- aripiprazole 10mg odt (New Starts Only)

- aripiprazole 15mg odt (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Both of the following: A) Member is unable to swallow aripiprazole tablet and B) Member is unable to use aripiprazole oral solution. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- tretinoin 0.01% gel
- tretinoin 0.025% gel
- tretinoin 0.1% cream

- tretinoin 0.025% cream

— tretinoin 0.05% cream

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ACTEMRA 162MG/0.9ML AUTO-INJECTOR

-ACTEMRA 162MG/0.9ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Enbrel, b) Hadlima, c) Rinvoq OR d) Xeljanz. For polyarticular juvenile idiopathic arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel OR c) Xeljanz. For giant cell arteritis (all requests): Trial of other agents not required. For systemic sclerosis-associated interstitial lung disease (initial requests): a) Diagnosis is confirmed with documentation provided of both of the following: i) HRCT scan AND ii) pulmonary function tests AND b) Trial of mycophenolate was ineffective or not tolerated. For systemic juvenile idiopathic arthritis (all requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, polyarticular juvenile idiopathic arthritis, systemic juvenile idiopathic arthritis, and giant cell arteritis: Prescribed by, or in consultation with, a rheumatology specialist. For systemic sclerosis-associated interstitial lung disease: Prescribed by, or in consultation with, a pulmonologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ACTIMMUNE 2000000UNIT/0.5ML INJ (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-alyq 20mg tab

- tadalafil 20mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Diagnosis confirmed by right heart catheterization. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ADEMPAS 0.5MG TAB
- ADEMPAS 1MG TAB
- ADEMPAS 2MG TAB

ADEMPAS 1.5MG TABADEMPAS 2.5MG TAB

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For all indications: Diagnosis confirmed by right heart catheterization. For pulmonary arterial hypertension: Both of the following were ineffective or not tolerated: one ERA (ambrisentan, bosentan or macitentan (Opsumit)) AND one PDE5-inhibitor (sildenafil or tadalafil). C) For persistent/recurrent chronic thromboembolic pulmonary hypertension (WHO Group 4): Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

Prior Authorization Criteria Last Updated 11/15/2024

Products Affected

- everolimus 10mg tab (New Starts Only)
- -everolimus 2mg tab for oral susp (New Starts Only)
- everolimus 5mg tab (New Starts Only)
- -everolimus 7.5mg tab (New Starts Only)
- torpenz 2.5mg tab (New Starts Only)
- torpenz 7.5mg tab (New Starts Only)

- -everolimus 2.5mg tab (New Starts Only)
- everolimus 3mg tab for oral susp (New Starts Only)
- everolimus 5mg tab for oral susp (New Starts Only)
- torpenz 10mg tab (New Starts Only)
- torpenz 5mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AJOVY 225MG/1.5ML AUTO-INJECTOR

- AJOVY 225MG/1.5ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Both of the following: A) Member has 4 or more migraine days per month for the previous 3 months or longer AND B) An 8-week or greater trial of two of the three following drug classes was ineffective or not tolerated: a) anticonvulsants, b) vasoactive agents, OR c) antidepressants. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AKEEGA 500-100MG TAB (New Starts Only)

- AKEEGA 500-50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ALECENSA 150MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-NITAZOXANIDE 500MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For diarrhea due to giardiasis: One of the following was ineffective or not tolerated: a) metronidazole OR b) tinidazole. For diarrhea due to cryptosporidiosis: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- PROLASTIN 1000MG INJ

-ZEMAIRA 1000MG INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | A) Diagnosis of congenital alpha1-antitrypsin deficiency is confirmed by both of the following: a) circulating baseline alpha1-antitrysin level is below the standard protective threshold (less than 11 micromol/L OR less than 50 mg per deciliter by nephelometry) AND b) high risk alpha1-antitrypsin deficiency genotype (SS, SZ, ZZ, or null/null) AND B) Prescriber attests that member does not have IgA deficiency with known anti-IgA antibody. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a pulmonologist |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ALUNBRIG 180MG TAB (New Starts Only)
- ALUNBRIG 90MG TAB (New Starts Only)

ALUNBRIG 30MG TAB (New Starts Only) ALUNBRIG TAB INITIATION PACK (30) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- APTIOM 200MG TAB (New Starts Only)

- APTIOM 600MG TAB (New Starts Only)

APTIOM 400MG TAB (New Starts Only)APTIOM 800MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) lamotrigine b) carbamazepine c) levetiracetam d) oxcarbazepine e) phenytoin f) topiramate OR g) lacosamide. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ARCALYST 220MG INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ARIKAYCE 590MG/8.4ML INH SUSP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, an infectious disease specialist or pulmonologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AUGTYRO 40MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AUSTEDO 12MG TAB
- -AUSTEDO 9MG TAB
- AUSTEDO XR 18MG TAB
- AUSTEDO XR 30MG TAB
- -AUSTEDO XR 42MG TAB
- AUSTEDO XR 6-12-24MG TAB TITRATION PACK (42)
- AUSTEDO XR TAB ONCE DAILY 4 WEEK TITRATION PACK

- AUSTEDO 6MG TAB
- AUSTEDO XR 12MG TAB
- -AUSTEDO XR 24MG TAB
- AUSTEDO XR 36MG TAB
- AUSTEDO XR 48MG TAB
- AUSTEDO XR 6MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For tardive dyskinesia (initial requests): A) One of the following: i) Member has failed to respond to a change in current antidopaminergic therapy OR ii) Member is unable to switch current antidopaminergic therapy OR iii) Member has symptoms of tardive dyskinesia and is not using antidopaminergic therapy AND B) Member has a functional disability due to tardive dyskinesia. For chorea associated with Huntington's disease (initial requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist or psychiatrist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AUVELITY 105-45MG ER TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) escitalopram, b) sertraline, c) fluoxetine, d) citalopram, e) paroxetine, f) fluvoxamine, g) bupropion, h) venlafaxine i) desvenlafaxine, or j) duloxetine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- AYVAKIT 100MG TAB (New Starts Only)
- AYVAKIT 25MG TAB (New Starts Only)
- AYVAKIT 50MG TAB (New Starts Only)

AYVAKIT 200MG TAB (New Starts Only)AYVAKIT 300MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- BALVERSA 3MG TAB (New Starts Only)

- BALVERSA 4MG TAB (New Starts Only)

-BALVERSA 5MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- rufinamide 200mg tab (New Starts Only)

- rufinamide 400mg tab (New Starts Only)

-rufinamide 40mg/ml oral susp (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of at least one anti-epileptic medication was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- BENLYSTA 200MG/ML AUTO-INJECTOR

- BENLYSTA 200MG/ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For systemic lupus erythematosus initial requests: Two of the following were ineffective or not tolerated: a) hydroxychloroquine, b) methotrexate, c) azathioprine, d) mycophenolate OR e) a corticosteroid. For all requests: Prescriber attests that member does not have severe active CNS lupus AND member is not taking other biologics. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a rheumatology specialist, nephrologist, or dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For lupus erythematosus initial therapy: Diagnosis of active systemic lupus erythematosus is confirmed by one of the following: A) anti-double stranded DNA value greater than 30 IU/mL OR B) low complement (C3/C4) OR C) positive for anti-Smith antibodies. For systemic lupus erythematosus (all requests): Will not be given in combination with other biologics. For active lupus nephritis (all requests): Will not be used in combination with voclosporin (Lupkynis). |

- BESREMI 500MCG/ML SYRINGE (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | One of the following was ineffective or not tolerated: A) hydroxyurea OR B) peginterferon alfa-2a. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -BOSULIF 100MG CAP (New Starts Only)
- -BOSULIF 400MG TAB (New Starts Only)
- -BOSULIF 50MG CAP (New Starts Only)

BOSULIF 100MG TAB (New Starts Only)BOSULIF 500MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-BRAFTOVI 75MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -BRIVIACT 100MG TAB (New Starts Only)
- BRIVIACT 10MG/ML ORAL SOLN (New Starts Only)
- -BRIVIACT 50MG TAB (New Starts Only)

BRIVIACT 10MG TAB (New Starts Only)
BRIVIACT 25MG TAB (New Starts Only)
BRIVIACT 75MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) lamotrigine b) carbamazepine c) levetiracetam d) oxcarbazepine e) phenytoin f) topiramate OR g) lacosamide. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- BRUKINSA 80MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- BYDUREON 2MG/0.85ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CABOMETYX 20MG TAB (New Starts Only)

- CABOMETYX 40MG TAB (New Starts Only)

- CABOMETYX 60MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- calcipotriene 0.005% cream

— calcipotriene 0.005% ointment

- CALCIPOTRIENE 0.005% TOPICAL SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CALQUENCE 100MG CAP (New Starts Only)

- CALQUENCE 100MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CAMZYOS 10MG CAP

-CAMZYOS 2.5MG CAP

- CAMZYOS 15MG CAP - CAMZYOS 5MG CAP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Member is symptomatic despite a maximally tolerated dose of one of the following: a) a beta blocker OR b) a non-dihydropyridine calcium channel blocker. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a cardiologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CAPLYTA 10.5MG CAP (New Starts Only)

- CAPLYTA 21MG CAP (New Starts Only)

- CAPLYTA 42MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For schizophrenia: Two of the following were ineffective or not tolerated: a) aripiprazole, b) olanzapine, c) quetiapine, d) risperidone, e) ziprasidone, or f) lurasidone. For bipolar depression: Both of the following were ineffective or not tolerated: a) lurasidone AND b) quetiapine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CAPRELSA 100MG TAB (New Starts Only)

- CAPRELSA 300MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-carglumic acid 200mg tab for oral susp

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CAYSTON 75MG/ML INH SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- tadalafil 2.5mg tab

— tadalafil 5mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

Prior Authorization Criteria Last Updated 11/15/2024

Products Affected

- CIMZIA 200MG INJ

- CIMZIA 200MG/ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel c) Rinvoq OR d) Xeljanz. For ankylosing spondylitis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Cosentyx d) Rinvoq OR e) Xeljanz. For psoriatic arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Cosentyx, d) Stelara, e) Otezla, f) Skyrizi, g) Tremfya, h) Rinvoq OR i) Xeljanz. For plaque psoriasis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Cosentyx, d) Skyrizi, e) Stelara, f) Tremfya OR g) Otezla. For Crohn's disease (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Stelara, c) Skyrizi, OR d) Rinvoq. For Non-radiographic axial spondyloarthritis (initial requests): Trial of two non-steroidal anti-inflammatory drugs (NSAIDs) was ineffective or not tolerated. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, psoriatic arthritis, non-radiographic axial spondyloarthritis or ankylosing spondylitis: Prescribed by, or in consultation, with a rheumatology specialist. For Crohn's disease: Prescribed by, or in consultation with, a gastroenterologist. For plaque psoriasis: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- COMETRIQ CAP 100MG DAILY DOSE PACK (56) (New Starts Only)

