

Prior Authorization Criteria 2026 MCORE

Last Updated: 10/7/2025

### **ABRYSVO**

### **Products Affected**

#### • Abrysvo

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | All indications: Vaccine is being used for the prevention of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV). Immunization in age 60 years of age and older: Patient has not received an RSV vaccine (i.e., Abrysvo, Arexvy) in their lifetime. Age greater than or equal to 60 years. Immunization in pregnancy: Patient has not received Abrysvo vaccine for the current pregnancy. Both of the following: 1) Will be used for active immunization of pregnant individuals at 32 through 36 weeks gestational age and 2) Will be used for the prevention of severe LRTD caused by RSV in infants from birth through 6 months of age. Immunization in age 18 through 59 years: Patient has not received an RSV vaccine (i.e., Abrysvo, Arexvy) in their lifetime. Age 18 through 59 years. Patient is at increased risk for LRTD caused by RSV. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | 6 months  |
| Other Criteria                     | N/A   |

| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug. |
|-------------------------------------|--|
|-------------------------------------|--|

### **ADALIMUMAB-AATY**

#### **Products Affected**

- Adalimumab-aaty 1-pen Kit INJ 80MG/0.8ML
- Adalimumab-aaty 2-pen Kit

- Adalimumab-aaty 2-syringe Kit
- Adalimumab-aaty Cd/uc/hs Starter

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Rheumatoid Arthritis (RA) (Initial): Diagnosis (Dx) of moderately to severely active RA. Minimum duration of a 3-month trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies at maximally tolerated doses: methotrexate (MTX), leflunomide, sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Dx of moderately to severely active PJIA. Minimum duration of a 6-week TF/C/I to one of the following conventional therapies at maximally tolerated doses: leflunomide or methotrexate. Psoriatic Arthritis (PsA) (Initial): Dx of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Plaque psoriasis (PsO) (Initial): Dx of moderate to severe chronic PSO. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), C/I to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, OR calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Ankylosing Spondylitis (AS) (Initial): Dx of active AS. Minimum duration of a one-month TF/C/I to one NSAID (eg, ibuprofen, naproxen) at maximally tolerated doses. Crohn's Disease (CD) (Initial): Dx of moderately to severely active CD. One of the following: frequent diarrhea and abdominal pain, at least 10% weight loss, complications (eg, obstruction, fever, abdominal mass), abnormal lab values (eg, CRP), OR CD Activity Index (CDAI) greater than 220. Uveitis (UV) (Initial): Dx of non-infectious uveitis classified as intermediate, posterior, or panuveitis. |

| Age Restrictions                    | N/A   |
|-------------------------------------|---|
| Prescriber<br>Restrictions          | RA, AS, JIA: (Initial) Prescribed by or in consultation with a rheumatologist. PsA: (Initial) Prescribed by or in consultation with a dermatologist or rheumatologist. Plaque Psoriasis, HS: (Initial) Prescribed by or in consultation with a dermatologist. CD, UC: (Initial) Prescribed by or in consultation with a gastroenterologist. UV (initial): Prescribed by or in consultation with a rheumatologist or an ophthalmologist.   |
| Coverage<br>Duration                | All uses (initial, reauth): plan year.  |
| Other Criteria                      | Ulcerative Colitis (UC) (Initial): Dx of moderately to severely active UC. One of the following: greater than 6 stools per day, frequent blood in the stools, frequent urgency, presence of ulcers, abnormal lab values (eg, hemoglobin, ESR, CRP), OR dependent on, or refractory to, corticosteroids. Hidradenitis suppurativa (HS) (Initial): Dx of moderate to severe hidradenitis suppurativa (ie, Hurley Stage II or III). RA, PJIA, PsA, AS, PsO, CD, HS, UV, UC (Reauth): Patient demonstrates positive clinical response to therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## ADALIMUMAB-ADBM

### **Products Affected**

#### Adalimumab-adbm

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Rheumatoid Arthritis (RA) (Initial): Diagnosis (Dx) of moderately to severely active RA. Minimum duration of a 3-month trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies at maximally tolerated doses: methotrexate (MTX), leflunomide, sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Dx of moderately to severely active PJIA. Minimum duration of a 6-week TF/C/I to one of the following conventional therapies at maximally tolerated doses: leflunomide or methotrexate. Psoriatic Arthritis (PsA) (Initial): Dx of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Plaque psoriasis (PsO) (Initial): Dx of moderate to severe chronic PSO. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), C/I to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, OR calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Ankylosing Spondylitis (AS) (Initial): Dx of active AS. Minimum duration of a one-month TF/C/I to one NSAID (eg, ibuprofen, naproxen) at maximally tolerated doses. Crohn's Disease (CD) (Initial): Dx of moderately to severely active CD. One of the following: frequent diarrhea and abdominal pain, at least 10% weight loss, complications (eg, obstruction, fever, abdominal mass), abnormal lab values (eg, CRP), OR CD Activity Index (CDAI) greater than 220. Uveitis (UV) (Initial): Dx of non-infectious uveitis classified as intermediate, posterior, or panuveitis. |
| Age Restrictions                   | N/A  |

| Prescriber<br>Restrictions          | RA, AS, JIA: (Initial) Prescribed by or in consultation with a rheumatologist. PsA: (Initial) Prescribed by or in consultation with a dermatologist or rheumatologist. Plaque Psoriasis, HS: (Initial) Prescribed by or in consultation with a dermatologist. CD, UC: (Initial) Prescribed by or in consultation with a gastroenterologist. UV (initial): Prescribed by or in consultation with a rheumatologist or an ophthalmologist.   |
|-------------------------------------|---|
| Coverage<br>Duration                | All uses (initial, reauth): plan year.  |
| Other Criteria                      | Ulcerative Colitis (UC) (Initial): Dx of moderately to severely active UC. One of the following: greater than 6 stools per day, frequent blood in the stools, frequent urgency, presence of ulcers, abnormal lab values (eg, hemoglobin, ESR, CRP), OR dependent on, or refractory to, corticosteroids. Hidradenitis suppurativa (HS) (Initial): Dx of moderate to severe hidradenitis suppurativa (ie, Hurley Stage II or III). RA, PJIA, PsA, AS, PsO, CD, HS, UV, UC (Reauth): Patient demonstrates positive clinical response to therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **ADEMPAS**

## **Products Affected**

### • Adempas

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH AND PAH is symptomatic AND One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH. Chronic thromboembolic pulmonary hypertension (CTEPH): One of the following: A) Both of the following: 1) Diagnosis of inoperable or persistent/recurrent CTEPH and 2) CTEPH is symptomatic OR B) Patient is currently on any therapy for the diagnosis of CTEPH. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PAH, CTEPH: Prescribed by or in consultation with a pulmonologist or cardiologist.  |
| Coverage<br>Duration                | PAH, CTEPH: plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **AIMOVIG**

## **Products Affected**

### • Aimovig

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Episodic Migraines (EM) (initial): Diagnosis of EM. Patient has greater than or equal to 4 migraine days per month. Chronic Migraines (CM) (initial): Diagnosis of CM. Patient has greater than or equal to 8 migraine days per month. All Indications (initial): Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | EM, CM (initial, reauth): Plan year.  |
| Other Criteria                      | EM, CM (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# AKEEGA

### **Products Affected**

### • Akeega

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of prostate cancer. Disease is all of the following: a) metastatic, b) castration-resistant, and c) deleterious or suspected deleterious BRCA-mutated (BRCAm). Used in combination with prednisone. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **ALECENSA**

## **Products Affected**

#### • Alecensa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is anaplastic lymphoma kinase (ALK)-positive. One of the following: 1) Disease is one of the following: a) Recurrent, b) Advanced, or c) Metastatic, or 2) Used as adjuvant treatment following tumor resection. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## **ALOSETRON**

### **Products Affected**

### • Alosetron Hydrochloride

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Irritable bowel syndrome (IBS) (Initial): Exclude if patient is of the male gender.  |
| Required<br>Medical<br>Information  | IBS (Initial): Diagnosis of chronic severe diarrhea-predominant IBS. IBS (Reauthorization): Symptoms of IBS continue to persist. Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | IBS (Initial): 18 years and older.   |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | IBS (Initial): 12 weeks. IBS (Reauthorization): 6 months.  |
| Other Criteria                      | IBS (initial): Trial and failure, contraindication, or intolerance to an anti-<br>diarrheal agent [eg, loperamide].  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# ALPHA - 1 PROTEINASE INHIBITORS

### **Products Affected**

• Prolastin-c INJ 1000MG/20ML

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Alpha-1 antitrypsin (AAT) deficiency (initial): Diagnosis of congenital AAT deficiency. Diagnosis of emphysema. One of the following: Pi*ZZ, Pi*Z(null) or Pi*(null)(null) protein phenotypes (homozygous) or Other rare AAT disease genotypes associated with pre-treatment serum AAT level less than 11 µmol/L [eg, Pi(Malton, Malton), Pi(SZ)]. One of the following: 1) Circulating pre-treatment serum AAT level less than 11 µmol/L (which corresponds to less than 80 mg/dL if measured by radial immunodiffusion or less than 57 mg/dL if measured by nephelometry), or 2) Patient has a concomitant diagnosis of necrotizing panniculitis. Continued conventional treatment for emphysema (eg, bronchodilators). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | AAT deficiency (initial, reauth): plan year   |
| Other Criteria                      | AAT deficiency (reauth): Patient demonstrates positive clinical response to therapy. Continued conventional treatment for emphysema (e.g., bronchodilators).  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **ALUNBRIG**

## **Products Affected**

## • Alunbrig

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of metastatic, recurrent, or advanced NSCLC and tumor is anaplastic lymphoma kinase (ALK)-positive. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **AMBRISENTAN**

### **Products Affected**

#### • Ambrisentan

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH. PAH is symptomatic. One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PAH: Prescribed by or in consultation with a pulmonologist or cardiologist.   |
| Coverage<br>Duration                | PAH: plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **APREPITANT**

### **Products Affected**

### • Aprepitant

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute Chemotherapy-induced Nausea and Vomiting (CINV): Patient is currently receiving moderately or highly emetogenic chemotherapy. Patient is concurrently on both corticosteroid [eg, Decadron (dexamethasone)] and 5-HT3 receptor antagonist [eg, Aloxi (palonosetron), Anzemet (dolasetron), Kytril (granisetron), Zofran (ondansetron)]. Delayed Chemotherapy-induced Nausea and Vomiting Prevention: Patient is currently receiving highly emetogenic chemotherapy and corticosteroid [eg, Decadron (dexamethasone)], or patient is receiving an anthracycline [eg, Adriamycin (doxorubicin), Ellence (epirubicin)] and Cytoxan (cyclophosphamide), or patient is currently receiving moderately emetogenic chemotherapy and was given aprepitant (oral or IV) on day 1 of chemotherapy. Postoperative Nausea and Vomiting (PONV): For the prevention of postoperative nausea and vomiting when administered prior to the induction of anesthesia. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Acute CINV, Delayed CINV: plan year. PONV: 1 month   |
| Other Criteria                      | Subject to Part B vs. Part D review.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **ARANESP**

#### **Products Affected**

 Aranesp Albumin Free INJ 100MCG/0.5ML, 100MCG/ML, 10MCG/0.4ML, 150MCG/0.3ML, 200MCG/0.4ML, 200MCG/ML, 25MCG/0.42ML, 25MCG/ML, 300MCG/0.6ML, 40MCG/0.4ML, 40MCG/ML, 500MCG/ML, 60MCG/0.3ML, 60MCG/ML

