Affected

• venlafaxine hcl oral tablet 100 mg, 25 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 |

Products Affected

• venlafaxine hcl oral tablet 37.5 mg

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

Products Affected

• venlafaxine hcl oral tablet 50 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 |

Products Affected

• venlafaxine hcl oral tablet 75 mg

| QL Criteria | 5 tablets Per 1 day |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Products Affected

• venlafaxine hcl er oral capsule extended release 24 hour 150 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 |

Venlafaxine HCl ER

Products Affected

• venlafaxine hcl er oral capsule extended release 24 hour 37.5 mg, 75 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 |

Venlafaxine HCl ER

Products Affected

• venlafaxine hcl er oral tablet extended release 24 hour 225 mg

| QL Criteria | 1 tabs Per 1 DAYS |
|----------------------|---|
| 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/pulmon aryhypertensionagents.html |
| Exclusion Criteria | |
| Required Medical 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 capsule 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 capsules 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 |

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 tablet 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 |

| 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 | 1 box-2 or 3 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 |

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 |
| 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 |

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 tablet Per 1 Day |
| Notes/ References | Annual Review: 05/2017 |

| | Prior Authorization: August 25, 2015 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 | Diagnosis of major depressive disorder |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | 3 years |
| Other Criteria | |
| 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: October 10, 2016 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 |
|---------------|---|
|---------------|---|

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/lys osomal_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 DAYS |
|----------------------|---|
| 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 tablets 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 |

Viorele

Products Affected

• viorele

| 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 |

Viread

Products Affected

• VIREAD ORAL TABLET

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

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 |

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 tablets 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/ga ucher_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 |

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 |

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 |

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 |

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 tablet 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 |

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 |

Xalkori

Products Affected

XALKORI

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

Xatmep

Products Affected

XATMEP

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Treatment of acute lymphoblastic leukemia (ALL) or polyarticular juvenile idiopathic arthritis (pJIA) in pediatric patients |
| Exclusion Criteria | |
| Required Medical Information | A documented diagnosis of Acute Lymphoblastic Leukemia (ALL) in a pediatric patient (18 years and younger) as part of a multi-phase, combination chemotherapy maintenance regimen or a diagnosis of Polyarticular Juvenile Idiopathic Arthritis (PJIA) in pediatric patients (18 years and younger) who have had an insufficient therapeutic response to, or are intolerant of, an adequate trial of first-line therapy including full dose non-steroidal anti-inflammatory agents (NSAIDs). Regardless of diagnosis, the patient must have a documented inability to swallow tablets/capsules. |
| Age Restrictions | Approved for those 18 years of age or younger |
| Prescriber Restrictions | |
| Coverage Duration | 1 year |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: July 06, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Xeljanz

Products Affected

• XELJANZ

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

Xeljanz XR

Products Affected

XELJANZ XR

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

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/botu linum_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/Xer melo.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/bon e_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/bon e_disease_agents.html |
| 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 tablets Per 1 fill |
|----------------------|---|
| Notes/ References | Annual Review: 04/2017 |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Xifaxan

Products Affected

• XIFAXAN ORAL TABLET 550 MG

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Hepatic Encephalopathy, Irritable Bowel Syndrome (IBS) with Diarrhea. |
| Exclusion Criteria | |
| Required Medical Information | FOR HEPATIC 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 TAB Per 1 DAYS |
|----------------------|---|
| 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 |

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/Xola ir.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber 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/Xola ir.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 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 |

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

Xulane

Products Affected

• XULANE

| QL Criteria | 1 box (3 patches) Per 1 month |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Xultophy

Products Affected

XULTOPHY

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one of the following: Victoza, Byetta, Bydureon, Tanzeum, Trulicity, Adylixin, Lantus, Toujeo, Levemir, Tresiba, Basaglar |
|----------------------|--|
| QL Criteria | 5 pens Per 1 month |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Xuriden

Products Affected

• XURIDEN

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

Xylon

Products Affected

• XYLON

| QL Criteria | 120 tablets Per 30 Days |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: November 09, 2017 |