- COMETRIQ CAP 60MG DAILY DOSE PACK (84) (New Starts Only)

- COMETRIQ CAP 140MG DAILY DOSE PACK (112) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- COPIKTRA 15MG CAP (New Starts Only)

- COPIKTRA 25MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ivabradine 5mg tab

— ivabradine 7.5mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For adults (18 years and older), one of the following: A) Member is on a maximally tolerated dose of beta blocker OR B) Member has a history of intolerance, contraindication, or a hypersensitivity to beta blocker. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a cardiologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- COSENTYX 150MG/ML AUTO-INJECTOR

- COSENTYX 75MG/0.5ML SYRINGE

- COSENTYX 150MG/ML SYRINGE - COSENTYX UNOREADY 300MG/2ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For ankylosing spondylitis (initial requests): Trial of sulfasalazine was ineffective or not tolerated (Trial of sulfasalazine not required for AS with predominant axial involvement). For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin (trial of other agents not required for patients under 18 years of age). For non-radiographic axial spondyloarthritis (initial requests): Trial of two non-steroidal anti-inflammatory drugs (NSAIDs) was ineffective or not tolerated. For hidradenitis suppurativa (initial requests): Member must have both of the following: a) At least 3 cysts AND b) Trial of one oral antibiotic was ineffective or not tolerated. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For psoriatic arthritis, non-radiographic axial spondyloarthritis or ankylosing spondylitis: Prescribed by, or in consultation with, a rheumatology specialist. For plaque psoriasis and hidradenitis suppurativa: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- COTELLIC 20MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- CYSTADROPS 0.37% OPHTH SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-CYSTARAN 0.44% OPHTH SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

— pyrimethamine 25mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- DAURISMO 100MG TAB (New Starts Only)

- DAURISMO 25MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- metyrosine 250mg cap

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- DIACOMIT 250MG CAP (New Starts Only)

- DIACOMIT 500MG CAP (New Starts Only)

DIACOMIT 250MG POWDER FOR ORAL SUSP (New Starts Only) DIACOMIT 500MG POWDER FOR ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of at least one anti-epileptic medication was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- DIFICID 200MG TAB

- DIFICID 40MG/ML ORAL SUSP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of oral vancomycin was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 1 month. |
| Other Criteria | |

- DOJOLVI 100% ORAL SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Diagnosis of a long-chain fatty acid oxidation disorder confirmed by two or more of the following: a) newborn blood screening/acylcarnitine profile b) molecular or genetic test OR c) fibroblast test. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a medical geneticist or prescriber specializing in the treatment of long-chain fatty acid oxidation disorders. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- DOPTELET 20MG TAB

- DOPTELET TAB 40MG DAILY DOSE PACK (10)

- DOPTELET TAB 60MG DAILY DOSE PACK (15)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For thrombocytopenia with chronic liver disease and scheduled to undergo a procedure: Member has a platelet count from the prior two weeks that shows less than 50,000 platelets per microliter. For chronic immune thrombocytopenia initial requests: Both of the following: A) Relapsed or refractory to at least one prior treatment for chronic immune thrombocytopenia B) Platelet count less than 30,000 platelets per microliter. For chronic immune thrombocytopenia continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For chronic immune thrombocytopenia: Prescribed by, or in consultation with, a hematologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- DRIZALMA 20MG DR SPRINKLE CAP (New Starts Only)

- DRIZALMA 40MG DR SPRINKLE CAP (New Starts Only)

DRIZALMA 30MG DR SPRINKLE CAP (New Starts Only) DRIZALMA 60MG DR SPRINKLE CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For all diagnoses: Member is unable to swallow solid dosage forms of duloxetine (Cymbalta equivalent). For major depressive disorder and generalized anxiety disorder: One of the following was ineffective or not tolerated: 1) citalopram oral solution, 2) escitalopram oral solution, 3) fluoxetine oral solution, 4) paroxetine oral suspension, or 5) sertraline oral solution. For diabetic peripheral neuropathy and fibromyalgia: One of the following was ineffective or not tolerated: 1) gabapentin oral solution or 2) pregabalin oral solution. For chronic musculoskeletal pain: Trial of additional agents (other than duloxetine [Cymbalta equivalent]) not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-dronabinol 10mg cap

-dronabinol 5mg cap

-dronabinol 2.5mg cap

- DUPIXENT 100MG/0.67ML SYRINGE
- DUPIXENT 200MG/1.14ML SYRINGE
- DUPIXENT 300MG/2ML SYRINGE

DUPIXENT 200MG/1.14ML AUTO-INJECTOR DUPIXENT 300MG/2ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: For atopic dermatitis: Two of the following were ineffective or not tolerated: a) a medium to very high potency topical steroid, b) a topical calcineurin inhibitor OR c) an oral immunosuppressant (trial of other agents not required for patients under 2 years of age). For asthma: History, within the last year of at least 1 asthma exacerbation requiring one of the following, despite regular use of inhaled corticosteroids plus an additional controller(s): a) treatment with systemic corticosteroids, b) emergency department visit OR c) hospitalization. For nasal polyps: Both of the following were ineffective or not tolerated: a) an oral corticosteroid AND b) a nasal corticosteroid. For eosinophilic esophagitis: Trial of topical corticosteroid was ineffective or not tolerated. For prurigo nodularis: Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For atopic dermatitis: Prescribed by, or in consultation with, an allergist, immunologist, or dermatologist. For asthma: Prescribed by, or in consultation with, an allergist, pulmonologist, or immunologist. For nasal polyps: Prescribed by, or in consultation with, an allergist, immunologist, or otolaryngologist. For eosinophilic esophagitis: Prescribed by, or in consultation with, an allergist or gastroenterologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For initial requests: For atopic dermatitis: Member has moderate to severe atopic dermatitis defined as: 1) One of the following: a) body surface area involvement of 10 percent or more OR b) Chart documentation is provided of severity with involvement of the face, head, neck, hands, feet, groin, or intertriginous areas. AND 2) At least two (2) of the following: a) intractable pruritus (itching), b) cracking and oozing/bleeding of skin OR c) impaired activities of daily living. For asthma: One of the following: 1) Eosinophilic phenotype with baseline blood eosinophil concentration greater than or equal to 150 cells/microliter) OR 2) Oral corticosteroid-dependent asthma requiring daily doses of 5 mg or greater prednisone (or equivalent). For nasal polyps, both of the following: A) Bilateral nasal polyposis confirmed with sinus CT scan AND B) Prescriber attests to moderate to severe symptoms of nasal congestion, blockage, or obstruction (such as loss of smell, rhinorrhea, or facial pain). For eosinophilic esophagitis, both of the following: A) endoscopic biopsy with at least 15 |

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eosinophils per high-power field (hpf) AND B) symptoms of esophageal dysfunction (e.g. dysphagia). For prurigo nodularis: Both of the following apply: a) diagnosis has persisted for at least 6 weeks, AND b) at least 20 nodules present at baseline. For all indications (all requests): Will not be used in combination with another targeted immunomodulator product for the prescribed indication.

- EMGALITY 100MG/ML SYRINGE

- EMGALITY 120MG/ML SYRINGE

PA Criteria **Criteria Details** Covered Uses All FDA-approved indications not otherwise excluded from Part D. **Exclusion** Criteria Required Medical Info For migraine initial requests: Both of the following: A) Member has 4 or more migraine days per month for the previous 3 months or longer AND B) An 8-week or greater trial of two of the three following drug classes was ineffective or not tolerated: a) anticonvulsants, b) vasoactive agents, OR c) antidepressants. For episodic cluster headache prophylaxis initial requests: Trial of verapamil was ineffective or not tolerated. For continuation requests (all diagnoses): Member has benefited with use of this medication. Age Restrictions Prescriber Restriction **Coverage Duration** Approved for duration of 1 year. Other Criteria

- EMGALITY 120MG/ML AUTO-INJECTOR

- EMSAM 12MG/24HR PATCH (New Starts Only)

- EMSAM 6MG/24HR PATCH (New Starts Only)

- EMSAM 9MG/24HR PATCH (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) escitalopram, b) sertraline, c) fluoxetine, d) citalopram, e) paroxetine, f) fluvoxamine, g) bupropion, h) venlafaxine i) desvenlafaxine, or j) duloxetine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ENBREL 25MG/0.5ML INJ
- ENBREL 50MG/ML AUTO-INJECTOR
- ENBREL 50MG/ML SYRINGE

ENBREL 25MG/0.5ML SYRINGEENBREL 50MG/ML CARTRIDGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Trial of methotrexate at a dose of at least 20mg/week (or maximally tolerated dose) was ineffective or not tolerated. For juvenile idiopathic arthritis (initial requests): Trial of methotrexate at a dose of at least 15 mg/week required (or maximally tolerated dose) was ineffective or not tolerated. For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin (trial of other agents not required for patients under 18 years of age). For ankylosing spondylitis (AS)(initial requests): Trial of sulfasalazine was ineffective or not tolerated (Trial of sulfasalazine not required for AS with predominant axial involvement). For psoriatic arthritis (initial requests): One of the following was ineffective OR b) sulfasalazine. For juvenile psoriatic arthritis: Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, psoriatic arthritis, juvenile psoriatic arthritis, juvenile idiopathic arthritis or ankylosing spondylitis: Prescribed by, or in consultation with, a rheumatology specialist. For plaque psoriasis: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-glutamine 5000mg powder for oral soln

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Criteria 1 and 2 must be met or criteria 3 must be met: 1. Trial of a maximally tolerated hydroxyurea dose was ineffective or not tolerated. 2. Member has had at least 1 vaso-occlusive crisis in the prior 12 months, while on hydroxyurea (if applicable). 3. Prescriber is a hematologist. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a hematologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SOFOSBUVIR/VELPATASVIR 400-100MG TAB

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | 1) Current HCV-RNA titer is provided 2) Member has not had prior treatment with a direct-acting antiviral for current hepatitis C infection 3) One of the following: a) Member does not have cirrhosis or b) Member has compensated cirrhosis and one of the following: i) Does not have genotype 3 or ii) has genotype 3 but no NS5A resistance-associated substitution Y93H or c) Member has decompensated cirrhosis AND will receive weight-based ribavirin. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a gastroenterologist, hepatologist, infectious disease or transplant specialist. |
| Coverage Duration | Coverage duration of 12 to 24 weeks. Applied consistent with current AASLD-IDSA guidance. |
| Other Criteria | |