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Anemia due to Chronic Kidney Disease (CKD) (Initial): Diagnosis of CKD. Anemia by lab values (Hct less than 30% or Hgb less than 10 g/dL) within 30 days of request. The rate of hemoglobin decline indicates the likelihood of requiring a red blood cell (RBC) transfusion, and reducing the risk of alloimmunization and/or other RBC transfusion-related risks is a goal. (Reauth): Diagnosis of CKD. Most recent or average (avg) Hct over 3 mo is 33% or less (Hgb 11 g/dL or less) for patients on dialysis, without ESRD OR Most recent or average (avg) Hct over 3 mo is 30% or less (Hgb 10 g/dL or less) for patients not on dialysis OR Most recent or average (avg) Hct over 3 mo is 36% or less (Hgb 12 g/dL or less) for pediatric patients. Patient demonstrates positive clinical response to therapy from pre-treatment level. Anemia w/ chemo (Initial): Other causes of anemia ruled out. Anemia w/ labs (Hct less than 30%, Hgb less than 10 g/dL) within prior 2 weeks of request. Cancer is non-myeloid malignancy. Patient is receiving chemo. (Reauth): Anemia by labs (Hgb less than 10 g/dl or Hct less than 30%) collected within the prior 2 weeks of request. Patient demonstrates positive clinical response to therapy from pre-treatment level. Patient is receiving chemo. Anemia in MDS (Init): Diagnosis of MDS. Serum erythropoietin 500 mU/mL or less, or transfusion dependent MDS. (Reauth): Most recent or avg Hct over 3 months was 36% or less, OR most recent or avg Hgb over 3 months is 12 g/dl or less. Patient demonstrates positive clinical response to therapy from pre-treatment level. |
| Age Restrictions                   | N/A   |

| Prescriber<br>Restrictions          | N/A  |
|-------------------------------------|--|
| Coverage<br>Duration                | CKD(Init): 6 mo. CKD(reauth):plan yr. Chemo(init, reauth): 3 mo. MDS(init): 3 mo,(reauth): plan yr   |
| Other Criteria                      | ESRD patients: Coverage is excluded under Medicare Part D for patients with ESRD on dialysis for any indication related or unrelated to treatment of ESRD since the payment for the drug is included in the ESRD PPS payment bundle. NON-ESRD PATIENTS for the following indications: Off-label uses (except Anemia in Myelodysplastic Syndrome (MDS): Will not be approved if patient has Hgb greater than 10 g/dL or Hct greater than 30%.CKD (init, reauth), Chemo (init), MDS (init): Adequate iron stores confirmed by both of the following: a) Patient's ferritin level is greater than 100 mcg/L and b) Patient's transferrin saturation (TSAT) is greater than 20%. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## **A**RCALYST

### **Products Affected**

### • Arcalyst

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Cryopyrin-Associated Period Syndromes (CAPS): Diagnosis of CAPS, Familial Cold Autoinflammatory Syndrome (FCAS) or Muckle-Wells Syndrome (MWS). Deficiency of Interleukin-1 Receptor Antagonist (DIRA): Diagnosis of DIRA. Patient weighs at least 10 kg. Patient is currently in remission (e.g., no fever, skin rash, and bone pain/no radiological evidence of active bone lesions/C-reactive protein [CRP] less than 5 mg/L). Recurrent Pericarditis (initial): Diagnosis of recurrent pericarditis as evidenced by at least 2 episodes that occur a minimum of 4 to 6 weeks apart. Trial and failure, contraindication, or intolerance (TF/C/I) to at least one of the following: nonsteroidal anti-inflammatory drugs (e.g., ibuprofen, naproxen), colchicine, or corticosteroids (e.g., prednisone). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | CAPS, DIRA: plan year. Recurrent Pericarditis (initial, reauth): plan year.   |
| Other Criteria                      | Recurrent Pericarditis (reauth): Patient demonstrates positive clinical response to therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **AREXVY**

## **Products Affected**

### • Arexvy

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Immunization in age 60 years of age and older: Vaccine is being used for the prevention of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV). Patient has not received an RSV vaccine (i.e., Abrysvo, Arexvy) in their lifetime. Age greater than or equal to 60 years. Immunization in age 50 through 59 years: Vaccine is being used for the prevention of LRTD caused by RSV. Patient has not received an RSV vaccine (i.e., Abrysvo, Arexvy) in their lifetime. Age 50 through 59 years. Patient is at increased risk for LRTD caused by RSV. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 6 months   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **A**RIKAYCE

## **Products Affected**

### • Arikayce

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Mycobacterium avium complex (MAC) lung disease: Diagnosis of Mycobacterium avium complex (MAC) lung disease. Used as part of a combination antibacterial drug regimen. Used in patients who do not achieve at least two negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy (e.g., a macrolide, a rifamycin, ethambutol, etc). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with an infectious disease specialist or pulmonologist.  |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# ARMODAFINIL

## **Products Affected**

#### • Armodafinil

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Obstructive sleep apnea (OSA) (Initial): Diagnosis (dx) of OSA defined by one of the following: a) 15 or more obstructive respiratory events per hour of sleep confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible), or b) both of the following: 5 or more obstructive respiratory events per hour of sleep confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible), AND 1 of the following symptoms: unintentional sleep episodes during wakefulness, daytime sleepiness, unrefreshing sleep, fatigue, insomnia, waking up breath holding/gasping/choking, loud snoring, or breathing interruptions during sleep. Shift-work disorder (SWD) (Initial): Dx of SWD confirmed by one of the following: 1) Symptoms of excessive sleepiness or insomnia for at least 3 months, which is associated with a work period (usually night work) that occurs during the normal sleep period, OR 2) A sleep study demonstrating loss of a normal sleep-wake pattern (ie, disturbed chronobiologic rhythmicity). Confirmation that no other medical conditions or medications are causing the symptoms of excessive sleepiness or insomnia. Narcolepsy (initial): Dx of narcolepsy as confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible). |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | OSA, SWD: Initial, Reauth: 6 mo. Narcolepsy: Initial, Reauth: Plan Year   |
| Other Criteria                     | OSA, Narcolepsy (Reauth): Patient demonstrates positive clinical response to therapy. SWD (Reauth): Patient demonstrates positive clinical response to therapy.   |

| Prerequisite<br>Therapy | Criteria DOES NOT require use of a prerequisite Part D drug. |
|-------------------------|--|
| Required                |  |

# **A**UGTYRO

### **Products Affected**

### • Augtyro

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: a) advanced, or b) metastatic. Disease is ROS1-positive. Solid Tumors: Diagnosis of solid tumors. Disease is positive for neutrophic tyrosine receptor kinase (NTRK) gene fusion (e.g., ETV6-NTRK3, TPM3-NTRK1, LMNA-NTRK1). Disease is one of the following: a) Locally advanced, b) Metastatic, or c) Unresectable (including cases where surgical resection is likely to result in severe morbidity). One of the following: a) Disease has progressed following treatment, or b) There is no satisfactory alternative therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year.   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **AUSTEDO**

### **Products Affected**

#### • Austedo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Chorea associated with Huntington's disease: Diagnosis of Chorea associated with Huntington's disease. Tardive dyskinesia: Diagnosis of tardive dyskinesia. One of the following: 1) Patient has persistent symptoms of tardive dyskinesia despite a trial of dose reduction, tapering, or discontinuation of the offending medication or 2) Patient is not a candidate for a trial of dose reduction, tapering, or discontinuation of the offending medication. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Huntington's disease chorea: Prescribed by or in consultation with a neurologist. Tardive dyskinesia: Prescribed by or in consultation with a neurologist or psychiatrist.   |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# AVMAPKI FAKZYNJA

### **Products Affected**

• Avmapki Fakzynja Co-pack

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of recurrent low-grade serous ovarian cancer (LGSOC). Tumor is KRAS-mutated. Patient has received prior systemic therapy (e.g., chemotherapy). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **A**YVAKIT

### **Products Affected**

### • Ayvakit

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Gastrointestinal stromal tumor (GIST): Diagnosis of GIST. One of the following: 1) Patient has unresectable, recurrent, or metastatic disease after failure on one approved therapy (e.g., imatinib, sunitinib, dasatinib, regorafenib, ripretinib) OR 2) Both of the following: a) Disease is one of the following: i) unresectable, ii) metastatic, iii) recurrent, iv) gross residual disease (R2 resection), v) residual disease with significant morbidity, vi) limited progression, or vii) resectable with significant morbidity AND b) Presence of platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation, including PDGFRA D842V mutations. Systemic Mastocytosis: Both of the following: 1) Diagnosis of one of the following: a) advanced systemic mastocytosis (AdvSM), b) aggressive systemic mastocytosis (ASM), c) systemic mastocytosis with an associated hematological neoplasm (SM-AHN), or d) mast cell leukemia (MCL) AND 2) platelet count is greater than 50 x 10^9/L. Ayvakit 25 mg - Indolent Systemic Mastocytosis (ISM): Diagnosis of ISM. Platelet count is greater than 50 x 10^9/L. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **B**ALVERSA

## **Products Affected**

#### • Balversa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Urothelial Carcinoma: Diagnosis of urothelial carcinoma (UC). One of the following: Locally advanced or metastatic. Presence of fibroblast growth factor receptor (FGFR) 3 genetic alterations. Disease has progressed on or after at least one line of prior systemic therapy (e.g., chemotherapy). One of the following: 1) Patient has received prior systemic therapy containing an immune checkpoint inhibitor (e.g., pembrolizumab, nivolumab, avelumab) or 2) Patient is not eligible for immune checkpoint inhibitor therapy (e.g., pembrolizumab, nivolumab, avelumab). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **B**ENLYSTA

### **Products Affected**

• Benlysta INJ 200MG/ML

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Systemic Lupus Erythematosus (SLE) (Initial): Diagnosis of active SLE. Autoantibody positive (ie, anti-nuclear antibody [ANA] titer greater than or equal to 1:80 or anti-dsDNA level greater than or equal to 30 IU/mL). Currently receiving at least one standard of care treatment for active SLE (eg, antimalarials [eg, Plaquenil (hydroxychloroquine)], corticosteroids [eg, prednisone], or immunosuppressants [eg, methotrexate, Imuran (azathioprine)]). Lupus Nephritis (Initial): Diagnosis of active lupus nephritis. Currently receiving standard of care treatment for active lupus nephritis (e.g., corticosteroids [e.g., prednisone] with mycophenolate or cyclophosphamide). SLE, Lupus Nephritis (Reauthorization): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | SLE, Lupus Nephritis (initial, reauth): 6 months   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **BENZODIAZEPINES**

#### **Products Affected**

- Alprazolam TABS
- Clorazepate Dipotassium TABS
- Diazepam ORAL SOLN 5MG/5ML
- Diazepam TABS
- Diazepam Intensol

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | The drug is being prescribed for a medically accepted indication.  Prescriber acknowledges safety concerns (e.g., drowsiness, increased risk of falls, altered mental status) with the use of the prescribed benzodiazepine. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **B**ESREMI

### **Products Affected**

#### • Besremi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of polycythemia vera as confirmed by: A) All of the following: 1) One of the following: a) Hemoglobin greater than 16.5 g/dL for men or hemoglobin greater than 16.0 g/dL for women, b) Hematocrit greater than 49% for men or hematocrit greater than 48% for women, AND 2) Bone marrow biopsy showing hypercellularity for age with trilineage growth (panmyelosis) including prominent erythroid, granulocytic and megakaryocytic proliferation with pleomorphic, mature megakaryocytes, AND 3) One of the following: a) Presence of JAK2 V617F or JAK2 exon 12 mutation or b) Subnormal serum erythropoietin level, OR B) All of the following: 1) Patient with sustained absolute erythrocytosis as demonstrated by one of the following: a) Hemoglobin greater than 18.5 g/dL for men or greater than 16.5 g/dL for women, or b) Hematocrit greater than 55.5% for men or greater than 49.5% for women, AND 2) Presence of JAK2 V617F or JAK2 exon 12 mutation, AND 3) Subnormal serum erythropoietin level. For high-risk polycythemia vera only (patient greater than or equal to 60 years old and/or prior thrombosis history), trial and inadequate response, contraindication or intolerance to hydroxyurea. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# BEXAROTENE