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/catapl exy-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 |

Yervoy

Products Affected

YERVOY

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/yervoy.html |
| Exclusion Criteria | |
| Required Medical 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 |

Zafirlukast

Products Affected

• zafirlukast

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

Zaleplon

Products Affected

• zaleplon

| 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 |

Zaltrap

Products Affected

• ZALTRAP

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/z altrap.html |
| Exclusion Criteria | |
| Required Medical 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 |

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/ga ucher_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 capsules 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| QL Criteria | 8 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zemaira

Products Affected

ZEMAIRA

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ENDO/Alp ha-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 |

Zenatane

Products Affected

• zenatane oral capsule 10 mg, 20 mg, 40 mg

| QL Criteria | 2 capsules 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 |

Zenatane

Products Affected

• ZENATANE ORAL CAPSULE 30 MG

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

Zenzedi

Products Affected

• ZENZEDI ORAL TABLET 10 MG, 5 MG

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

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 capsules 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 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 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/Gon adotropins.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| Notes/ References | |
| Revision Date | Prior Authorization: February 20, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zoledronic Acid

Products Affected

zoledronic acid intravenous concentrate
 zoledronic acid intravenous solution

| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/MUSC/bon e_disease_agents.html |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zolinza

Products Affected

ZOLINZA

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/ Antineoplastics.html |
| Exclusion Criteria | |
| Required Medical Information | |
| Age Restrictions | |
| Prescriber Restrictions | |
| Coverage Duration | Refer to the clinical policy bulletin above for details. |
| Other Criteria | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| 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 |

Products Affected

• zolmitriptan 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 |

Products Affected

• zolmitriptan 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 |

Products Affected

• zolmitriptan oral tablet dispersible 2.5 mg

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

Products Affected

• zolmitriptan oral tablet dispersible 5 mg

| QL Criteria | 30 tablet Per 30 days |
|----------------------|---|
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zolpidem Tartrate

Products Affected

• zolpidem tartrate oral

| 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 |

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 tablet 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 SOLUTION 5 MG

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one month of three of the following: naratriptan, rizatriptan, sumatriptan, or zolmitriptan |
|----------------------|--|
| QL Criteria | 1 box (6 doses) Per 1 month |
| 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 tablet Per 1 Day |
| Notes/ References | Annual Review: 06/2017 |
| Revision Date | Prior Authorization: July 19, 2017 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zorbtive

Products Affected

ZORBTIVE

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

Zorvolex

Products Affected

ZORVOLEX

| ST Criteria | A documented contraindication, intolerance, allergy, or failure of one generic non steroidal anti-inflammatory drug (NSAID) |
|----------------------|---|
| QL Criteria | 3 CAPS Per 1 DAYS |
| Notes/ References | |
| Revision Date | Prior Authorization: August 25, 2015 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zubsolv

Products Affected

• ZUBSOLV SUBLINGUAL TABLET SUBLINGUAL 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, 11.4-2.9 MG, 2.9-0.71 MG, 5.7-1.4 MG, 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 |

Zyban

Products Affected

• ZYBAN

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

Zydelig

Products Affected

• ZYDELIG

| PA Criteria | Criteria Details |
|---------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 CAP Per 1 DAYS |
| 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 | |
| ST Criteria | http://www.aetna.com/products/rxnonmedicare/data/2017/ANEOPL/Antineoplastics.html |
| QL Criteria | 5 CAP Per 1 DAYS |
| 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 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 tablets Per 1 day |
| Notes/ References | |
| Revision Date | Prior Authorization: September 21, 2016 Step Therapy: August 25, 2015 Quantity Limits: August 25, 2015 |

Zytiga

Products Affected

• ZYTIGA ORAL TABLET 500 MG

| PA Criteria | Criteria Details |
|------------------------------|---|
| Covered Uses | Refer to the clinical policy bulletin for details: http://www.aetna.com/products/rxnonmedicare/data/2017/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 |

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