- EPIDIOLEX 100MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of at least one anti-epileptic medication was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- EPRONTIA 25MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of topiramate. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ERIVEDGE 150MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ERLEADA 240MG TAB (New Starts Only)

- ERLEADA 60MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For metastatic castration-sensitive prostate cancer: Trial of abiraterone was ineffective or not tolerated. For nonmetastatic castration-resistant prostate cancer: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-pirfenidone 267mg cap

-pirfenidone 801mg tab

-pirfenidone 267mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For idiopathic pulmonary fibrosis initial requests: Diagnosis confirmed by one of the following: 1) Surgical lung biopsy revealing histopathological pattern of unspecified interstitial pneumonia (UIP) 2) High-resolution computed tomography indicates definite UIP pattern 3) Both High-resolution computed tomography indicates possible UIP pattern AND surgical lung biopsy reveals a histopathological pattern of probable UIP. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- EVRYSDI 0.75MG/ML ORAL SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Documentation is provided of a genetic test confirming diagnosis of spinal muscular atrophy. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For all requests: Will not be used in combination with nusinersen (Spinraza). |

- FANAPT 10MG TAB (New Starts Only)
- FANAPT 1MG TAB (New Starts Only)
- FANAPT 4MG TAB (New Starts Only)
- FANAPT 8MG TAB (New Starts Only)

- FANAPT 12MG TAB (New Starts Only)
- FANAPT 2MG TAB (New Starts Only)
- FANAPT 6MG TAB (New Starts Only)
- FANAPT TAB TITRATION PACK (8) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) aripiprazole, b) olanzapine, c) quetiapine, d) risperidone, e) ziprasidone, or f) lurasidone. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FASENRA 10MG/0.5ML SYRINGE

- FASENRA 30MG/ML SYRINGE

PA Criteria **Criteria Details** Covered Uses All FDA-approved indications not otherwise excluded from Part D. **Exclusion** Criteria Required Medical Info For asthma (initial requests): History within the last year of at least 1 asthma exacerbation requiring one of following, despite regular use of inhaled corticosteroids plus an additional controller(s): a) treatment with systemic corticosteroids, b) emergency department visit OR c) hospitalization. For asthma (continuation requests): Member has benefited with use of this medication. Age Restrictions Prescriber Restriction For asthma: Prescribed by, or in consultation with, an allergist, immunologist, or pulmonologist. **Coverage Duration** Approved for duration of 1 year. For asthma (initial requests): Eosinophilic phenotype with baseline blood eosinophil concentration greater than or equal to Other Criteria 150 cells/microliter. For asthma (all requests): Will not be used in combination with another targeted immunomodulator product for the prescribed indication.

- FASENRA 30MG/ML AUTO-INJECTOR

- FETZIMA 120MG ER CAP (New Starts Only)

- FETZIMA 40MG ER CAP (New Starts Only)

- FETZIMA ER CAP TITRATION PACK (28) (New Starts Only)

FETZIMA 20MG ER CAP (New Starts Only)FETZIMA 80MG ER CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) escitalopram, b) sertraline, c) fluoxetine, d) citalopram, e) paroxetine, f) fluvoxamine, g) bupropion, h) venlafaxine i) desvenlafaxine, or j) duloxetine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FINTEPLA 2.2MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of at least one anti-epileptic medication was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FIRDAPSE 10MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | Diagnosis of Lambert-Eaton myasthenic syndrome (LEMS) confirmed by one of the following: a) Presence of voltage-gated calcium channel antibodies OR b) electrophysiologic compound muscle action potential test findings are consistent with LEMS. |

- FIRMAGON 120MG INJ (New Starts Only)

- FIRMAGON 80MG INJ (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FOTIVDA 0.89MG CAP (New Starts Only)

- FOTIVDA 1.34MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FRUZAQLA 1MG CAP (New Starts Only)

- FRUZAQLA 5MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- FYCOMPA 0.5MG/ML ORAL SUSP (New Starts Only)
- -FYCOMPA 12MG TAB (New Starts Only)
- FYCOMPA 4MG TAB (New Starts Only)
- FYCOMPA 8MG TAB (New Starts Only)

FYCOMPA 10MG TAB (New Starts Only)
FYCOMPA 2MG TAB (New Starts Only)
FYCOMPA 6MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For partial-onset seizures: Two of the following were ineffective or not tolerated: a) lamotrigine b) carbamazepine c) levetiracetam d) oxcarbazepine e) phenytoin f) topiramate OR g) lacosamide. For primary generalized tonic-clonic seizures: Two of the following were ineffective or not tolerated: a) lamotrigine, b) levetiracetam, c) primidone OR d) topiramate. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- GATTEX 5MG INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is dependent on parenteral support for at least 12 months and at least 3 days per week. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- GAVRETO 100MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- GILOTRIF 20MG TAB (New Starts Only)

- GILOTRIF 30MG TAB (New Starts Only)

- GILOTRIF 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- NORDITROPIN 10MG/1.5ML PEN INJ
- NORDITROPIN 30MG/3ML PEN INJ
- OMNITROPE 10MG/1.5ML CARTRIDGE
- OMNITROPE 5MG/1.5ML CARTRIDGE
- SOGROYA 15MG/1.5ML PEN INJ

- NORDITROPIN 15MG/1.5ML PEN INJ
- NORDITROPIN 5MG/1.5ML PEN INJ
- OMNITROPE 5.8MG INJ
- SOGROYA 10MG/1.5ML PEN INJ
- SOGROYA 5MG/1.5ML PEN INJ

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Documentation is provided of failure to stimulate growth hormone secretion (peak growth hormone level of 10mcg/L or less) by one of the acceptable provocative tests. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- HADLIMA 40MG/0.4ML AUTO-INJECTOR

- HADLIMA 40MG/0.8ML AUTO-INJECTOR

HADLIMA 40MG/0.4ML SYRINGE HADLIMA 40MG/0.8ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Trial of methotrexate at a dose of at least 20mg/week (or maximally tolerated dose) was ineffective or not tolerated. For juvenile idiopathic arthritis (initial requests): Trial of methotrexate at a dose of at least 15 mg/week (or maximally tolerated dose) was ineffective or not tolerated. For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin. For ankylosing spondylitis (AS)(initial requests): Trial of sulfasalazine was ineffective or not tolerated (Trial of sulfasalazine not required for AS with predominant axial involvement). For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For ulcerative colitis or Crohn's disease (all requests): Trial of other agents not required. For hidradenitis suppurativa (initial requests): Member must have both of the following: a) At least 3 cysts AND b) Trial of one oral antibiotic was ineffective or not tolerated. For uveitis (initial requests): Both of the following were ineffective or not tolerated: a) a corticosteroid AND b) an immunosuppressant (methotrexate, mycophenolate mofetil, or cyclosporine). For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, psoriatic arthritis, juvenile idiopathic arthritis or ankylosing spondylitis: Prescribed by, or in consultation with, a rheumatology specialist. For plaque psoriasis and hidradenitis suppurativa: Prescribed by, or in consultation with, a dermatologist. For Crohn's disease and ulcerative colitis: Prescribed by, or in consultation with, a gastroenterologist. For uveitis: Prescribed by, or in consult with, a rheumatology specialist OR ophthalmologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- BERINERT 500UNIT INJ
- -HAEGARDA 3000UNIT INJ
- sajazir 30mg/3ml syringe
- TAKHZYRO 300MG/2ML SYRINGE

- HAEGARDA 2000UNIT INJ
- *icatibant 10mg/ml syringe*
- TAKHZYRO 300MG/2ML INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For medications indicated for long-term prophylaxis (all requests): Will not be used in combination with another agent for long-term prophylaxis of hereditary angioedema attacks. |

- IBRANCE 100MG CAP (New Starts Only)
- IBRANCE 125MG CAP (New Starts Only)
- IBRANCE 75MG CAP (New Starts Only)

– IBRANCE 100MG TAB (New Starts Only)
– IBRANCE 125MG TAB (New Starts Only)
– IBRANCE 75MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Intolerance or contraindication to therapy with both of the following: a) Verzenio AND b) Kisqali. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ICLUSIG 10MG TAB (New Starts Only)
- ICLUSIG 30MG TAB (New Starts Only)

ICLUSIG 15MG TAB (New Starts Only)ICLUSIG 45MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- IDHIFA 100MG TAB (New Starts Only)

- IDHIFA 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ARISTADA 1064MG/3.9ML SYRINGE (New Starts Only)
- ARISTADA 662MG/2.4ML SYRINGE (New Starts Only)
- ARISTADA 882MG/3.2ML SYRINGE (New Starts Only)
- INVEGA HAFYERA 1560MG/5ML SYRINGE (New Starts Only)
- INVEGA SUSTENNA 156MG/ML SYRINGE (New Starts Only)
- INVEGA SUSTENNA 39MG/0.25ML SYRINGE (New Starts Only)
- INVEGA TRINZA 273MG/0.875ML SYRINGE (New Starts Only)
- INVEGA TRINZA 546MG/1.75ML SYRINGE (New Starts Only)
- risperidone 37.5mg inj (New Starts Only)
- risperidone microspheres 12.5mg inj (New Starts Only)

- ARISTADA 441MG/1.6ML SYRINGE (New Starts Only)
- ARISTADA 675MG/2.4ML SYRINGE (New Starts Only)
- INVEGA HAFYERA 1092MG/3.5ML SYRINGE (New Starts Only)
- INVEGA SUSTENNA 117MG/0.75ML SYRINGE (New Starts Only)
- INVEGA SUSTENNA 234MG/1.5ML SYRINGE (New Starts Only)
- INVEGA SUSTENNA 78MG/0.5ML SYRINGE (New Starts Only)
- INVEGA TRINZA 410MG/1.315ML SYRINGE (New Starts Only)
- INVEGA TRINZA 819MG/2.625ML SYRINGE (New Starts Only)
- risperidone 50mg inj (New Starts Only)
- risperidone microspheres 25mg inj (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Patient has established tolerability with the oral version of medication being requested. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- IMBRUVICA 140MG CAP (New Starts Only)
- IMBRUVICA 70MG CAP (New Starts Only)

– IMBRUVICA 420MG TAB (New Starts Only) – IMBRUVICA 70MG/ML ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- INCRELEX 40MG/4ML INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- INGREZZA 40MG CAP
- INGREZZA 60MG CAP
- INGREZZA 80MG CAP
- INGREZZA CAP THERAPY PACK (28)