### **Products Affected**

#### • Bexarotene

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Cutaneous T-cell lymphoma (CTCL): Diagnosis of cutaneous T-cell lymphoma (CTCL). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.           |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                     |

# **B**EXSERO

### **Products Affected**

#### • Bexsero INJ 0.5ML

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **BOSENTAN**

## **Products Affected**

#### • Bosentan TABS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH AND PAH is symptomatic AND One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | PAH: Prescribed by or in consultation with a pulmonologist or cardiologist   |
| Coverage<br>Duration                | PAH: Plan year   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **BOSULIF**

## **Products Affected**

#### • Bosulif

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of chronic myelogenous/myeloid leukemia (CML).               |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# **B**RAFTOVI

### **Products Affected**

#### • Braftovi CAPS 75MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma. Patient is positive for BRAF V600 mutation. Used in combination with Mektovi (binimetinib). Colorectal Cancer: One of the following diagnoses: Colon Cancer or Rectal Cancer. One of the following: 1) Unresectable or advanced disease or 2) Metastatic disease. One of the following: 1) Patient has received prior therapy or 2) Used in combination with mFOLFOX6 (fluorouracil, leucovorin, and oxaliplatin). Patient is positive for BRAF V600E mutation. Used in combination with one of the following: 1) Erbitux (cetuximab) or 2) Vectibix (panitumumab). Non-Small Cell Lung Cancer: Diagnosis of metastatic NSCLC. Patient is positive for BRAF V600 mutation. Used in combination with Mektovi (binimetinib). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **B**RIVIACT

### **Products Affected**

• Briviact ORAL SOLN

#### • Briviact TABS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Partial-onset seizures: Diagnosis of partial-onset seizures.           |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# **B**RUKINSA

### **Products Affected**

#### • Brukinsa CAPS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Mantle Cell Lymphoma (MCL): Diagnosis of MCL. Contraindication or intolerance to Calquence (acalabrutinib). Waldenstrom's Macroglobulinemia (WM)/Lymphoplasmacytic Lymphoma (LPL): Diagnosis of WM/LPL. Contraindication or intolerance to Imbruvica (ibrutinib). Marginal Zone Lymphoma (MZL): Diagnosis of MZL. Disease is relapsed, refractory, or progressive. Patient has received at least one anti-CD20-based regimen (e.g., rituximab, obinutuzumab). Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL. Contraindication or intolerance to Calquence (acalabrutinib). Follicular Lymphoma (FL): Diagnosis of FL. Disease is relapsed or refractory. Used in combination with Gazyva (obinutuzumab). Patient has received at least two prior lines of systemic therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year.   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **B**YLVAY

# Products Affected Bylvay

• Bylvay (pellets)

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Progressive familial intrahepatic cholestasis (PFIC) (initial): Diagnosis of progressive familial intrahepatic cholestasis (PFIC) type 1, 2, or 3, confirmed by one of the following: 1) Diagnostic test (e.g., liver function test, liver ultrasound and biopsy, bile analysis) or 2) Genetic testing. Patient is experiencing moderate to severe pruritus. Patient has had an inadequate response to one of the following treatments used for the relief of pruritus: a) Ursodeoxycholic acid (e.g., Ursodiol), b) Antihistamines (e.g., diphenhydramine, hydroxyzine), c) Rifampin, or d) Bile acid sequestrants (e.g., Questran). Alagille syndrome (ALGS) (initial): Diagnosis of Alagille Syndrome (ALGS). Molecular genetic testing confirms mutations in the JAG1 or NOTCH2 gene. Patient is experiencing both of the following: 1) Moderate to severe cholestatic pruritus, and 2) Patient has a serum bile acid concentration above the upper limit of the normal reference for the reporting laboratory. Patient has had an inadequate response to one of the following treatments used for the relief of pruritus: a) Ursodeoxycholic acid (e.g., Ursodiol), b) Antihistamines (e.g., diphenhydramine, hydroxyzine), c) Rifampin, or d) Bile acid sequestrants (e.g., Questran). |
| Age Restrictions                   | PFIC (initial): Patient is 3 months of age or older  |
| Prescriber<br>Restrictions         | PFIC, ALGS (initial): Prescribed by or in consultation with a hepatologist or gastroenterologist   |
| Coverage<br>Duration               | PFIC, ALGS (initial): 6 months. PFIC, ALGS (reauth): Plan year.  |
| Other Criteria                     | PFIC, ALGS (reauth): Patient demonstrates positive clinical response to therapy.   |

# **C**ABLIVI

### **Products Affected**

#### • Cablivi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acquired thrombotic thrombocytopenic purpura (aTTP): Diagnosis of aTTP. First dose was/will be administered by a healthcare provider as a bolus intravenous injection. Used in combination with immunosuppressive therapy (e.g. rituximab, glucocorticoids). One of the following: 1) Used in combination with plasma exchange or 2) both of the following: patient has completed plasma exchange and less than 59 days have or will have elapsed beyond the last plasma exchange. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 3 months   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **CABOMETYX**

### **Products Affected**

#### • Cabometyx

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Renal cell carcinoma (RCC): Diagnosis of one of the following: a) Stage IV or relapsed RCC or b) Hereditary leiomyomatosis and RCC (HLRCC). Hepatocellular Carcinoma (HCC): Diagnosis of HCC. Used as subsequent-line systemic therapy. One of the following: a) Patient has liver-confined, unresectable disease and is not a transplant candidate or b) Patient has extrahepatic/metastatic disease and deemed ineligible for resection, transplant, or locoregional therapy. Differentiated Thyroid Cancer (DTC): Diagnosis of DTC. Disease is one of the following: a) locally advanced or b) metastatic. Disease has progressed following prior VEGFR-targeted therapy (e.g., Lenvima [lenvatinib], Nexavar [sorafenib]). Disease is radioactive iodine-refractory or ineligible. Pancreatic neuroendocrine tumors (pNET), Extra-pancreatic neuroendocrine tumors (epNET): Diagnosis of pNET or epNET. Disease is one of the following: a) unresectable, b) locally advanced, or c) metastatic. Tumors are well-differentiated. Patient has received at least one prior line of therapy (e.g., octreotide, lanreotide, chemotherapy). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **CALQUENCE**

### **Products Affected**

### • Calquence TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Mantle Cell Lymphoma: Diagnosis of mantle cell lymphoma (MCL). One of the following: 1) All of the following: a) Patient has received no prior therapy for MCL (e.g., bortezomib, rituximab), b) Patient is ineligible for autologous hematopoietic stem cell transplantation (HSCT), and c) Used in combination with bendamustine and rituximab, or 2) Patient has received at least one prior therapy for MCL. Chronic Lymphocytic Leukemia (CLL) or Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **CAYSTON**

### **Products Affected**

#### • Cayston

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Cystic Fibrosis (CF): Diagnosis of CF and lung infection with positive culture demonstrating Pseudomonas aeruginosa infection. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **CHENODAL**

#### **Products Affected**

#### • Chenodal

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Initial: Diagnosis of radiolucent gallstones. Patient has a well-opacifying gallbladder visualized by oral cholecystography. Trial and failure, contraindication or intolerance to ursodiol. Patient is not a candidate for surgery. Stones are not calcified (radiopaque) or radiolucent bile pigment stones. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Initial, reauth: Plan year   |
| Other Criteria                      | Reauth: Patient's disease status has been re-evaluated since the last authorization to confirm the patient's condition warrants continued treatment as evidenced by Oral cholecystograms or ultrasonograms.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **C**HOLBAM

### **Products Affected**

#### • Cholbam

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Bile acid synthesis disorders due to single enzyme defects (BAS) (initial): diagnosis of a bile acid synthesis disorder due to a single enzyme defect. Peroxisomal disorders (PD) (initial): All of the following: 1) diagnosis of peroxisomal disorder, 2) patient exhibits at least one of the following: a) liver disease (eg, jaundice, elevated serum transaminases), OR b) steatorrhea, OR c) complications from decreased fat-soluble vitamin absorption (eg, poor growth), AND 3) Will be used as an adjunctive treatment. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Initial: Prescribed by a hepatologist, medical geneticist, gastroenterologist, OR other specialist that treats inborn errors of metabolism.  |
| Coverage<br>Duration                | Initial: 3 months Reauth: Plan year  |
| Other Criteria                      | All uses (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **CLOBAZAM**

#### **Products Affected**

• Clobazam SUSP 2.5MG/ML

#### • Clobazam TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Lennox-Gastaut syndrome: Diagnosis of Lennox-Gastaut syndrome. Used for the adjunctive treatment of seizures associated with Lennox-Gastaut syndrome. Dravet syndrome: Diagnosis of seizures associated with Dravet syndrome (DS). Used in combination with Diacomit. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **CLONIDINE ER**

### **Products Affected**

• Clonidine Hydrochloride Er TB12

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.                           |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of attention deficit hyperactivity disorder (ADHD). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **COBENFY**

### **Products Affected**

• Cobenfy

• Cobenfy Starter Pack

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of schizophrenia. Trial and failure, contraindication, or intolerance to two of the following: a) aripiprazole, b) asenapine, c) olanzapine, d) paliperidone, e) quetiapine (IR or ER), f) risperidone, or g) ziprasidone. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **C**OMETRIQ

### **Products Affected**

#### • Cometriq

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Medullary Thyroid Cancer (MTC): Diagnosis of medullary thyroid cancer (MTC). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.       |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                 |

# **C**OPIKTRA

### **Products Affected**

### • Copiktra

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL. Disease is relapsed or refractory. Trial and failure, contraindication, or intolerance to at least two prior therapies for CLL/SLL [e.g., Leukeran (chlorambucil), Gazyva (obinutuzumab), Arzerra (ofatumumab), Bendeka (bendamustine), Imbruvica (ibrutinib), etc.]. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **CORLANOR**

### **Products Affected**

### • Ivabradine Hydrochloride

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **COSENTYX**

#### **Products Affected**

• Cosentyx Unoready

- Cosentyx INJ 150MG/ML, 75MG/0.5ML
- Cosentyx Sensoready Pen

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Plaque psoriasis (Initial): Diagnosis of moderate to severe plaque psoriasis. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance (TF/C/I) to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Psoriatic Arthritis (PsA) (Initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Ankylosing Spondylitis (AS) (Initial): Diagnosis of active AS. Minimum duration of a one-month TF/C/I to one non-steroidal anti-inflammatory drug (NSAID) (eg, ibuprofen, naproxen) at maximally tolerated doses. Non-radiographic axial spondyloarthritis (nr-axSpA, initial): Dx of active nr-axSpA with signs of inflammation. Minimum duration of a one-month TF/C/I to two non-steroidal anti-inflammatory drugs (NSAIDs) (eg, ibuprofen, naproxen) at maximally tolerated doses. Hidradenitis suppurativa (HS) (Initial): Diagnosis of moderate to severe HS. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | Plaque psoriasis, HS (Initial): Prescribed by or in consultation with a dermatologist. Psoriatic Arthritis (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. AS, nr-axSpA, ERA (initial): Prescribed by or in consultation with a rheumatologist.  |
| Coverage<br>Duration               | All uses (Initial, Reauth): plan year.   |

| Other Criteria                      | Enthesitis-Related Arthritis (ERA) (Initial): Diagnosis of active ERA. Minimum duration of a one-month TF/C/I to two NSAIDs (eg, ibuprofen, naproxen) at maximally tolerated doses. All indications (Reauth): Patient demonstrates positive clinical response to therapy. |
|-------------------------------------|---|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **C**OTELLIC

### **Products Affected**

#### • Cotellic

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Melanoma: Diagnosis of unresectable or metastatic melanoma. Disease is positive for BRAF V600E or V600K mutation. Used in combination with Zelboraf (vemurafenib). Histiocytic Neoplasm: Diagnosis of histiocytic neoplasm. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# CRESEMBA ORAL

### **Products Affected**

#### • Cresemba CAPS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.                           |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of invasive aspergillosis or invasive mucormycosis. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | 3 months  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **CRINONE**

### **Products Affected**

#### • Crinone GEL 8%

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | All indications: Excluded if for fertility uses.   |
| Required<br>Medical<br>Information  | Secondary amenorrhea: Diagnosis of secondary amenorrhea (the absence of menses in women who have already started menstruation who are not pregnant, breastfeeding, or in menopause). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **C**TEXLI

### **Products Affected**

• Ctexli

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **DANZITEN**

### **Products Affected**

#### • Danziten

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                            |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of chronic myeloid leukemia.                         |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | For continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **D**AURISMO

### **Products Affected**

#### • Daurismo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute myeloid leukemia (AML): Diagnosis of newly-diagnosed acute myeloid leukemia (AML) AND Daurismo therapy to be given in combination with low-dose cytarabine AND One of the following: 1) Patient is greater than or equal to 75 years old, or 2) Patient has significant comorbidities that preclude the use of intensive induction chemotherapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **D**EFERASIROX

#### **Products Affected**

• Deferasirox TABS

• Deferasirox TBSO

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Chronic Iron Overload due to Blood Transfusions (initial): Diagnosis of chronic iron overload (eg, sickle cell anemia, thalassemia, etc.) due to blood transfusion. Patient has blood transfusion of at least 100 mL/kg of packed red blood cells (eg, at least 20 units of packed red blood cells for a 40-kg person or more in individuals weighing more than 40 kg) prior to initiation of treatment with deferasirox. Patient has serum ferritin levels consistently greater than 1000 mcg/L prior to initiation of treatment with deferasirox. Chronic Overload in non-transfusion dependent thalassemia syndromes (initial): Diagnosis of chronic iron overload in non-transfusion dependent thalassemia syndrome. Patient has liver iron (Fe) concentration (LIC) levels consistently greater than or equal to 5 mg Fe per gram of dry weight prior to initiation of treatment with deferasirox. Patient has serum ferritin levels consistently more than 300 mcg/L prior to initiation of treatment with deferasirox. Chronic Iron Overload due to Blood Transfusions, Chronic Overload in non-transfusion dependent thalassemia syndromes (reauthorization): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Initial: Prescribed by a hematologist/oncologist or hepatologist.   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **DEGARELIX**

### **Products Affected**

• Firmagon INJ 120MG/VIAL, 80MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prostate Cancer: Diagnosis of prostate cancer.                         |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# DICLOFENAC GEL 3%

#### **Products Affected**

• Diclofenac Sodium GEL 3%

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Actinic Keratosis (initial): Diagnosis of Actinic Keratosis. Actinic Keratosis (reauthorization): Patient demonstrates positive clinical response to therapy. At least 30 days have elapsed since cessation of diclofenac sodium 3% topical gel therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 90 days  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **DICLOFENAC TOPICAL SOLUTION 1.5%**

#### **Products Affected**

• Diclofenac Sodium EXTERNAL SOLN 1.5%

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                          |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of osteoarthritis of the knee(s).                  |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug. |

# **DOPTELET**

### **Products Affected**

• Doptelet

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Thrombocytopenia Prior to Planned Procedure (TPPP): Diagnosis (dx) of thrombocytopenia. Patient has chronic liver disease and is scheduled to undergo a procedure. Baseline platelet count is less than 50,000/mcL. Chronic Immune Thrombocytopenia (ITP) (initial): Diagnosis of chronic ITP or relapsed/refractory ITP. Baseline platelet count is less than 30,000/mcL. Trial and failure, contraindication, or intolerance to at least one of the following: corticosteroids (e.g., prednisone, methylprednisolone), immunoglobulins [e.g., Gammagard, immune globulin (human)], or splenectomy. Patient's degree of thrombocytopenia and clinical condition increase the risk of bleeding. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | TPPP: 1 month. ITP (initial, reauth): Plan year   |
| Other Criteria                      | ITP (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **DRONABINOL**

### **Products Affected**

#### • Dronabinol

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **D**ROXIDOPA

### **Products Affected**

### • Droxidopa

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Neurogenic orthostatic hypotension (NOH): (Initial): Diagnosis of symptomatic NOH. NOH is caused by one of the following conditions: primary autonomic failure (eg, Parkinson's disease, multiple system atrophy, pure autonomic failure), dopamine beta-hydroxylase deficiency, non-diabetic autonomic neuropathy. (Reauth): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | NOH (init): Prescribed by or in consultation with one of the following specialists: cardiologist, neurologist, nephrologist.  |
| Coverage<br>Duration                | Initial: 1 month. Reauth: plan year   |
| Other Criteria                      | Trial and failure, contraindication, or intolerance to one of the following agents: Fludrocortisone acetate, midodrine.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **D**UPIXENT

### **Products Affected**

• Dupixent INJ 200MG/1.14ML, 300MG/2ML

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **EBGLYSS**

### **Products Affected**

• Ebglyss

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Initial: Diagnosis of moderate to severe atopic dermatitis. Patient weighs at least 40 kg. One of the following: a) Involvement of at least 10% body surface area (BSA) or b) SCORing Atopic Dermatitis (SCORAD) index value of at least 25. Trial and failure, contraindication, or intolerance to one of the following: Pimecrolimus cream or Tacrolimus ointment. |
| Age Restrictions                    | Initial: Patient is 12 years of age or older.  |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with a dermatologist or allergist/immunologist.  |
| Coverage<br>Duration                | Initial, Reauth: Plan Year.  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **ELIGARD**

### **Products Affected**

### • Eligard

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prostate Cancer: Treatment of advanced prostate cancer.                |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# **E**MFLAZA

### **Products Affected**

#### • Deflazacort

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Initial: Submission of medical records (e.g., chart notes, laboratory values) documenting diagnosis of Duchenne muscular dystrophy (DMD). Patient has received genetic testing for a mutation of the dystrophin gene. Submission of medical records (e.g., chart notes, laboratory values) documenting one of the following: A) Documentation of a confirmed mutation of the dystrophin gene or B) Muscle biopsy confirmed an absence of dystrophin protein. Patient has had a trial and failure or intolerance to prednisone or prednisolone. |
| Age Restrictions                    | Initial: 2 years of age or older   |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with a neurologist.  |
| Coverage<br>Duration                | Initial, Reauth: Plan year   |
| Other Criteria                      | Reauth: Patient has experienced a benefit from therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### **E**MGALITY

### **Products Affected**

### • Emgality

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Episodic Migraines (EM) (120 mg/mL strength only) (initial): Diagnosis of EM. Patient has greater than or equal to 4 migraine days per month. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines. Chronic Migraines (CM) (120 mg/mL strength only) (initial): Diagnosis of CM. Patient has greater than or equal to 8 migraine days per month. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines. Episodic Cluster Headaches (ECH) (100 mg/mL strength only) (initial): Diagnosis of ECH. Patient has experienced at least 2 cluster periods lasting from 7 days to 365 days, separated by pain-free periods lasting at least three months. Medication will not be used in combination with another injectable CGRP inhibitor. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | EM, CM, ECH (initial, reauth): Plan year.   |
| Other Criteria                      | ECH (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another injectable CGRP inhibitor. EM, CM (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### **ENBREL**

#### **Products Affected**

- Enbrel INJ 25MG/0.5ML, 50MG/ML
- Enbrel Mini
- Enbrel Sureclick

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Rheumatoid Arthritis (RA) (Initial): Diagnosis of moderately to severely active RA. Minimum duration of a 3-month trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies at maximally tolerated doses: methotrexate, leflunomide, sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Diagnosis of moderately to severely active PJIA. Minimum duration of a 6-week trial and failure, contraindication, or intolerance to one of the following conventional therapies at maximally tolerated doses: leflunomide or methotrexate. Psoriatic Arthritis (PsA) (Initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Plaque psoriasis (Initial): Diagnosis of moderate to severe chronic plaque psoriasis. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), C/I to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Ankylosing Spondylitis (AS) (Initial): Diagnosis of active AS. Minimum duration of a one-month TF/C/I to one nonsteroidal anti-inflammatory drug (NSAID) (eg, ibuprofen, naproxen) at maximally tolerated doses. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | RA, JIA, AS (Initial): Prescribed by or in consultation with a rheumatologist. PsA (Initial): Prescribed by or in consultation with a rheumatologist or dermatologist. Plaque Psoriasis (Initial): Prescribed by or in consultation with a dermatologist.   |

| Coverage<br>Duration                | All indications (initial, reauth): plan year  |
|-------------------------------------|---|
| Other Criteria                      | All indications (Reauth): Patient demonstrates positive clinical response to therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.                              |

# **ENDARI**

## **Products Affected**

## • L-glutamine PACK

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Sickle cell disease (initial): Diagnosis of sickle cell disease. Used to reduce acute complications of sickle cell disease. Patient has had 2 or more painful sickle cell crises within the past 12 months. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Sickle cell disease (initial, reauth): plan year  |
| Other Criteria                      | Sickle cell disease (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **EPCLUSA**

## **Products Affected**

## • Sofosbuvir/velpatasvir

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **E**PIDIOLEX

## **Products Affected**

## • Epidiolex

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Lennox-Gastaut syndrome (LGS): Diagnosis of seizures associated with LGS. Trial of, contraindication, or intolerance to two formulary anticonvulsants (e.g., topiramate, lamotrigine, valproate). Dravet syndrome (DS): Diagnosis of seizures associated with DS. Tuberous sclerosis complex (TSC): Diagnosis of seizures associated with TSC. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **EPOETIN ALFA (PREFERRED)**

## **Products Affected**

• Procrit

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Anemia due to Chronic Kidney Disease (CKD) (Initial): Diagnosis (dx) of CKD. Anemia by lab values (Hct less than 30% or Hgb less than 10 g/dL) within 30 days of request. The rate of hemoglobin decline indicates the likelihood of requiring a red blood cell (RBC) transfusion, and reducing the risk of alloimmunization and/or other RBC transfusion-related risks is a goal. (Reauth): Diagnosis of CKD. Most recent or average (avg) Hct over 3 months is 33% or less (Hgb 11 g/dL or less) for patients on dialysis, without ESRD OR Most recent or average (avg) Hct over 3 months is 30% or less (Hgb 10 g/dL or less) for patients not on dialysis OR Most recent or average (avg) Hct over 3 months is 36% or less (Hgb 12 g/dL or less) for pediatric patients. Patient demonstrates positive clinical response to therapy from pre-treatment level. Anemia w/ HIV (Initial): Anemia by labs (Hgb less than 12 g/dL or Hct less than 36%) within 30 days of request. Serum erythropoietin level less than or equal to 500 mU/mL. Receiving zidovudine therapy or dx of HIV. (Reauth): Most recent or avg Hct over 3 months is below 36% or most recent or avg Hgb over 3 months is below 12 g/dl. Patient demonstrates positive clinical response to therapy from pre-treatment level. Anemia with chemo (Initial):Other causes of anemia ruled out. Anemia with labs (Hct less than 30%, Hgb less than 10 g/dL) within prior 2 weeks of request. Cancer is non-myeloid malignancy. Patient is receiving chemo. (Reauth): Anemia by labs (Hgb less than 10 g/dl or Hct less than 30%) collected within the prior 2 wks of request. Patient demonstrates positive clinical response to therapy from pre-treatment level. Patient is receiving chemo. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | CKD,HIV(Init):6mo.(reauth):plan yr.Chemo(init, reauth):3mo.MDS(init):3mo,(reauth):plan yr.Preop:1mo.   |

#### Other Criteria Anemia in Myelodysplastic Syndrome (MDS) (Init): Diagnosis of MDS. Serum erythropoietin 500 mU/mL or less, or transfusion dependent MDS. (Reauth): Most recent or avg Hct over 3 months was 36% or less, OR most recent or avg Hgb over 3 months is 12 g/dl or less. Patient demonstrates positive clinical response to therapy from pre-treatment level. Preoperative for reduction of allogeneic blood transfusion: Patient is scheduled to undergo elective, non-cardiac, non-vascular surgery. Hgb is greater than 10 to less than or equal to 13 g/dL. Patient is at high risk for perioperative transfusions. Patient is unwilling or unable to donate autologous blood pre-operatively. ESRD patients: Coverage is excluded under Medicare Part D for patients with ESRD on dialysis for any indication related or unrelated to treatment of ESRD since the payment for the drug is included in the ESRD PPS payment bundle. NON-ESRD PATIENTS for the following indications: Off-label uses (except Anemia in MDS and Hep C patients being treated with combo ribavirin and interferon/peginterferon): Will not be approved if patient has Hgb greater than 10 g/dL or Hct greater than 30%.CKD (init, reauth), Chemo (init), preop, MDS (init): Adequate iron stores confirmed by both of the following: a) Patient's ferritin level is greater than 100 mcg/L and b) Patient's transferrin saturation (TSAT) is greater than 20%.