– INGREZZA 40MG SPRINKLE CAP – INGREZZA 60MG SPRINKLE CAP – INGREZZA 80MG SPRINKLE CAP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For tardive dyskinesia (initial requests): A) One of the following: i) Member has failed to respond to a change in current antidopaminergic therapy OR ii) Member is unable to switch current antidopaminergic therapy OR iii) Member has symptoms of tardive dyskinesia and is not using antidopaminergic therapy AND B) Member has a functional disability due to tardive dyskinesia. For chorea associated with Huntington's disease (initial requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist or psychiatrist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- INLYTA 1MG TAB (New Starts Only)

- INLYTA 5MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- INQOVI 35-100MG TAB PACK (5) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- INREBIC 100MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of Jakafi was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- gefitinib 250mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

— ivermectin 3mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 1 month. |
| Other Criteria | |

- GAMUNEX 1GM/10ML INJ

- OCTAGAM 2GM/20ML INJ

OCTAGAM 1GM/20ML INJPRIVIGEN 20GM/200ML INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | Approval will be based off BvD coverage determination. |

- IWILFIN 192MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- deferasirox 180mg tab

- deferasirox 90mg tab

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaRequired Medical InfoAge RestrictionsPrescriber RestrictionPrescribed by or in consultation with a hematologistCoverage DurationApproved for duration of 1 year.Other Criteria

- deferasirox 360mg tab

- JAKAFI 10MG TAB (New Starts Only)
- JAKAFI 20MG TAB (New Starts Only)
- JAKAFI 5MG TAB (New Starts Only)

JAKAFI 15MG TAB (New Starts Only)JAKAFI 25MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- JAYPIRCA 100MG TAB (New Starts Only)

- JAYPIRCA 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- JYLAMVO 2MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of methotrexate. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | Approval will be based off BvD coverage determination. |

- KALYDECO 13.4MG ORAL GRANULES
- KALYDECO 25MG ORAL GRANULES
- KALYDECO 50MG ORAL GRANULES

KALYDECO 150MG TAB KALYDECO 5.8MG ORAL GRANULES KALYDECO 75MG ORAL GRANULES

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-KERENDIA 10MG TAB

- KERENDIA 20MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- KEVZARA 150MG/1.14ML AUTO-INJECTOR

- KEVZARA 200MG/1.14ML AUTO-INJECTOR

KEVZARA 150MG/1.14ML SYRINGEKEVZARA 200MG/1.14ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Rinvoq OR d) Xeljanz. For polymyalgia rheumatica (initial requests), one of the following: a) a trial of a corticosteroid was ineffective OR b) member was unable to tolerate a corticosteroid taper to less than or equal to 5 mg prednisone equivalent per day. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis and polymyalgia rheumatica: Prescribed by, or in consultation with, a rheumatology specialist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -KISQALI TAB 200MG DAILY DOSE PACK (21) (New Starts Only)
- -KISQALI TAB 600MG DAILY DOSE PACK (63) (New Starts Only)
- KISQALI/FEMARA 400 CO-PACK (70) (New Starts Only)

- -KISQALI TAB 400MG DAILY DOSE PACK (42) (New Starts Only)
- -KISQALI/FEMARA 200 CO-PACK (49) (New Starts Only)
- -KISQALI/FEMARA 600 CO-PACK (91) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- mifepristone 300mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- KOSELUGO 10MG CAP (New Starts Only)

-KOSELUGO 25MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Chart notes documentation is provided that indicates inoperable and symptomatic disease |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- KRAZATI 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- javygtor 100mg powder for oral soln
- javygtor 500mg powder for oral soln
- sapropterin 100mg tab

- *javygtor 100mg tab*
- -sapropterin 100mg powder for oral soln
- sapropterin 500mg powder for oral soln

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For continuation therapy: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a medical geneticist or metabolic physician. |
| Coverage Duration | Initial approval of 3 months. Continuing therapy approved for 1 year. |
| Other Criteria | |

- LENVIMA 10MG DAILY DOSE PACK (30) (New Starts Only)
- LENVIMA 14MG DAILY DOSE PACK (60) (New Starts Only)
- LENVIMA 20MG DAILY DOSE PACK (60) (New Starts Only)
- LENVIMA 4MG DAILY DOSE PACK (30) (New Starts Only)

- LENVIMA 12MG DAILY DOSE PACK (90) (New Starts Only)
- LENVIMA 18MG DAILY DOSE PACK (90) (New Starts Only)
- LENVIMA 24MG DAILY DOSE PACK (90) (New Starts Only)
- LENVIMA 8MG DAILY DOSE PACK (60) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ambrisentan 10mg tab

-ambrisentan 5mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Diagnosis confirmed by right heart catheterization. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LIBERVANT 10MG BUCCAL FILM (New Starts Only)
- LIBERVANT 15MG BUCCAL FILM (New Starts Only)
- -LIBERVANT 7.5MG BUCCAL FILM (New Starts Only)

-LIBERVANT 12.5MG BUCCAL FILM (New Starts Only)

-LIBERVANT 5MG BUCCAL FILM (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

— lidocaine 5% patch

— tridacaine 5% patch

— lidocan 5% patch

| PA Criteria | Criteria Details |
|------------------------|-------------------------------------|
| Covered Uses | All Medically-accepted Indications. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

— lidocaine 5% ointment

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-LITFULO 50MG CAP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For alopecia areata (initial requests): Hair loss impacts 50% or greater of the scalp. For alopecia areata (continuation requests): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-LIVTENCITY 200MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Prescriber attests that the medication will not be used for CMV infection prophylaxis. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a hematologist, oncologist, transplant or infectious disease specialist. |
| Coverage Duration | Approved for 3 months. |
| Other Criteria | |

- LOKELMA 10GM POWDER FOR ORAL SUSP

- LOKELMA 5GM POWDER FOR ORAL SUSP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Member has baseline persistent potassium level greater than 5.0 mmol/L. For continuing requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a nephrologist, cardiologist, hematologist or endocrinologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LONSURF 6.14-15MG TAB (New Starts Only)

-LONSURF 8.19-20MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LORBRENA 100MG TAB (New Starts Only)

-LORBRENA 25MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LUMAKRAS 120MG TAB (New Starts Only)

-LUMAKRAS 320MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LUMRYZ 4.5GM GRANULES FOR ORAL SUSP

-LUMRYZ 7.5GM GRANULES FOR ORAL SUSP

LUMRYZ 6GM GRANULES FOR ORAL SUSP LUMRYZ 9GM GRANULES FOR ORAL SUSP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For excessive daytime sleepiness with narcolepsy in adults: Both of the following were ineffective or not tolerated: a) Sunosi AND b) either modafinil or armodafinil. For cataplexy with narcolepsy: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For excessive daytime sleepiness with narcolepsy: Documentation is provided of full nocturnal polysomnogram used to confirm diagnosis. For cataplexy with narcolepsy: Documentation is provided of one of the following to confirm diagnosis: a) full nocturnal polysomnogram OR b) low cerebrospinal fluid orexin-A concentration. |

-LUPKYNIS 7.9MG CAP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For continuation therapy: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a rheumatology specialist or nephrologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For all requests: Will not be used in combination with belimumab (Benlysta). |

- LYNPARZA 100MG TAB (New Starts Only)

-LYNPARZA 150MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LYTGOBI TAB 12MG DAILEY DOSE PACK (21) (New Starts Only)

-LYTGOBI TAB 20MG DAILEY DOSE PACK (35) (New Starts Only)

- LYTGOBI TAB 16MG DAILEY DOSE PACK (28) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

Prior Authorization Criteria Last Updated 11/15/2024

Products Affected

- MAVYRET 100-40MG TAB

- MAVYRET 50-20MG ORAL PELLET

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | 1) Current HCV-RNA titer is provided 2) Member does not have decompensated cirrhosis 3) One of the following: a) member has not had prior treatment with a direct-acting antiviral for current hepatitis C infection or b) prior treatment with sofosbuvir-based regimen and all of the following: i) Member does not have genotype 3 and ii) No prior treatment with an NS3/4A protease inhibitor. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a gastroenterologist, hepatologist, infectious disease or transplant specialist. |
| Coverage Duration | Coverage duration of 8 to 16 weeks. Applied consistent with current AASLD-IDSA guidance. |
| Other Criteria | |

- MEGESTROL ACETATE 125MG/ML SUSP

- megestrol acetate 40mg/ml oral susp

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- megestrol acetate 20mg tab (New Starts Only)

- megestrol acetate 40mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- MEKINIST 0.05MG/ML ORAL SOLN (New Starts Only)

- MEKINIST 0.5MG TAB (New Starts Only)

- MEKINIST 2MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- MEKTOVI 15MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- dihydroergotamine mesylate 0.5mg/act nasal inhaler

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of two different triptans was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- MOUNJARO 10MG/0.5ML AUTO-INJECTOR
- MOUNJARO 15MG/0.5ML AUTO-INJECTOR
- MOUNJARO 5MG/0.5ML AUTO-INJECTOR

MOUNJARO 12.5MG/0.5ML AUTO-INJECTOR MOUNJARO 2.5MG/0.5ML AUTO-INJECTOR MOUNJARO 7.5MG/0.5ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- MOVANTIK 12.5MG TAB

- MOVANTIK 25MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -carisoprodol 350mg tab
- -cyclobenzaprine 10mg tab
- metaxalone 800mg tab
- methocarbamol 750mg tab

- chlorzoxazone 500mg tab
- *cyclobenzaprine 5mg tab*
- methocarbamol 500mg tab
- orphenadrine citrate 100mg er tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | Prior Authorization applies to patients 65 years or older. |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

Prior Authorization Criteria Last Updated 11/15/2024

Products Affected

- ABELCET 5MG/ML INJ
- acetylcysteine 200mg/ml inh soln
- albuterol 0.21mg/ml (0.63mg/3ml) inh soln
- albuterol 1.25mg/3ml neb soln
- AMPHOTERICIN B 50MG INJ
- aprepitant 125mg/80mg cap therapy pack (3)
- -aprepitant 80mg cap
- -azathioprine 50mg tab
- budesonide 0.5mg/2ml inh susp
- CLINIMIX 4.25/10 INJ
- CLINIMIX 5/15 INJ
- -clinisol 15% inj
- CYCLOPHOSPHAMIDE 50MG TAB
- -cyclosporine 25mg cap
- cyclosporine modified 100mg/ml oral soln
- cyclosporine modified 50mg cap
- ENGERIX-B 10MCG/0.5ML SYRINGE
- ENGERIX-B 20MCG/ML SYRINGE
- ENVARSUS XR 1MG TAB
- -everolimus 0.25mg tab
- -everolimus 0.75mg tab
- -gengraf 100mg cap
- -gengraf 25mg cap
- GLUCOSE 100MG/ML/SODIUM CHLORIDE 2MG/ML INJ
- -granisetron 1mg tab
- HUMULIN R 500UNIT/ML INJ
- INSULIN LISPRO 100UNIT/ML INJ
- ipratropium bromide 0.02% inh soln
- -levalbuterol 0.31mg/3ml neb soln