Criteria DOES NOT require use of a prerequisite Part D drug.

**Prerequisite** 

Therapy Required

# **E**RIVEDGE

## **Products Affected**

## • Erivedge

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Metastatic basal cell carcinoma (BCC): Diagnosis of metastatic basal cell carcinoma. Advanced basal cell carcinoma: Diagnosis of locally advanced basal cell carcinoma. One of the following: Cancer has recurred following surgery, Patient is not a candidate for surgery, or Patient is not a candidate for radiation. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **E**RLEADA

## **Products Affected**

#### • Erleada

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-metastatic castration-resistant prostate cancer (NM-CRPC): Diagnosis of non-metastatic, castration-resistant (chemical or surgical) or recurrent prostate cancer. One of the following: 1) Used in combination with a gonadotropin-releasing hormone (GnRH) analog (e.g., Zoladex [goserelin], Vantas [histrelin], Lupron [leuprolide], Trelstar [triptorelin], Firmagon [degarelix]) OR 2) Patient received a bilateral orchiectomy. Metastatic castration-sensitive prostate cancer (M-CSPC): Diagnosis of metastatic, castration-sensitive or naive prostate cancer. One of the following: 1) Used in combination with a gonadotropin-releasing hormone (GnRH) analog (e.g., Lupron[leuprolide], Trelstar[triptorelin], Zoladex [goserelin], Vantas [histrelin], Firmagon [degarelix]) OR 2) Patient received bilateral orchiectomy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **E**RLOTINIB

## **Products Affected**

## • Erlotinib Hydrochloride TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-Small Cell Lung Cancer (NSCLC): 1) Diagnosis of NSCLC. Disease is one of the following: a) advanced, b) metastatic, or c) recurrent. Tumors are positive for epidermal growth factor receptor (EGFR) exon 19 deletion mutations or exon 21 (L858R) substitution mutations or a known sensitizing EGFR mutation (e.g., in-frame exon 20 insertions, exon 18 G719 mutation, exon 21 L861Q mutation). Pancreatic cancer: Diagnosis of locally advanced, unresectable or metastatic pancreatic cancer. Used in combination with Gemzar (gemcitabine). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **EVEROLIMUS**

## **Products Affected**

• Everolimus TABS 10MG, 2.5MG, 5MG, 7.5MG

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Advanced Neuroendocrine Tumors (NET): Diagnosis (Dx) of neuroendocrine tumors of pancreatic origin, gastrointestinal origin, lung origin, or thymic origin. Disease is progressive. Disease is unresectable, locally advanced or metastatic. Advanced Renal Cell Carcinoma/Kidney Cancer: Dx of advanced renal cell cancer/kidney cancer. Disease is one of the following: (1) relapsed or (2) stage IV disease. Renal angiomyolipoma with tuberous sclerosis complex (TSC): Dx of renal angiomyolipoma and TSC, not requiring immediate surgery. Subependymal Giant Cell Astrocytoma (SEGA) with tuberous sclerosis (TS): Dx of SEGA associated with TS. Patient is not a candidate for curative surgical resection. Breast Cancer: Dx of recurrent or metastatic breast cancer. One of the following: Disease is hormone receptor positive (HR+) [i.e., estrogen-receptor-positive (ER+) or progesterone-receptor-positive (PR+)] OR both of the following: disease is HR- and disease has clinical characteristics that predict a HR+ tumor. Disease is HER2-negative. One of the following: Patient is a postmenopausal woman, patient is a premenopausal woman being treated with ovarian ablation/suppression, or patient is male. One of the following: A) Both of the following: a) one of the following: 1) Disease progressed while on or within 12 months of non-steroidal aromatase inhibitor [e.g., Arimidex (anastrozole), Femara (letrozole)] therapy or 2) patient was treated with tamoxifen at any time AND b) Used in combination with Aromasin (exemestane) OR B) Used in combination with Fulvestrant or Tamoxifen. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan year  |

| Other Criteria                      | All uses: Approve for continuation of prior therapy if within the past 120 days. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.                         |

# **EVEROLIMUS SOLUTION**

## **Products Affected**

#### • Everolimus TBSO

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Advanced Neuroendocrine Tumors (NET): Diagnosis (Dx) of neuroendocrine tumors of pancreatic origin, gastrointestinal origin, lung origin, or thymic origin. Disease is progressive. Disease is unresectable, locally advanced or metastatic. Advanced Renal Cell Carcinoma/Kidney Cancer: Dx of advanced renal cell cancer/kidney cancer. Disease is one of the following: (1) relapsed or (2) stage IV disease. Renal angiomyolipoma with tuberous sclerosis complex (TSC): Dx of renal angiomyolipoma and TSC, not requiring immediate surgery. Subependymal Giant Cell Astrocytoma (SEGA) with tuberous sclerosis (TS): Dx of SEGA associated with TS. Patient is not a candidate for curative surgical resection. Breast Cancer: Dx of recurrent or metastatic breast cancer. One of the following: Disease is hormone receptor positive (HR+) [i.e., estrogen-receptor-positive (ER+) or progesterone-receptor-positive (PR+)] OR both of the following: disease is HR- and disease has clinical characteristics that predict a HR+ tumor. Disease is HER2-negative. One of the following: Patient is a postmenopausal woman, patient is a premenopausal woman being treated with ovarian ablation/suppression, or patient is male. One of the following: A) Both of the following: a) one of the following: 1) Disease progressed while on or within 12 months of non-steroidal aromatase inhibitor [e.g., Arimidex (anastrozole), Femara (letrozole)] therapy or 2) patient was treated with tamoxifen at any time AND b) Used in combination with Aromasin (exemestane) OR B) Used in combination with Fulvestrant or Tamoxifen. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan year  |

| Other Criteria                      | TSC Associated Partial-Onset Seizures: Dx of TSC associated partial-onset seizures. Used as adjunctive therapy. All uses: Approve for continuation of prior therapy if within the past 120 days. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **FASENRA**

## **Products Affected**

• Fasenra

#### • Fasenra Pen

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Severe asthma (Initial): Diagnosis of severe asthma. Asthma is an eosinophilic phenotype as defined by a baseline (pre-treatment) peripheral blood eosinophil level greater than or equal to 150 cells per microliter. One of the following: a) Patient has had two or more asthma exacerbations requiring systemic corticosteroids (e.g., prednisone) within the past 12 months, OR b) Prior asthma-related hospitalization within the past 12 months. One of the following: 1) Both of the following: a) Patient is 6 years of age or older but less than 12 years of age, and b) Patient is currently being treated with one of the following unless there is a contraindication or intolerance to these medications: i) Medium-dose inhaled corticosteroid [ICS] (eg, greater than 100 - 200 mcg fluticasone propionate equivalent/day) AND Additional asthma controller medication (eg, leukotriene receptor antagonist [LTRA] [eg, montelukast], long-acting beta-2 agonist [LABA] [eg, salmeterol], long-acting muscarinic antagonist [LAMA] [eg, tiotropium]), OR ii) One medium dosed combination ICS/LABA product (eg, Wixela Inhub [fluticasone propionate 100mcg/salmeterol 50mcg], budesonide 80mcg/formoterol 4.5mcg, Breo Ellipta [fluticasone furoate 50 mcg/vilanterol 25 mcg]), OR 2) Both of the following: a) Patient is 12 years of age or older, and b) Patient is currently being treated with one of the following unless there is a contraindication or intolerance to these medications: a) Both of the following: 1) High-dose ICS (eg, greater than 500 mcg fluticasone propionate equivalent/day) and 2) additional asthma controller medication (e.g., LTRA [eg, montelukast], LABA [e.g., salmeterol], tiotropium) OR b) One maximally-dosed combination ICS/LABA product [e.g., Wixela (fluticasone propionate 500mcg/salmeterol 50mcg), budesonide 160mcg/formoterol 4.5mcg, Breo Ellipta (fluticasone 200mcg/vilanterol 25mcg)]. |
| Age Restrictions                   | N/A  |

| Prescriber<br>Restrictions          | Severe Asthma: Initial: Prescribed by or in consultation with a pulmonologist or allergy/immunology specialist. EGPA (init): Prescribed by or in consultation with pulmonologist, rheumatologist, or allergist/immunologist  |
|-------------------------------------|--|
| Coverage<br>Duration                | Severe asthma: Initial: 6 mo, Reauth: Plan year EPGA (init, reauth): Plan year.  |
| Other Criteria                      | Eosinophilic granulomatosis with polyangiitis (EGPA) (initial): Diagnosis of EGPA. Patient's disease has relapsed or is refractory to standard of care therapy (ie, corticosteroid treatment with or without immunosuppressive therapy). Patient is currently receiving corticosteroid therapy (eg, prednisolone, prednisone) unless there is a contraindication or intolerance to corticosteroid therapy. EPGA (reauth): Patient demonstrates positive clinical response to therapy. Severe asthma (Reauth): Patient demonstrates positive clinical response to therapy. Patient continues to be treated with an inhaled corticosteroid (ICS) (e.g., fluticasone, budesonide) with or without additional asthma controller medication (e.g., leukotriene receptor antagonist [e.g., montelukast], long-acting beta-2 agonist [LABA] [e.g., salmeterol], tiotropium) unless there is a contraindication or intolerance to these medications. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **FINTEPLA**

## **Products Affected**

## • Fintepla

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | One of the following: 1) Diagnosis of seizures associated with Dravet syndrome, OR 2) Diagnosis of seizures associated with Lennox-Gastaut syndrome. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a neurologist  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **FLUCYTOSINE**

## **Products Affected**

• Flucytosine CAPS 250MG, 500MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **FOTIVDA**

## **Products Affected**

#### • Fotivda

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of advanced renal cell carcinoma. Disease is one of the following: relapsed or refractory. Patient has received two or more prior systemic therapies (e.g., chemotherapy). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# FRUZAQLA

## **Products Affected**

## • Fruzaqla

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of colorectal cancer. Disease is one of the following: Advanced or Metastatic. Patient has been previously treated with both of the following: A) Fluoropyrimidine-, oxaliplatin-, irinotecan-based chemotherapy, and B) Anti-VEGF biological therapy (e.g., bevacizumab, ramucirumab). One of the following: A) Patient has RAS mutant tumors, OR B) Both of the following: a) Patient has RAS wild type tumors, AND b) Patient has been previously treated with an anti-EGFR biological therapy (e.g., panitumumab, cetuximab) |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **GAVRETO**

## **Products Affected**

#### • Gavreto

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: recurrent, advanced, or metastatic. Presence of RET (rearranged during transfection) gene fusion-positive or RET rearrangement positive tumors. Thyroid Cancer: Diagnosis of thyroid cancer. Disease is one of the following: advanced or metastatic. Disease is rearranged during transfection (RET) gene fusion-positive. Disease requires treatment with systemic therapy. One of the following: patient is radioactive iodine-refractory or radioactive iodine therapy is not appropriate. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **G**ILOTRIF

## **Products Affected**

#### • Gilotrif

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of metastatic NSCLC. One of the following: 1) Tumors have non-resistant epidermal growth factor (EGFR) mutations as detected by an FDA-approved test OR 2) squamous disease progressing after previous platinum-based chemotherapy (e.g., cisplatin, carboplatin) OR 3) tumors are positive for a known sensitizing EGFR mutation (e.g., in-frame exon 20 insertions, exon 18 G719 mutation, exon 21 L861Q mutation). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **G**LYCOPYRROLATE

## **Products Affected**

• Glycopyrrolate TABS 1MG, 2MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of peptic ulcer. One of the following: 1) Patient is receiving concomitant treatment therapy with a proton-pump inhibitor (PPI) (e.g., lansoprazole, omeprazole), OR 2) Both of the following: a) Patient has a contraindication or intolerance to PPIs, and b) Patient is receiving concomitant treatment therapy with an H2-receptor antagonist (e.g., famotidine, nizatidine).  |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Initial, Reauth: Prescribed by or in consultation with a gastroenterologist.  |
| Coverage<br>Duration                | Initial, Reauth: 3 months.  |
| Other Criteria                      | Reauth: One of the following: 1) Patient's peptic ulcer has not healed, OR 2) Patient has a new peptic ulcer. One of the following: 1) Patient is receiving concomitant treatment therapy with a proton-pump inhibitor (PPI) (e.g., lansoprazole, omeprazole), OR 2) Both of the following: a) Patient has a contraindication or intolerance to PPIs, and b) Patient is receiving concomitant treatment therapy with an H2-receptor antagonist (e.g., famotidine, nizatidine). Patient experienced a reduction in peptic ulcer symptoms while on therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **G**OMEKLI

## **Products Affected**

#### • Gomekli

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **GROWTH HORMONES**

### **Products Affected**

• Genotropin

• Genotropin Miniquick

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# HAEGARDA

## **Products Affected**

## Haegarda

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prophylaxis of hereditary angioedema (HAE) attacks (initial): Diagnosis of HAE. Diagnosis has been confirmed by both of the following: 1) C4 level below the lower limit of normal, and 2) C1 inhibitor (C1-INh) deficiency or dysfunction (Type I or II HAE) as documented by ONE of the following: a) C1-INH antigenic level below the lower limit of normal OR b) C1-INH functional level below the lower limit of normal. For prophylaxis against HAE attacks. Not used in combination with other approved treatments for prophylaxis against HAE attacks. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | HAE (prophylaxis) (initial): Prescribed by or in consultation with an immunologist or an allergist   |
| Coverage<br>Duration                | HAE (prophylaxis) (initial, reauth): Plan year   |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy. Not used in combination with other approved treatments for prophylaxis against HAE attacks   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **HERNEXEOS**

## **Products Affected**

#### • Hernexeos

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **IBRANCE**

## **Products Affected**

#### • Ibrance

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Breast cancer: Diagnosis of breast cancer.                             |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# **IBTROZI**

## **Products Affected**

• Ibtrozi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **ICATIBANT**

## **Products Affected**

#### • Icatibant Acetate

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Treatment of hereditary angioedema (HAE) attacks (initial): Diagnosis of HAE. Diagnosis has been confirmed by both of the following: 1) C4 level below the lower limit of normal, and 2) C1 inhibitor (C1-INh) deficiency or dysfunction (Type I or II HAE) as documented by ONE of the following: a) C1-INH antigenic level below the lower limit of normal OR b) C1-INH functional level below the lower limit of normal. For the treatment of acute HAE attacks. Not used in combination with other approved treatments for acute HAE attacks. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | HAE (initial): Prescribed by or in consultation with an immunologist or an allergist  |
| Coverage<br>Duration                | Initial, Reauth: Plan year  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy. Not used in combination with other approved treatments for acute HAE attacks  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **ICLUSIG**

## **Products Affected**

• Iclusig

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **IDHIFA**

## **Products Affected**

• Idhifa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute Myeloid Leukemia (AML): Diagnosis of AML. AML is isocitrate dehydrogenase-2 (IDH2) mutation-positive. One of the following: A) Disease is relapsed or refractory, B) Patient is not a candidate for or patient declines intensive induction therapy, C) Used for post induction therapy following response to low intensity induction therapy, or D) Used for consolidation therapy as continuation of low-intensity regimen used for induction. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **IMBRUVICA**

#### **Products Affected**

- Imbruvica CAPS
- Imbruvica SUSP

• Imbruvica TABS 140MG, 280MG, 420MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL. Waldenstrom's Macroglobulinemia (WM)/Lymphoplasmacytic Lymphoma (LPL): Diagnosis of WM/LPL. Chronic Graft Versus Host Disease (cGVHD): Diagnosis of cGVHD AND trial and failure of at least one other systemic therapy (e.g., corticosteroids like prednisone or methylprednisolone, mycophenolate, etc.). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **I**MKELDI

## **Products Affected**

#### • Imkeldi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Chronic Myelogenous/Myeloid Leukemia (CML): Diagnosis of CML. Acute Lymphoblastic Leukemia/Acute Lymphoblastic Lymphoma (ALL): Diagnosis of Philadelphia chromosome-positive ALL. Myelodysplastic Disease (MDS)/Myeloproliferative Disease (MPD): Diagnosis of MDS/MPD. Platelet-derived growth factor receptor (PDGFR) gene rearrangements. Aggressive Systemic Mastocytosis (ASM): Diagnosis of ASM. One of the following: a) KIT D816V mutation negative or unknown, b) Well-differentiated SM [WDSM], or c) Eosinophilia is present with FIP1L1-PDGFRA fusion gene. Hypereosinophilic Syndrome (HES) and/or Chronic Eosinophilic Leukemia (CEL): Diagnosis of at least one of the following: a) HES or b) CEL. Dermatofibrosarcoma Protuberans (DFSP): Diagnosis of unresectable, recurrent, or metastatic DFSP. Gastrointestinal Stromal Tumors (GIST): Diagnosis of GIST. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **I**NBRIJA

## **Products Affected**

• Inbrija

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Parkinson's disease (PD) (initial): Diagnosis of PD. Patient is experiencing intermittent OFF episodes. Used in combination with carbidopa/levodopa. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | PD (initial, reauth): Plan year  |
| Other Criteria                      | PD (reauth): Patient demonstrates positive clinical response to therapy. Used in combination with carbidopa/levodopa.                                |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **INGREZZA**

## **Products Affected**

• Ingrezza

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Tardive Dyskinesia: Diagnosis of tardive dyskinesia. One of the following: a) patient has persistent symptoms of tardive dyskinesia despite a trial of dose reduction, tapering, or discontinuation of the offending medication OR b) patient is not a candidate for a trial of dose reduction, tapering, or discontinuation of the offending medication. Chorea associated with Huntington's disease: Diagnosis of chorea in patients with Huntington's disease. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Tardive Dyskinesia: Prescribed by or in consultation with a neurologist or psychiatrist. Chorea associated with Huntington's disease: Prescribed by or in consultation with a neurologist.  |
| Coverage<br>Duration                | All Indications: Plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **INLYTA**

## **Products Affected**

• Inlyta

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Advanced Renal Cell Carcinoma: Diagnosis of renal cell cancer. One of the following: (1) diagnosis of stage IV disease, or (2) Both of the following: a) Disease is advanced, and b) One of the following: i) Patient has failed one prior systemic therapy (e.g., chemotherapy), or ii) Inlyta will be used in combination with Bavencio (avelumab) or Keytruda (pembrolizumab). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# INQOVI

#### **Products Affected**

• Inqovi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | One of the following: 1) Both of the following: a) Diagnosis of myelodysplastic syndrome (MDS) and b) Patient is intermediate-1, intermediate-2, or high-risk per the International Prognostic Scoring System (IPSS), OR 2) Diagnosis of chronic myelomonocytic leukemia (CMML). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **INREBIC**

### **Products Affected**

#### • Inrebic

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Myelofibrosis: Diagnosis of one of the following: primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# INSULIN - LIKE GROWTH FACTOR

#### **Products Affected**

#### • Increlex

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial therapy: IGF-1 deficiency: Diagnosis of severe primary IGF-1 deficiency. Height standard deviation score of -3.0 or less. Basal IGF-1 standard deviation score of -3.0 or less. Normal or elevated growth hormone (GH). GH gene deletion: Diagnosis of growth hormone gene deletion who have developed neutralizing antibodies to GH. Reauthorization: Patient demonstrates positive clinical response to therapy. Both of the following: (1) Expected adult height is not obtained and (2) Documentation of expected adult height goal |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with an endocrinologist   |
| Coverage<br>Duration                | Initial, reauth: plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### **IRESSA**

### **Products Affected**

#### • Gefitinib

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of metastatic NSCLC, and One of the following: tumors are positive for epidermal growth factor receptor (EGFR) exon 19 deletions or tumors are positive for EGFR exon 21 (L858R) substitution mutations or tumors are positive for a known sensitizing EGFR mutation (e.g., in-frame exon 20 insertions, exon 18 G719 mutation, exon 21 L861Q mutation). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **ISOTRETINOIN**

#### **Products Affected**

- Accutane CAPS 10MG, 20MG, 40MG
- Amnesteem

- Claravis
- Isotretinoin CAPS
- Zenatane

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | (initial): Diagnosis of severe recalcitrant nodular acne unresponsive to conventional therapy OR diagnosis of treatment resistant acne. Trial and failure, contraindication or intolerance to an adequate trial (at least 6 weeks) on two of the following conventional therapy regimens: a) topical retinoid or retinoid-like agent [eg, Retin-A/Retin-A Micro (tretinoin)], b) oral antibiotic [eg, Ery-Tab (erythromycin), Minocin (minocycline)], c) topical antibiotic with or without benzoyl peroxide [eg, Cleocin-T (clindamycin), erythromycin, BenzaClin (benzoyl peroxide/clindamycin), Benzamycin (benzoyl peroxide/erythromycin)]. Retreatment (Reauthorization): After greater than or equal to 2 months off therapy, persistent or recurring severe recalcitrant nodular acne is still present. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Initial: 6 months. Reauth: Retreatment - 6 months  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### **ITOVEBI**

### **Products Affected**

#### • Itovebi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of breast cancer. Disease is one of the following: a) Locally advanced, or b) Metastatic. Disease is all of the following: a) PIK3CA-mutated, b) Hormone receptor (HR)-positive, and c) Human epidermal growth-factor receptor 2 (HER2)-negative. Used following recurrence on or after completing adjuvant endocrine therapy (e.g. Zoladex [goserelin], Arimidex [anastrozole], Nolvadex [tamoxifen]). Used in combination with both of the following: a) Ibrance (Palbociclib), and b) Fulvestrant. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# ITRACONAZOLE (CAPSULES)