- -acetylcysteine 100mg/ml inh soln
- acyclovir 50mg/ml inj
- albuterol 0.83mg/ml (0.083%) inh soln
- albuterol 5mg/ml (0.5%) inh soln
- -aprepitant 125mg cap
- -aprepitant 40mg cap
- arformoterol tartrate 15mcg/2ml neb soln
- budesonide 0.25mg/2ml inh susp
- -budesonide 1mg/2ml inh susp
- CLINIMIX 4.25/5 INJ
- CLINIMIX 5/20 INJ
- CYCLOPHOSPHAMIDE 25MG TAB
- -cyclosporine 100mg cap
- cyclosporine modified 100mg cap
- cyclosporine modified 25mg cap
- DIPHTHERIA/TETANUS TOXOID INJ
- ENGERIX-B 20MCG/ML INJ
- ENVARSUS XR 0.75MG TAB
- ENVARSUS XR 4MG TAB
- -everolimus 0.5mg tab
- -everolimus 1mg tab
- -gengraf 100mg/ml oral soln
- -glucose 100mg/ml inj
- GLUCOSE 100MG/ML/SODIUM CHLORIDE 4.5MG/ML INJ
- HEPLISAV-B 20MCG/0.5ML SYRINGE
- IMOVAX 2.5UNIT/ML INJ
- INTRALIPID 20GM/100ML INJ
- ipratropium/albuterol 0.5-2.5mg/3ml inh soln
- levalbuterol 0.63mg/3ml inh soln

| | Last Opaalea 11/15/2024 |
|---|--|
| - levalbuterol 1.25mg/3ml neb soln | <i>— methylprednisolone 16mg tab</i> |
| - methylprednisolone 32mg tab | <i>— methylprednisolone 4mg tab</i> |
| – methylprednisolone 8mg tab | — mycophenolate mofetil 200mg/ml oral susp |
| — mycophenolate mofetil 250mg cap | - mycophenolate mofetil 500mg tab |
| - mycophenolic acid 180mg dr tab | — mycophenolic acid 360mg dr tab |
| - NUTRILIPID 20GM/100ML INJ | — ondansetron 0.8mg/ml oral soln |
| <i>— ondansetron 4mg odt</i> | <i>— ondansetron 4mg tab</i> |
| - ondansetron 8mg odt | - ondansetron 8mg tab |
| <i>— pentamidine isethionate 300mg/6ml inh soln</i> | — plenamine 15% inj |
| — prednisolone 1mg/ml oral soln | - prednisolone 3mg/ml oral soln |
| — prednisolone 5mg/ml oral soln | - prednisone 10mg tab |
| -prednisone 1mg tab | - PREDNISONE 1MG/ML ORAL SOLN |
| -prednisone 2.5mg tab | -prednisone 20mg tab |
| -prednisone 50mg tab | -prednisone 5mg tab |
| – PREHEVBRIO 10MCG/ML INJ | - PROGRAF 0.2MG GRANULES FOR ORAL SUSP |
| - PROGRAF 1MG GRANULES FOR ORAL SUSP | – PROSOL 20% INJ |
| – PULMOZYME 1MG/ML INH SOLN | - RABAVERT 2.5UNIT/ML INJ |
| - RECOMBIVAX 10MCG/ML INJ | - RECOMBIVAX 10MCG/ML SYRINGE |
| - RECOMBIVAX 40MCG/ML INJ | - RECOMBIVAX 5MCG/0.5ML INJ |
| - RECOMBIVAX 5MCG/0.5ML SYRINGE | — sirolimus 0.5mg tab |
| – sirolimus 1mg tab | — sirolimus 1mg/ml oral soln |
| — sirolimus 2mg tab | — tacrolimus 0.5mg cap |
| - tacrolimus 1mg cap | — tacrolimus 5mg cap |
| - TDVAX 4-4UNIT/ML INJ | - TENIVAC 4-10UNIT/ML INJ |
| - TENIVAC 4-10UNIT/ML SYRINGE | - TPN ELECTROLYTES INJ |
| - TRAVASOL 10% INI | |

- TRAVASOL 10% INJ

| PA Criteria | Criteria Details |
|--------------|--|
| Covered Uses | This drug may be covered under Medicare Part B or D depending upon the circumstances. Information may need to be submitted describing the use and setting of the drug to make the determination. |
| | |

| | Prior Authorization Criteria Last Updated 11/15/2024 |
|------------------------|---|
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | |
| Other Criteria | |

- NERLYNX 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- sorafenib 200mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- NEXLETOL 180MG TAB

-NEXLIZET 180-10MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- NINLARO 2.3MG CAP (New Starts Only)

- NINLARO 3MG CAP (New Starts Only)

- NINLARO 4MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- droxidopa 100mg cap

- droxidopa 300mg cap

- droxidopa 200mg cap

-posaconazole 100mg dr tab

-posaconazole 40mg/ml oral susp

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- NUBEQA 300MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For metastatic hormone-sensitive prostate cancer: Trial of abiraterone was ineffective or not tolerated. For non-metastatic castration-resistant prostate cancer: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

Prior Authorization Criteria Last Updated 11/15/2024

Products Affected

- NUCALA 100MG INJ

- NUCALA 100MG/ML SYRINGE

NUCALA 100MG/ML AUTO-INJECTOR NUCALA 40MG/0.4ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For asthma initial requests: History within the last year of at least 1 asthma exacerbation requiring one of the following, despite regular use of inhaled corticosteroids plus an additional controller(s): a) treatment with systemic corticosteroids, b) emergency department visit OR c) hospitalization. For eosinophilic granulomatosis with polyangiitis (EGPA) initial requests: All of the following: A) One of the following: 1) baseline blood eosinophil count greater than 1000 cells per microliter OR 2) baseline blood eosinophil count greater than 10% of the total leukocyte count B) Trial of oral corticosteroid therapy was ineffective or not tolerated C) One of the following was ineffective or not tolerated: a) cyclophosphamide OR b) methotrexate. For hypereosinophilic syndrome (HES) initial requests: Both of the following: A) Diagnosis confirmed by blood eosinophil count greater than 1000 cells per microliter AND B) Hypereosinophilic syndrome has persisted for at least six months. For nasal polyps initial requests: Both of the following were ineffective or not tolerated: a) an oral corticosteroid AND b) a nasal corticosteroid. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with: For asthma: an allergist, pulmonologist, or immunologist. For nasal polyps: an allergist, immunologist, or otolaryngologist. For EGPA: a rheumatology specialist, allergist, pulmonologist, pulmonologist, or immunologist. For HES: a rheumatology specialist, allergist, pulmonologist, gastroenterologist, hematologist, or other specialist experienced in the diagnosis and treatment of HES |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For asthma (initial requests): Eosinophilic phenotype with baseline blood eosinophil concentration greater than or equal to 150 cells/microliter. For asthma (all requests): Will not be used in combination with another targeted immunomodulator product for the prescribed indication. |

- NUEDEXTA 20-10MG CAP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: A) Documentation is provided (in the form of chart notes or imaging) of structural neurological condition as the cause of pseudobulbar affect. For continuation requests, both of the following: A) Documentation is provided (in the form of chart notes or imaging) of structural neurological condition as the cause of pseudobulbar affect AND B) Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- NUPLAZID 10MG TAB (New Starts Only)

- NUPLAZID 34MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-armodafinil 150mg tab

-armodafinil 250mg tab

armodafinil 200mg tab
armodafinil 50mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- octreotide 0.05mg/ml inj
- *octreotide 0.2mg/ml inj*
- octreotide 1mg/ml inj

- octreotide 0.1mg/ml inj - octreotide 0.5mg/ml inj

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ODOMZO 200MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OFEV 100MG CAP

-OFEV 150MG CAP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | 1) For idiopathic pulmonary fibrosis initial requests: A) Diagnosis confirmed by one of the following: i) Surgical lung biopsy revealing histopathological pattern of unspecified interstitial pneumonia (UIP) ii) High-resolution computed tomography (HRCT) indicates definite UIP pattern iii) Both HRCT indicates possible UIP pattern AND surgical lung biopsy reveals a histopathological pattern of probable UIP AND B) Trial of pirfenidone was ineffective or not tolerated. 2) For systemic sclerosis-associated interstitial lung disease (ILD) initial requests: A) Diagnosis confirmed with documentation provided of both of the following: i) HRCT scan AND ii) pulmonary function tests AND B) Trial of mycophenolate mofetil was ineffective or not tolerated. 3) For chronic fibrosing ILDs with a progressive phenotype initial requests: A) Disease is progressive, defined by one of the following: a) forced vital capacity (FVC) decline of 5% or more OR b) corrected hemoglobin decline of 10% or more OR iii) radiological evidence of disease progression AND B) Progression occurred despite treatment with one of the following: i) azathioprine ii) cyclosporine iii) mycophenolate mofetil iv) tacrolimus v) oral corticosteroids equivalent to 20 mg or more per day of prednisone vi) cyclophosphamide vii) rituximab. 4) For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OGSIVEO 100MG TAB 7-DAY PACK (14) (New Starts Only)

- OGSIVEO 150MG TAB 7-DAY PACK (14) (New Starts Only)

- OGSIVEO 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OJEMDA 100MG TAB (New Starts Only)

- OJEMDA 100MG TAB PACK (600MG ONCE WEEKLY) (24) (New Star - OJEMDA 25MG/ML POWDER FOR ORAL SUSP (New Starts Only)

OJEMDA 100MG TAB PACK (400MG ONCE WEEKLY) (16) (New Star OJEMDA 25MG/ML POWDER FOR ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OJJAARA 100MG TAB (New Starts Only)

- OJJAARA 150MG TAB (New Starts Only)

- OJJAARA 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OLUMIANT 1MG TAB

- OLUMIANT 4MG TAB

- OLUMIANT 2MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Rinvoq OR d) Xeljanz. For alopecia areata (initial requests): Hair loss impacts 50% or greater of the scalp. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis: Prescribed by or in consultation with, a rheumatology specialist. For alopecia areata: Prescribed by or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ONUREG 200MG TAB (New Starts Only)

- ONUREG 300MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OPSUMIT 10MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Diagnosis confirmed by right heart catheterization. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-fentanyl 1200mcg lozenge