### **Products Affected**

#### • Itraconazole CAPS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Systemic Fungal Infections: Diagnosis of blastomycosis, histoplasmosis, or aspergillosis. Onychomycosis: Diagnosis of fingernail or toenail onychomycosis confirmed by one of the following: KOH test, fungal culture, or nail biopsy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Systemic Fungal infxns: plan year. Onychomycosis: (Fingernail) 2 mo. (Toenail) 3 mo.   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **IVERMECTIN TABLETS**

#### **Products Affected**

#### • Ivermectin TABS 3MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Any medically accepted indication [e.g., Onchocerciasis due to nematode parasite, Pediculosis, Strongyloidiasis, Ascariasis, Scabies (including crusted scabies), Cutaneous larva migrans (hook worm disease), Enterobiasis, Filariasis, Trichuriasis, or Gnathostomiasis]. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | 1 month   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# IVIG (PREFERRED)

#### **Products Affected**

• Gamunex-c INJ 1GM/10ML

• Octagam INJ 1GM/20ML, 2GM/20ML

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Primary immunodeficiency syndrome (PIS): patients with PIS. Idiopathic Thrombocytopenic Purpura (ITP): diagnosis (dx) of ITP. TF/C/I to corticosteroid therapy (e.g., prednisone, dexamethasone) OR platelet count of less than 30,000 cells/mm3. Kawasaki disease (KD): dx of KD. B-cell Chronic Lymphocytic Leukemia (CLL): dx of B-Cell CLL. Doc hypogammaglobulinemia (IgG less than 500mg/dL) or history of bacterial infections associated with B-cell CLL. Bone Marrow Transplant (BMT): Confirmed allogeneic BMT within the last 100 days. Doc severe hypogammaglobulinemia (IgG less than 400 mg/dL). HIV:dx of HIV. Doc hypogammaglobulinemia (IgG less than 400 mg/dL) or functional antibody deficiency demonstrated by poor specific antibody titers or recurrent bacterial infections. Guillain-Barre Syndrome (GBS) initial: dx of GBS. severe disease requiring aid to walk. Onset of neuropathic symptoms in the last 4 weeks. Myasthenia Gravis (MG): dx of generalized MG. Evidence of myasthenic exacerbation, defined by 1 of the following sxs in the last month: difficulty swallowing, acute respiratory failure, or major functional disability responsible for the discontinuation of physical activity. Dermatomyositis and Polymyositis (D/P) initial: dx of dermatomyositis or polymyositis. Trial and failure, contraindication or intolerance (TF/C/I) to immunosuppressive tx (eg corticosteroids, methotrexate, azathioprine, cyclophosphamide). Stiff person syndrome (SPS) initial:dx of SPS. TF/C/I to GABAergic medication (eg, baclofen). Lambert-Eaton myasthenic syndrome (LEMS) initial: dx of LEMS. TF/C/I to immunomodulator monotherapy (eg, azathioprine, corticosteroids). |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | All uses: Prescribed by or in consultation with a physician who has specialized expertise in managing patients on immune globulin therapy (e.g., immunologist, hematologist, neurologist).   |

| Coverage |  |
|----------|--|
| Duration |  |

KD: 1 mo. GBS,CIDP (initial), MG: 3 mo. ITP: 6 mo. CIDP,GBS (reauth), other uses: plan year.

#### **Other Criteria**

Subject to Part B vs. Part D review. Pt does not meet criteria for Part B or patient is in a long-term care facility. PIS: Clinically significant functional deficiency of humoral immunity as evidenced by doc failure to produce antibodies to specific antigens or hx of significant recurrent infxns. Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) initial: dx of CIDP confirmed by: 1) progressive sxs present for at least 2 mo, 2) symptomatic polyradiculoneuropathy as indicated by progressive or relapsing motor impairment of more than 1 limb, OR Progressive or relapsing sensory impairment of more than 1 limb, 3) Electrophysiologic findings when 1 of the following criteria are present: Partial conduction block of 1 or more motor nerve, Reduced conduction velocity of 2 or more motor nerves, Prolonged distal latency of 2 or more motor nerves, Prolonged F-wave latencies of 2 or more motor nerves, Absence of F waves of 2 or motor nerves, Abnormal Temporal Dispersion of 2 or more motor nerves, Distal compound muscle action potential (CMAP) duration increase of 1 or more motor nerves. Multifocal motor neuropathy (MMN) initial: dx of MMN confirmed by all of the following: 1) weakness with slowly progressive or stepwise progressive course over at least 1 month, 2) asymmetric involvement of 2 or more nerves, AND 3) absence of motor neuron signs and bulbar signs. CIDP, MMN reauth: documentation of positive clinical response to tx as measured by an objective scale [eg, Rankin, Modified Rankin, Medical Research Council (MRC) scale]. Documentation of titration to the minimum dose and frequency needed to maintain a sustained clinical effect. Relapsing remitting Multiple Sclerosis (MS) initial: dx of relapsing remitting form of MS (RRMS). Documentation of an MS exacerbation or progression (worsening) of the patient's clinical status from the visit prior to the one prompting the decision to initiate immune globulin tx. TF/c/I to 1 of the following: teriflunomide, interferon beta-1b, interferon beta-1a, glatiramer acetate, natalizumab, dimethyl fumarate, fingolimod. RRMS reauth: The prescriber maintains and provides chart documentation of the patient's evaluation, including all of the following: findings of interval examination including neurological deficits incurred, and assessment of disability (eg, Expanded Disability Status Score [EDSS], Functional Systems Score [FSS], Multiple Sclerosis Functional Composite [MSFC], Disease Steps [DS]). Stable or improved disability score (eg, EDSS, FSS, MSFC, DS). Documentation of decreased number of relapses since starting immune globulin tx. Dx continues to be the relapsing-remitting form of MS. Documentation of titration to the minimum dose and frequency needed to maintain a sustained clinical effect. GBS, D/P, SPS, LEMS reauth: Documentation of titration to the minimum dose and frequency needed to maintain a sustained clinical effect.

### **IWILFIN**

#### **Products Affected**

#### • Iwilfin

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of high-risk neuroblastoma (HRNB). Patient has shown at least a partial response to prior multiagent, multimodality therapy. Prior therapy included anti-GD2 immunotherapy (e.g., Danyelza [naxitamab-gqgk], Unituxin [dinutuximab]). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

### **J**AKAFI

### **Products Affected**

#### • Jakafi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Myelofibrosis: One of the following: Primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis. Polycythemia vera: Diagnosis of polycythemia vera and trial and failure, contraindication, or intolerance to one of the following: hydroxyurea or interferon therapy (e.g., Intron A, pegasys, etc.). Graft versus host disease (GVHD): One of the following: A) Both of the following: Diagnosis of acute (aGVHD). Disease is steroid-refractory, or B) Both of the following: Diagnosis of chronic graft versus host disease (cGVHD). Trial and failure of at least one or more lines of systemic therapy (e.g., corticosteroids, mycophenolate). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 6 months   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### **JAYPIRCA**

### **Products Affected**

• Jaypirca

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of mantle cell lymphoma (MCL). Disease is one of the following: a) relapsed, or b) refractory. Patient has received at least two prior systemic therapies [e.g., chemotherapy] for MCL, one of which is a Bruton Tyrosine Kinase (BTK) inhibitor therapy [e.g., Calquence (acalabrutinib), Imbruvica (ibrutinib)]. Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of one of the following: a) CLL, or b) SLL. Patient has received treatment for CLL/SLL with both of the following therapies: a) BTK inhibitor therapy [e.g., Calquence (acalabrutinib), Imbruvica (ibrutinib)], and b) B-cell lymphoma 2 (BCL-2) inhibitor therapy [e.g., Venclexta (venetoclax)]. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### **J**AYTHARI

#### **Products Affected**

• Jaythari

| PA Criteria                         | Criteria Details     |
|-------------------------------------|----------------------|
| Indications                         | Pending CMS Review   |
| Off-Label Uses                      | Pending CMS Review   |
| Exclusion<br>Criteria               | Pending CMS Review   |
| Required<br>Medical<br>Information  | Pending CMS Review   |
| Age Restrictions                    | Pending CMS Review   |
| Prescriber<br>Restrictions          | Pending CMS Review . |
| Coverage<br>Duration                | Pending CMS Review   |
| Other Criteria                      | Pending CMS Review   |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review   |

# **JYNARQUE**

#### **Products Affected**

#### • Tolvaptan TABS

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Autosomal dominant polycystic kidney disease (ADPKD) (initial): Diagnosis of rapidly progressing ADPKD. One of the following: 1) Both of the following: A) Patient is new to therapy or has received Jynarque for less than or equal to 18 months AND B) Alanine transaminase (ALT), aspartate transaminase (AST), and bilirubin will be measured prior to initiation, at 2 weeks and 4 weeks after initiation, then monthly for the first 18 months of therapy OR 2) Both of the following: A) Patient has received Jynarque for longer than 18 months AND B) ALT, AST, and bilirubin will be measured at least every 3 months. Patient does not have a history of significant liver impairment or injury, not including uncomplicated polycystic liver disease. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Plan year   |
| Other Criteria                     | ADPKD (reauth): Patient demonstrates positive clinical response to therapy. Patient does not have signs or symptoms consistent with hepatic injury, not including uncomplicated polycystic liver disease. One of the following: 1) Both of the following: A) Patient has received Jynarque for less than or equal to 18 months AND B) Alanine transaminase (ALT), aspartate transaminase (AST), and bilirubin will be measured prior to initiation, at 2 weeks and 4 weeks after initiation, then monthly for the first 18 months of therapy OR 2) Both of the following: A) Patient has received Jynarque for longer than 18 months AND B) ALT, AST, and bilirubin will be measured at least every 3 months.   |

| Prerequisite Criteria DOES NOT require use of a prerequisite Part D drug.  Therapy Required |  |
|---|--|
|---|--|

### KALYDECO

### **Products Affected**

### • Kalydeco

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Cystic Fibrosis (CF) (Initial): Diagnosis of CF. Submission of laboratory records confirming patient has at least one mutation in the CFTR gene that is responsive to ivacaftor potentiation. |
| Age Restrictions                    | CF (initial): Patient is 1 month of age or older.   |
| Prescriber<br>Restrictions          | CF (initial, reauth): Prescribed by or in consultation with a specialist affiliated with a CF care center or pulmonologist  |
| Coverage<br>Duration                | CF (initial, reauth): plan year   |
| Other Criteria                      | CF (reauth): Patient demonstrates positive clinical response to therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# KERENDIA (SGLT2)

### **Products Affected**

#### • Kerendia

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

### KINERET

#### **Products Affected**

#### • Kineret

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Rheumatoid Arthritis (RA) (initial): Diagnosis of moderately to severely active RA. One of the following: Trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: Enbrel (etanercept), one Formulary adalimumab product, Rinvoq (upadacitinib), or Xeljanz/Xeljanz XR (tofacitinib), OR for continuation of prior therapy if within the past 120 days. Neonatal-Onset Multisystem Inflammatory Disease (NOMID) (initial): Diagnosis of NOMID. Dx of NOMID confirmed by one of the following: 1) NLRP-3 (nucleotide-binding domain, leucine rich family (NLR), pyrin domain containing 3-gene) (also known as Cold-Induced Auto-inflammatory Syndrome-1 [CIAS1]) mutation or 2) Both of the following: a) two of the following clinical symptoms: urticaria-like rash, cold/stress triggered episodes, sensorineural hearing loss, musculoskeletal symptoms (e.g., arthralgia, arthritis, myalgia), chronic aseptic meningitis, or skeletal abnormalities (e.g., epiphyseal overgrowth, frontal bossing) AND b) elevated acute phase reactants (eg, erythrocyte sedimentation rate [ESR], C-reactive protein [CRP], serum amyloid A [SAA]). Deficiency of Interleukin-1 Receptor Antagonist (DIRA): Diagnosis of DIRA. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | RA (initial): Prescribed by or in consultation with a rheumatologist. NOMID (initial): Prescribed by or in consultation with an allergist/immunologist, pediatrician, or rheumatologist.  |
| Coverage<br>Duration               | RA, NOMID (initial, reauth): plan year. DIRA: plan year.  |
| Other Criteria                     | RA, NOMID (reauth): Patient demonstrates positive clinical response to therapy.   |

| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug. |
|-------------------------------------|--|
|-------------------------------------|--|