- -fentanyl 200mcg lozenge
- -fentanyl 600mcg lozenge

-fentanyl 1600mcg lozenge -fentanyl 400mcg lozenge -fentanyl 800mcg lozenge

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Documented tolerance to opioids defined as patients taking around the clock medicine consisting of at least 60mg of oral morphine daily, at least 25mcg of transdermal fentanyl per hour, at least 30mg of oxycodone daily, at least 8mg of oral hydromorphone daily, or an equianalgesic dose of another opioid daily for a week or longer. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ORENCIA 125MG/ML AUTO-INJECTOR

- ORENCIA 50MG/0.4ML SYRINGE

ORENCIA 125MG/ML SYRINGE ORENCIA 87.5MG/0.7ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Enbrel, b) Hadlima, c) Rinvoq OR d) Xeljanz. For polyarticular juvenile idiopathic arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel OR c) Xeljanz. For adult psoriatic arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel, c) Cosentyx, d) Stelara, e) Otezla, f) Skyrizi, g) Tremfya, h) Rinvoq OR i) Xeljanz. For pediatric psoriatic arthritis (initial requests): Trial of Enbrel was ineffective or not tolerated. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, polyarticular juvenile idiopathic arthritis, and psoriatic arthritis (adult and pediatric): Prescribed by, or in consultation with a rheumatology specialist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ORGOVYX 120MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ORKAMBI 125-100MG ORAL GRANULES
- ORKAMBI 125-200MG TAB
- ORKAMBI 94-75MG ORAL GRANULES

- ORKAMBI 125-100MG TAB - ORKAMBI 188-150MG ORAL GRANULES

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ORSERDU 345MG TAB (New Starts Only)

- ORSERDU 86MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- OTEZLA 20MG TAB

- OTEZLA 30MG TAB

- OTEZLA TAB 28-DAY STARTER PACK (55)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For oral ulcers associated with Behcet's disease (initial requests): Trial of topical triamcinolone 0.1% oral paste was ineffective or not tolerated. For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of 10mg/week (or maximally tolerated dose) OR b) acitretin (trial of other agents not required for patients under 18 years of age). For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For oral ulcers associated with Behcet's disease and psoriatic arthritis: Prescribed by, or in consultation with, a rheumatology specialist. For Plaque Psoriasis: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For oral ulcers associated with Behcet's disease (initial requests): Diagnosis confirmed by the presence of oral ulcers AND at least two of the following: recurrent genital ulceration, eye lesions, skin lesions, positive pathergy test. For psoriatic arthritis and plaque psoriasis (all requests): Will not be used in combination with biologic therapy for the prescribed indication. |

- OXBRYTA 300MG TAB

-OXBRYTA 500MG TAB

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Criteria 1 and 2 must be met or criteria 3 must be met: 1. Trial of a maximally tolerated hydroxyurea dose was ineffective or not tolerated. 2. Member has had at least 1 vaso-occlusive crisis in the prior 12 months, while on hydroxyurea (if applicable). 3. Prescriber is a hematologist. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a hematologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For all requests: Will not be used in combination with crizanlizumab (Adakveo). |

- OXBRYTA 300MG TAB FOR ORAL SUSP

- OZEMPIC 2.68MG/ML PEN INJ

- OZEMPIC 4MG/3ML PEN INJ

PA Criteria Criteria Details Covered Uses All FDA-approved indications not otherwise excluded from Part D. **Exclusion** Criteria Required Medical Info Age Restrictions Prescriber Restriction **Coverage Duration** Approved for duration of 1 year. Other Criteria

- OZEMPIC 2MG/3ML PEN INJ

- PANRETIN 0.1% GEL (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- PEMAZYRE 13.5MG TAB (New Starts Only)

- PEMAZYRE 4.5MG TAB (New Starts Only)

- PEMAZYRE 9MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- PIQRAY TAB 200MG DAILY DOSE PACK (28) (New Starts Only)

- PIQRAY TAB 250MG DAILY DOSE PACK (56) (New Starts Only)

- PIQRAY TAB 300MG DAILY DOSE PACK (56) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- POMALYST 1MG CAP (New Starts Only)

- POMALYST 3MG CAP (New Starts Only)

POMALYST 2MG CAP (New Starts Only)POMALYST 4MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- PREVYMIS 240MG TAB

-PREVYMIS 480MG TAB

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member will/has initiated Prevymis within 30 days after an allogeneic hematopoietic stem cell transplant or 7 days after kidney transplant. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a hematologist, oncologist, transplant or infectious disease specialist. |
| Coverage Duration | Approved for 8 months for hematopoietic stem cell transplant or 8 months for kidney transplant. |
| Other Criteria | |

- PROMACTA 12.5MG POWDER FOR ORAL SUSP
- PROMACTA 25MG POWDER FOR ORAL SUSP
- PROMACTA 50MG TAB

PROMACTA 12.5MG TAB
PROMACTA 25MG TAB
PROMACTA 75MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- modafinil 100mg tab

- modafinil 200mg tab

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications, Some Medically-Accepted Indications |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- PURIXAN 2000MG/100ML ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of mercaptopurine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-QINLOCK 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-quinine sulfate 324mg cap

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 1 month. |
| Other Criteria | |

- RADICAVA 105MG/5ML ORAL SUSP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Member has a score of two or greater for each individual item on the Amyotrophic Lateral Sclerosis Functional Rating Scale-Revised (ALSFRS-R). For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- REGRANEX 0.01% GEL

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- REPATHA 140MG/ML AUTO-INJECTOR

- REPATHA 140MG/ML SYRINGE

- REPATHA 420MG/3.5ML CARTRIDGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- RETACRIT 10000UNIT/ML INJ
- RETACRIT 20000UNIT/ML INJ
- RETACRIT 3000UNIT/ML INJ
- RETACRIT 4000UNIT/ML INJ

RETACRIT 20000UNIT/2ML INJ RETACRIT 2000UNIT/ML INJ RETACRIT 40000UNIT/ML INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- RETEVMO 120MG TAB (New Starts Only)
- RETEVMO 40MG CAP (New Starts Only)
- RETEVMO 80MG CAP (New Starts Only)

- RETEVMO 160MG TAB (New Starts Only)
 RETEVMO 40MG TAB (New Starts Only)
- RETEVMO 80MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- sildenafil 20mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Diagnosis confirmed by right heart catheterization. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -lenalidomide 10mg cap (New Starts Only)
- -lenalidomide 2.5mg cap (New Starts Only)
- -lenalidomide 25mg cap (New Starts Only)

- -lenalidomide 15mg cap (New Starts Only)
- lenalidomide 20mg cap (New Starts Only)
- -lenalidomide 5mg cap (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- REXULTI 0.25MG TAB (New Starts Only)
- -REXULTI 1MG TAB (New Starts Only)
- REXULTI 3MG TAB (New Starts Only)

- REXULTI 0.5MG TAB (New Starts Only)
 REXULTI 2MG TAB (New Starts Only)
- REXULTI 4MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For schizophrenia: Two of the following were ineffective or not tolerated: a) aripiprazole, b) olanzapine, c) quetiapine, d) risperidone, e) ziprasidone, or f) lurasidone. For major depressive disorder: Trial of aripiprazole was ineffective or not tolerated. For agitation associated with dementia due to Alzheimer's disease: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- REZLIDHIA 150MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- REZUROCK 200MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- RINVOQ 15MG ER TAB

- RINVOQ 45MG ER TAB

- RINVOQ 30MG ER TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For atopic dermatitis (initial requests): Two of the following were ineffective or not tolerated: a) a medium to very high potency topical steroid, b) a topical calcineurin inhibitor OR c) an oral immunosuppressant. For ulcerative colitis (initial requests): Trial of Hadlima was ineffective or not tolerated. For ankylosing spondylitis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For non-radiographic axial spondyloarthritis (initial requests): Trial of Cimzia was ineffective or not tolerated. For Crohn's disease (initial requests): Trial of Hadlima was ineffective or not tolerated. For juvenile idiopathic arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, psoriatic arthritis, ankylosing spondylitis, or non-radiographic axial spondyloarthritis: Prescribed by, or in consultation with, a rheumatology specialist. For atopic dermatitis: Prescribed by, or in consultation with, an allergist, immunologist, or dermatologist. For Crohn's disease or ulcerative colitis: Prescribed by, or in consultation with a gastroenterologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For atopic dermatitis (initial requests): Member has moderate to severe atopic dermatitis defined as: 1) One of the following: a) body surface area involvement of 10 percent or more OR b) Chart documentation is provided of severity with involvement of the face, head, neck, hands, feet, groin, or intertriginous areas. AND 2) At least two (2) of the following: a) intractable pruritus (itching), b) cracking and oozing/bleeding of skin OR c) impaired activities of daily living. For atopic dermatitis (all requests): Will not be used in combination with another targeted immunomodulator product for the prescribed indication. |

- ROZLYTREK 100MG CAP (New Starts Only)

- ROZLYTREK 200MG CAP (New Starts Only)

-ROZLYTREK 50MG ORAL PELLET (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- RUBRACA 200MG TAB (New Starts Only)

-RUBRACA 250MG TAB (New Starts Only)

-RUBRACA 300MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- RYBELSUS 14MG TAB

- RYBELSUS 7MG TAB

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaRequired Medical InfoAge RestrictionsPrescriber RestrictionCoverage DurationApproved for duration of 1 year.Other Criteria

- RYBELSUS 3MG TAB

- RYDAPT 25MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -vigabatrin 500mg powder for oral soln (New Starts Only)
- -vigadrone 500mg powder for oral soln (New Starts Only)
- -vigpoder 500mg powder for oral soln (New Starts Only)

- -vigabatrin 500mg tab (New Starts Only)
- -vigadrone 500mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SCEMBLIX 100MG TAB (New Starts Only)

- SCEMBLIX 20MG TAB (New Starts Only)

- SCEMBLIX 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For T315I mutation: failure of or intolerance to Iclusig required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SECUADO 3.8MG/24HR PATCH (New Starts Only)

- SECUADO 5.7MG/24HR PATCH (New Starts Only)

- SECUADO 7.6MG/24HR PATCH (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) aripiprazole, b) olanzapine, c) quetiapine, d) risperidone, e) ziprasidone, f) lurasidone, or g) oral asenapine. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SIGNIFOR 0.3MG/ML INJ

- SIGNIFOR 0.9MG/ML INJ

PA Criteria Criteria Details Covered Uses All FDA-approved indications not otherwise excluded from Part D. **Exclusion** Criteria Required Medical Info Age Restrictions Prescriber Restriction **Coverage Duration** Approved for duration of 1 year. Other Criteria

- SIGNIFOR 0.6MG/ML INJ

- SIRTURO 100MG TAB

- SIRTURO 20MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SKYRIZI 150MG/ML AUTO-INJECTOR

- SKYRIZI 180MG/1.2ML CARTRIDGE

SKYRIZI 150MG/ML SYRINGESKYRIZI 360MG/2.4ML CARTRIDGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin. For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For Crohn's disease (all requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For plaque psoriasis: Prescribed by, or in consultation with, a dermatologist. For Psoriatic Arthritis: Prescribed by, or in consultation with, a rheumatology specialist. For Crohn's Disease: Prescribed by, or in consultation with, a gastroenterologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- diclofenac sodium 3% gel

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SOLTAMOX 10MG/5ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of tamoxifen. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SOMAVERT 10MG INJ
- SOMAVERT 20MG INJ
- SOMAVERT 30MG INJ

- SOMAVERT 15MG INJ - SOMAVERT 25MG INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SPRITAM 1000MG TAB FOR ORAL SUSP (New Starts Only)
- SPRITAM 500MG TAB FOR ORAL SUSP (New Starts Only)

SPRITAM 250MG TAB FOR ORAL SUSP (New Starts Only) SPRITAM 750MG TAB FOR ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of generic levetiracetam was ineffective or not tolerated |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- dasatinib 100mg tab (New Starts Only)
- dasatinib 20mg tab (New Starts Only)
- dasatinib 70mg tab (New Starts Only)

- dasatinib 140mg tab (New Starts Only)
- dasatinib 50mg tab (New Starts Only)
- dasatinib 80mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- STELARA 45MG/0.5ML INJ

- STELARA 90MG/ML SYRINGE

- STELARA 45MG/0.5ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin (trial of other agents not required for patients under 18 years of age). For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For ulcerative colitis and Crohn's disease (all requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For psoriatic arthritis: Prescribed by, or in consultation with, a rheumatology specialist. For Crohn's disease and ulcerative colitis: Prescribed by, or in consultation with, a gastroenterologist. For plaque psoriasis: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- STIVARGA 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SUCRAID 8500UNIT/ML ORAL SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SUNOSI 150MG TAB

- SUNOSI 75MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | One of the following was ineffective or not tolerated: a) modafinil OR b) armodafinil. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | A nocturnal polysomnogram was used to confirm diagnosis. |

- sunitinib 12.5mg cap (New Starts Only) - sunitinib 37.5mg cap (New Starts Only) - sunitinib 25mg cap (New Starts Only) - sunitinib 50mg cap (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SYMDEKO TAB 4-WEEK PACK (56)

- SYMDEKO TAB 50-75MG/75MG PACK (56)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- SYMPAZAN 10MG ORAL FILM (New Starts Only)

- SYMPAZAN 20MG ORAL FILM (New Starts Only)

- SYMPAZAN 5MG ORAL FILM (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Both of the following: A) Member is unable to swallow solid dosage forms of clobazam and B) Member is unable to use clobazam oral suspension. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

— trientine 250mg cap

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TABRECTA 150MG TAB (New Starts Only)

- TABRECTA 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TAFINLAR 10MG TAB FOR ORAL SUSP (New Starts Only)

- TAFINLAR 50MG CAP (New Starts Only)

- TAFINLAR 75MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TAGRISSO 40MG TAB (New Starts Only)

- TAGRISSO 80MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TALZENNA 0.1MG CAP (New Starts Only)
- TALZENNA 0.35MG CAP (New Starts Only)
- TALZENNA 0.75MG CAP (New Starts Only)

- TALZENNA 0.25MG CAP (New Starts Only)
- TALZENNA 0.5MG CAP (New Starts Only)
- TALZENNA 1MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- erlotinib 100mg tab (New Starts Only)

- erlotinib 25mg tab (New Starts Only)

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaExclusion CriteriaRequired Medical InfoAge RestrictionsPrescriber RestrictionPrescriber RestrictionCoverage DurationApproved for duration of 1 year.Other CriteriaImage: Coverage Duration

-erlotinib 150mg tab (New Starts Only)

- bexarotene 1% gel (New Starts Only)

-bexarotene 75mg cap (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TASIGNA 150MG CAP (New Starts Only)

- TASIGNA 50MG CAP (New Starts Only)

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaExclusion CriteriaRequired Medical InfoAge RestrictionsPrescriber RestrictionPrescriber RestrictionCoverage DurationApproved for duration of 1 year.Other CriteriaImage: Coverage Duration

- TASIGNA 200MG CAP (New Starts Only)

- tazarotene 0.1% cream

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TAZVERIK 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TEPMETKO 225MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- testosterone 1% (12.5mg/act) gel pump
- *testosterone 1% (50mg) gel packet*
- *testosterone 1.62% (2.5gm) gel packet*
- testosterone 30mg/act topical soln

- testosterone 1% (25mg) gel packet
- *testosterone 1.62% (1.25gm) gel packet*
- testosterone 1.62% (20.25mg/act) gel pump

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | A) For initial requests: documentation is provided of morning testosterone levels, from two separate days, that fall below the normal range for a healthy adult male. B) For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TIBSOVO 250MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- tobramycin 300mg/5ml inh soln

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | Approval will be based off BvD coverage determination. |

- bosentan 125mg tab

— bosentan 62.5mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Diagnosis confirmed by right heart catheterization. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TREMFYA 100MG/ML AUTO-INJECTOR

- TREMFYA 100MG/ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For plaque psoriasis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate at a dose of at least 10mg/week (or maximally tolerated dose) OR b) acitretin. For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) methotrexate OR b) sulfasalazine. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For Psoriatic Arthritis: Prescribed by, or in consultation, with a rheumatology specialist. For Plaque Psoriasis: Prescribed by, or in consultation with, a dermatologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TRIKAFTA 100-50-75MG/150MG TAB PACK (84)

- TRIKAFTA 50-37.5-25MG/75MG TAB PACK (84)

TRIKAFTA 100-50-75MG/75MG GRANULES PACK (56) TRIKAFTA 80-40-60MG/59.5MG GRANULES PACK (56)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TRULICITY 0.75MG/0.5ML AUTO-INJECTOR

- TRULICITY 3MG/0.5ML AUTO-INJECTOR

TRULICITY 1.5MG/0.5ML AUTO-INJECTOR TRULICITY 4.5MG/0.5ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TRUQAP 160MG TAB (New Starts Only)

- TRUQAP 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TUKYSA 150MG TAB (New Starts Only)

- TUKYSA 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TURALIO 125MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- TYENNE 162MG/0.9ML AUTO-INJECTOR

- TYENNE 162MG/0.9ML SYRINGE

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications, Some Medically-Accepted Indications |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Enbrel, b) Hadlima, c) Rinvoq OR d) Xeljanz. For polyarticular juvenile idiopathic arthritis (initial requests): Two of the following were ineffective or not tolerated: a) Hadlima, b) Enbrel OR c) Xeljanz. For giant cell arteritis (all requests): Trial of other agents not required. For systemic sclerosis-associated interstitial lung disease (initial requests): a) Diagnosis is confirmed with documentation provided of both of the following: i) HRCT scan AND ii) pulmonary function tests AND b) Trial of mycophenolate was ineffective or not tolerated. For systemic juvenile idiopathic arthritis (all requests): Trial of other agents not required. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, polyarticular juvenile idiopathic arthritis, systemic juvenile idiopathic arthritis, and giant cell arteritis: Prescribed by, or in consultation with, a rheumatology specialist. For systemic sclerosis-associated interstitial lung disease: Prescribed by, or in consultation with, a pulmonologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- lapatinib 250mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- UBRELVY 100MG TAB

- UBRELVY 50MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of one triptan was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- budesonide 2mg/act rectal foam

-budesonide 9mg er tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of mesalamine was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VALCHLOR 0.016% GEL (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VANFLYTA 17.7MG TAB (New Starts Only)

- VANFLYTA 26.5MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VELTASSA 16.8GM POWDER FOR ORAL SUSP

- VELTASSA 25.2GM POWDER FOR ORAL SUSP

- VELTASSA 8.4GM POWDER FOR ORAL SUSP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: Member has baseline persistent potassium level greater than 5.0 mmol/L. For continuing requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a nephrologist, cardiologist, hematologist or endocrinologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VENCLEXTA 100MG TAB (New Starts Only)
- VENCLEXTA 50MG TAB (New Starts Only)

VENCLEXTA 10MG TAB (New Starts Only)VENCLEXTA TAB STARTER PACK (42) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VERQUVO 10MG TAB

- VERQUVO 5MG TAB

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaRequired Medical InfoAge RestrictionsPrescriber RestrictionCoverage DurationApproved for duration of 1 year.Other Criteria

- VERQUVO 2.5MG TAB

- VERSACLOZ 50MG/ML ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Both of the following: A) Member is unable to swallow clozapine tablet and B) Member is unable to use clozapine ODT. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VERZENIO 100MG TAB (New Starts Only)
- VERZENIO 200MG TAB (New Starts Only)

VERZENIO 150MG TAB (New Starts Only)
VERZENIO 50MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- LIRAGLUTIDE 18MG/3ML PEN INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VIGAFYDE 100MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Both of the following: A) Member is unable to swallow vigabatrin tablet and B) Member is unable to use vigabatrin powder for oral solution. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-vilazodone 10mg tab (New Starts Only)

-vilazodone 40mg tab (New Starts Only)

PA CriteriaCriteria DetailsCovered UsesAll FDA-approved indications not otherwise excluded from Part D.Exclusion CriteriaExclusion CriteriaRequired Medical InfoTwo of the following were ineffective or not tolerated: a) escitalopram, b) sertraline, c) fluoxetine, d) citalopram, e)
paroxetine, f) fluvoxamine, g) bupropion, h) venlafaxine i) desvenlafaxine, or j) duloxetine.Age RestrictionsPrescriber RestrictionCoverage DurationApproved for duration of 1 year.Other CriteriaImage: Coverage Duration

-vilazodone 20mg tab (New Starts Only)

- VITRAKVI 100MG CAP (New Starts Only)

- VITRAKVI 20MG/ML ORAL SOLN (New Starts Only)

- VITRAKVI 25MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VIZIMPRO 15MG TAB (New Starts Only)

- VIZIMPRO 30MG TAB (New Starts Only)

- VIZIMPRO 45MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VONJO 100MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -voriconazole 200mg inj
- -voriconazole 40mg/ml oral susp

voriconazole 200mg tab
voriconazole 50mg tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 6 months. |
| Other Criteria | |

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | 1) Current HCV-RNA titer is provided 3) Member does not have decompensated cirrhosis 3) Previous Hepatitis C treatment(s) is provided. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a gastroenterologist, hepatologist, or infectious disease or transplant specialist. |
| Coverage Duration | Coverage duration of 12 weeks. |
| Other Criteria | Treatment regimen will be approved based on previous treatment experience as defined by current AASLD guidelines. |

-pazopanib 200mg tab (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VOWST 3000000UNIT CAP

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 1 month. |
| Other Criteria | For all requests: Will not be used in combination with fecal microbiota, live for rectal use (Rebyota) or bezlotoxumab (Zinplava). |

- VRAYLAR 1.5MG CAP (New Starts Only)

- VRAYLAR 4.5MG CAP (New Starts Only)

VRAYLAR 3MG CAP (New Starts Only)VRAYLAR 6MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) aripiprazole, b) olanzapine, c) quetiapine, d) risperidone, e) ziprasidone, or f) lurasidone. For major depressive disorder: Trial of aripiprazole was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- VYNDAMAX 61MG CAP

- VYNDAQEL 20MG CAP

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For Initial requests: Diagnosis confirmed by one of the following: A) Cardiac biopsy with positive congo red staining and ATTR confirmation by mass spectrometry or immunofluorescence staining OR B) All of the following: i) Serum kappa/lambda free light chain ratio 0.26 to 1.65 AND ii) Absence of monoclonal protein via serum protein immunofixation AND iii) Absence of monoclonal protein via urine protein immunofixation AND iv) Myocardial uptake of 99mTc-PYP demonstrated by a greater than 1.5 heart-to-contralateral ratio or grade 2 or greater visual evidence. For continuation requests: Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a cardiologist or other provider experienced in the treatment of cardiomyopathy of transthyretin-mediated amyloidosis. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For all requests: Will not be used in combination with Tegsedi, Onpattro, or Amvuttra. |

- WELIREG 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XALKORI 150MG ORAL PELLET (New Starts Only)
- XALKORI 20MG ORAL PELLET (New Starts Only)
- -XALKORI 50MG ORAL PELLET (New Starts Only)

XALKORI 200MG CAP (New Starts Only)
XALKORI 250MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XATMEP 2.5MG/ML ORAL SOLN (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of methotrexate. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | Approval will be based off BvD coverage determination. |

- XCOPRI 100MG TAB (New Starts Only)
- XCOPRI 200MG TAB (New Starts Only)
- -XCOPRI 50MG TAB (New Starts Only)
- XCOPRI TAB 12.5/25MG TITRATION PACK (28) (New Starts Only)
- XCOPRI TAB 150/200MG TITRATION PACK (28) (New Starts Only)
- XCOPRI 150MG TAB (New Starts Only)
- -XCOPRI 25MG TAB (New Starts Only)
- XCOPRI TAB 100/150MG MAINTENANCE PACK (56) (New Starts Onl
- XCOPRI TAB 150/200MG PACK (56) (New Starts Only)
- XCOPRI TAB 50/100MG TITRATION PACK (28) (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Two of the following were ineffective or not tolerated: a) lamotrigine b) carbamazepine c) levetiracetam d) oxcarbazepine e) phenytoin f) topiramate OR g) lacosamide. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XDEMVY 0.25% OPHTH SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, an ophthalmologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XELJANZ 10MG TAB
- XELJANZ 5MG TAB
- XELJANZ XR 22MG TAB

XELJANZ 1MG/ML ORAL SOLN XELJANZ XR 11MG TAB

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For rheumatoid arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For juvenile idiopathic arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For psoriatic arthritis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For ankylosing spondylitis (initial requests): One of the following was ineffective or not tolerated: a) Hadlima OR b) Enbrel. For ulcerative colitis (initial requests): Failure of, or intolerance to Hadlima. For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For rheumatoid arthritis, juvenile idiopathic arthritis, ankylosing spondylitis, or psoriatic arthritis: Prescribed by, or in consultation with a rheumatology specialist. For ulcerative colitis : Prescribed by, or in consultation with a gastroenterologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XERMELO 250MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XGEVA 120MG/1.7ML INJ

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-XIFAXAN 550MG TAB

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For diagnosis of IBS-D, approval will increase quantity limit to 42 tablets over 14 days, maximum of three fills per 1 year. |

- XOLAIR 150MG INJ
- -XOLAIR 150MG/ML SYRINGE
- XOLAIR 300MG/2ML SYRINGE
- XOLAIR 75MG/0.5ML SYRINGE

XOLAIR 150MG/ML AUTO-INJECTOR XOLAIR 300MG/2ML AUTO-INJECTOR XOLAIR 75MG/0.5ML AUTO-INJECTOR

| PA Criteria | Criteria Details |
|------------------------|---|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For initial requests: For asthma: History within the last year of at least 1 asthma exacerbation requiring one of the following, despite regular use of inhaled corticosteroids plus an additional controller(s): a) treatment with systemic corticosteroids, b) emergency department visit OR c) hospitalization. For chronic idiopathic urticaria: one of the following: a) patient remains symptomatic despite H1 antihistamine treatment OR b) has intolerance or contraindication to H1 antihistamine treatment. For nasal polyps: A) Confirmed diagnosis of nasal polyps (see other criteria) AND B) Trial of Dupixent was ineffective or not tolerated. For IgE-mediated food allergy: Confirmed diagnosis of IgE-mediated food allergy (see other criteria). For continuation requests (all diagnoses): Member has benefited with use of this medication. |
| Age Restrictions | |
| Prescriber Restriction | For asthma: Prescribed by, or in consultation with an allergist, pulmonologist, or immunologist. For chronic idiopathic urticaria: Prescribed by, or in consultation with an allergist, dermatologist, or immunologist. For nasal polyps: Prescribed by, or in consultation with, an allergist, immunologist, or otolaryngologist. For IgE-mediated food allergy: Prescribed by, or in consultation with an allergist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For asthma (initial requests): Documentation of diagnosis via skin test or RAST for specific allergy sensitivity. For nasal polyps (initial requests): Diagnosis is confirmed with a sinus CT scan AND at least four of the following apply: a) prior surgery for bilateral nasal polyposis b) evidence of type 2 inflammation c) two or more courses of oral corticosteroids required in the prior year d) significantly impaired quality of life e) significant loss of smell f) diagnosis of comorbid asthma. For IgE-mediated food allergy (initial requests): Both of the following: a) diagnosis supported by one of the following: i) positive skin prick test or ii) positive serum IgE test and b) diagnosis confirmed by one of the following: i) positive oral food challenge or ii) history of anaphylaxis to the suspected food allergen. For asthma (all requests): Will not be used in combination with another targeted immunomodulator product for the prescribed indication. For IgE-mediated |

food allergy (all requests): Will not be used in combination with peanut allergen powder (Palforzia).

- XOSPATA 40MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XPOVIO TAB 100MG ONCE WEEKLY CARTON (8) (New Starts Only)
- XPOVIO TAB 40MG TWICE WEEKLY CARTON (8) (New Starts Only)
- XPOVIO TAB 60MG TWICE WEEKLY CARTON (24) (New Starts Only

- XPOVIO TAB 80MG TWICE WEEKLY CARTON (32) (New Starts Only

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- XPOVIO TAB 40MG ONCE WEEKLY CARTON (4) (New Starts Only)
- XPOVIO TAB 60MG ONCE WEEKLY CARTON (4) (New Starts Only)
- XPOVIO TAB 80MG ONCE WEEKLY CARTON (8) (New Starts Only)

- XTANDI 40MG CAP (New Starts Only)

- XTANDI 80MG TAB (New Starts Only)

PA Criteria **Criteria Details** Covered Uses All FDA-approved indications, Some Medically-Accepted Indications **Exclusion** Criteria For metastatic castration-resistant prostate cancer and metastatic castration-sensitive prostate cancer: Trial of abiraterone Required Medical Info was ineffective or not tolerated. For nonmetastatic castration-resistant prostate cancer: Both of the following were ineffective or not tolerated: a) Nubeqa and b) Erleada. For non metastatic castration sensitive prostate cancer with biochemical recurrence at high risk for metastasis: Trial of other agents not required. For homologous recombination repair gene-mutated metastatic castration-resistant prostate cancer in combination with Talzenna: Trial of other agents not required. Age Restrictions Prescriber Restriction **Coverage Duration** Approved for duration of 1 year. Other Criteria

- XTANDI 40MG TAB (New Starts Only)

- SODIUM OXYBATE 500MG/ML ORAL SOLN

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | For excessive daytime sleepiness with narcolepsy in adults: Both of the following were ineffective or not tolerated: a) Sunosi AND b) either modafinil or armodafinil. Trial of other agents not required for patients aged 7 to 17 years. For cataplexy with narcolepsy: Trial of other agents not required. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | For excessive daytime sleepiness with narcolepsy: Documentation is provided of full nocturnal polysomnogram used to confirm diagnosis. For cataplexy with narcolepsy: Documentation is provided of one of the following to confirm diagnosis: a) full nocturnal polysomnogram OR b) low cerebrospinal fluid orexin-A concentration. |

- ZAVZPRET 10MG/ACT NASAL SPRAY

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial of one triptan was ineffective or not tolerated. |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ZEJULA 100MG TAB (New Starts Only)

- ZEJULA 300MG TAB (New Starts Only)

- ZEJULA 200MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ZELBORAF 240MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ZOLINZA 100MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- -zolpidem tartrate 10mg tab
- -zolpidem tartrate 5mg tab

-zolpidem tartrate 12.5mg er tab -zolpidem tartrate 6.25mg er tab

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Trial and failure of trazodone. |
| Age Restrictions | Prior Authorization applies to members 65 years or older. |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ZONISADE 100MG/5ML ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Member is unable to swallow solid dosage forms of zonisamide. |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ZTALMY 50MG/ML ORAL SUSP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | Documentation is provided of a CDKL5 gene mutation |
| Age Restrictions | |
| Prescriber Restriction | Prescribed by, or in consultation with, a neurologist. |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

- ZURZUVAE 20MG CAP (New Starts Only)

-ZURZUVAE 25MG CAP (New Starts Only)

-ZURZUVAE 30MG CAP (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for 1 month. |
| Other Criteria | |

-ZYDELIG 100MG TAB (New Starts Only)

-ZYDELIG 150MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |

-ZYKADIA 150MG TAB (New Starts Only)

| PA Criteria | Criteria Details |
|------------------------|--|
| Covered Uses | All FDA-approved indications not otherwise excluded from Part D. |
| Exclusion Criteria | |
| Required Medical Info | |
| Age Restrictions | |
| Prescriber Restriction | |
| Coverage Duration | Approved for duration of 1 year. |
| Other Criteria | |