# KISQALI

#### **Products Affected**

• Kisqali

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Advanced or Metastatic Breast cancer: Diagnosis of advanced, recurrent, or metastatic breast cancer. Cancer is hormone receptor (HR)-positive and human epidermal growth factor receptor 2 (HER2)-negative. One of the following: A) Used in combination with an aromatase inhibitor (e.g., anastrozole, letrozole, exemestane) or B) Used in combination with Faslodex (fulvestrant). Early Breast Cancer: Diagnosis of early breast cancer. Both of the following: A) Disease is HR-positive, and B) Disease is HER2-negative. Both of the following: A) Disease is stage II or stage III, and B) Patient is at high risk of recurrence. Used in combination with an aromatase inhibitor (e.g., anastrozole, letrozole, exemestane). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# KISQALI - FEMARA PACK

### **Products Affected**

• Kisqali Femara 400 Dose

• Kisqali Femara 600 Dose

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Advanced or Metastatic Breast cancer: Diagnosis of advanced, recurrent, or metastatic breast cancer. Cancer is hormone receptor (HR)-positive and human epidermal growth factor receptor 2 (HER2)-negative. Early Breast Cancer: Diagnosis of early breast cancer. Both of the following: A) Disease is HR-positive, and B) Disease is HER2-negative. Both of the following: A) Disease is stage II or stage III, and B) Patient is at high risk of recurrence. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **KORLYM**

### **Products Affected**

### • Mifepristone TABS 300MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | (Initial): Diagnosis of endogenous Cushing's syndrome (i.e., hypercortisolism is not a result of chronic administration of high dose glucocorticoids). Diagnosis of either type 2 diabetes mellitus or diagnosis of glucose intolerance. Patient has either failed surgery or patient is not a candidate for surgery. Patient is not pregnant. (Reauthorization): Documentation of one of the following: patient has improved glucose tolerance while on therapy or patient has stable glucose tolerance while on therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 6 months   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **KOSELUGO**

#### **Products Affected**

#### • Koselugo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Neurofibromatosis Type 1 (NF1): Diagnosis of NF1. Patient has plexiform neurofibromas that are both of the following: inoperable and causing significant morbidity (e.g., disfigurement, motor dysfunction, pain, airway dysfunction, visual impairment, bladder/bowel dysfunction). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### KRAZATI

#### **Products Affected**

#### • Krazati

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Presence of KRAS G12C mutation. Disease is recurrent, advanced, or metastatic. Patient has received at least one prior systemic therapy (e.g., chemotherapy). Colorectal Cancer (CRC): Diagnosis of CRC. Presence of KRAS G12C mutation. Disease is recurrent, advanced, or metastatic. Patient has received at least one prior systemic therapy (e.g., chemotherapy). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# LAPATINIB

#### **Products Affected**

### • Lapatinib Ditosylate

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Breast Cancer: One of the following: A) Diagnosis of recurrent or stage IV hormone receptor positive (HR+), human epidermal growth factor receptor 2-positive (HER2+) breast cancer. Used in combination with an aromatase inhibitor [eg, Aromasin (exemestane), Femara (letrozole), Arimedex (anastrozole)]. OR B) Diagnosis of recurrent or metastatic HER2+ breast cancer. Used in combination with trastuzumab or Xeloda (capecitabine). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# LAZCLUZE

#### **Products Affected**

#### • Lazcluze

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of non-small cell lung cancer (NSCLC). Disease is one of the following: a) recurrent b) advanced c) metastatic. Disease is positive for one of the following: a) Epidermal growth factor receptor (EGFR) exon 19 deletion OR b) EGFR exon 21 L858R mutation. Used in combination with Rybrevant (amivantamab-vmjw). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# LENALIDOMIDE

#### **Products Affected**

#### • Lenalidomide

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

### **LENVIMA**

#### **Products Affected**

- Lenvima 10 Mg Daily Dose
- Lenvima 12mg Daily Dose
- Lenvima 14 Mg Daily Dose
- Lenvima 18 Mg Daily Dose

- Lenvima 20 Mg Daily Dose
- Lenvima 24 Mg Daily Dose
- Lenvima 4 Mg Daily Dose
- Lenvima 8 Mg Daily Dose

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Differentiated thyroid cancer (DTC): Diagnosis of DTC. Renal cell carcinoma: Diagnosis of Renal cell carcinoma. Hepatocellular Carcinoma (HCC): Diagnosis of Hepatocellular Carcinoma or liver cell carcinoma. Endometrial Carcinoma: Diagnosis of endometrial carcinoma. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# LEUPROLIDE ACETATE

#### **Products Affected**

• Leuprolide Acetate INJ 1MG/0.2ML

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Prostate Cancer: Treatment of advanced prostate cancer.                                 |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Prostate Cancer: plan year  |
| Other Criteria                      | Prostate Cancer: Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                            |

# LIDOCAINE PATCH

#### **Products Affected**

#### • Lidocaine PTCH 5%

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.                           |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Post-herpetic neuralgia: Diagnosis of post-herpetic neuralgia |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### LIVTENCITY

#### **Products Affected**

### • Livtencity

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of cytomegalovirus (CMV) infection/disease. Patient is a recipient of one of the following: a) Hematopoietic stem cell transplant OR b) Solid organ transplant. Trial and failure of a minimum 2 weeks duration, contraindication, or intolerance to one of the following therapies at an appropriately indicated dose: a) Intravenous (IV) ganciclovir, b) Oral valganciclovir, c) IV foscarnet, OR d) IV cidofovir. Patient weighs greater than or equal to 35 kg. |
| Age Restrictions                    | Patient is 12 years of age or older.   |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a provider who specializes in one of the following areas: transplant, oncology, or infectious disease.   |
| Coverage<br>Duration                | 8 weeks  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# LONSURF

#### **Products Affected**

#### • Lonsurf

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Metastatic colorectal cancer (mCRC): Diagnosis of mCRC. Trial and failure, contraindication, or intolerance with all of the following: fluoropyrimidine-based chemotherapy, oxaliplatin-based chemotherapy, irinotecan-based chemotherapy, and anti-VEGF biological therapy (e.g., bevacizumab). One of the following: a) tumor is RAS mutant-type or b) tumor is RAS wild-type and Trial and failure, contraindication or intolerance to one anti-EGFR therapy (eg, Vectibix [panitumumab], Erbitux [cetuximab]). Gastric/Gastroesophageal Junction Adenocarcinoma: Diagnosis of metastatic gastric cancer or diagnosis of metastatic gastroesophageal junction adenocarcinoma. Trial and failure, contraindication or intolerance to at least two of the following: fluropyrimidine-based chemotherapy (e.g. fluorouracil), Platinum-based chemotherapy (e.g., carboplatin, cisplatin, oxaliplatin), Taxane (e.g., docetaxel, paclitaxel) or irinotecan-based chemotherapy, HER2/neutargeted therapy (e.g., trastuzumab) (if HER2 overexpression). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# LORBRENA

### **Products Affected**

#### • Lorbrena

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. One of the following: A) Disease is advanced, metastatic, or recurrent and anaplastic lymphoma kinase (ALK)-positive OR B) Both of the following: 1) Disease is both of the following: i) advanced, metastatic, or recurrent and ii) ROS proto-oncogene 1 (ROS1)-positive AND 2) Disease has progressed on at least one of the following therapies: Augtyro (repotrectinib), Xalkori (crizotinib), Rozlytrek (entrectinib), or Zykadia (ceritinib). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# LUMAKRAS

#### **Products Affected**

#### • Lumakras

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: a) recurrent, b) advanced or c) metastatic. Tumor is KRAS G12C-mutated. Patient has received at least one prior systemic therapy (e.g., immune checkpoint inhibitor, platinum-based chemotherapy). Metastatic Colorectal Cancer (mCRC): Diagnosis of metastatic colorectal cancer (mCRC). Tumor is KRAS G12C-mutated. Patient has received prior therapy with fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy. Used in combination with Vectibix (panitumumab). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## LUMRYZ

### **Products Affected**

• Lumryz

• Lumryz Starter Pack

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **LUPRON DEPOT**

#### **Products Affected**

- Lupron Depot (1-month)
- Lupron Depot (3-month)

- Lupron Depot (4-month)
- Lupron Depot (6-month)

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Prostate Cancer (7.5 mg, 22.5 mg, 30 mg, 45 mg): Treatment of advanced prostate cancer. Endometriosis (3.75 mg, 11.25 mg) (initial): Diagnosis of endometriosis. One of the following: a) Patient has had surgical ablation to prevent recurrence, or b) trial and failure, contraindication, or intolerance to one NSAID (e.g., diclofenac, ibuprofen, meloxicam, naproxen) and one oral contraceptive (e.g., norethindrone-ethinyl estradiol, estradiol and norethindrone). Endometriosis (3.75 mg, 11.25 mg) (reauthorization): Symptoms recur after one course. Used in combination with one of the following: norethindrone 5 mg daily, other "add -back" sex hormones (e.g., estrogen, medroxyprogesterone), or other bone-sparing agents (e.g., bisphosphonates such as alendronate, risedronate). Uterine Leiomyomata (3.75 mg, 11.25 mg) (fibroids): Either for use prior to surgery to reduce size of fibroids to facilitate a surgical procedure (eg, myomectomy, hysterectomy) Or all of the following: treatment of anemia, anemia caused by uterine leiomyomata (fibroids), and use prior to surgery. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Prostate CA: plan yr. Endometrosis(all), Uterine leiomyomata (anemia): 6 mo. (fibroids): 4 mo.  |
| Other Criteria                     | Prostate Cancer (7.5 mg, 22.5 mg, 30 mg, 45 mg): Approve for continuation of prior therapy if within the past 120 days.   |

## **LUPRON DEPOT PED**

#### **Products Affected**

- Lupron Depot-ped (1-month) INJ 7.5MG
- Lupron Depot-ped (3-month) INJ 11.25MG
- Lupron Depot-ped (6-month)

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Central Precocious Puberty (CPP) (initial): diagnosis of central precocious puberty (idiopathic or neurogenic). Onset of secondary sexual characteristics in one of the following: females less than age 8 or males less than age 9. Confirmation of diagnosis defined by one of the following: pubertal basal level of luteinizing hormone (based on laboratory reference ranges), a pubertal response to a GnRH stimulation test, or bone age advanced one year beyond the chronological age. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | CPP (ini, reauth): plan yr.   |
| Other Criteria                      | CPP (Reauthorization): Documentation of bone age monitoring (eg, radiographic imaging).   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# LYNPARZA (TABLETS)

### **Products Affected**

### • Lynparza TABS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## LYTGOBI

### **Products Affected**

### • Lytgobi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of cholangiocarcinoma (intrahepatic or extrahepatic). Disease is one of the following: a) unresectable, b) locally advanced, c) Resected gross residual (R2), or d) metastatic. Positive for fibroblast growth factor receptor 2 (FGFR2) fusions or other rearrangements. Used as second line or subsequent treatment. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **MAVYRET**

### **Products Affected**

• Mavyret

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **MEKINIST**

### **Products Affected**

#### • Mekinist

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Melanoma: Diagnosis of unresectable or metastatic melanoma AND cancer is BRAF V600 mutant type. Adjuvant Treatment for Melanoma: Diagnosis of melanoma. Cancer is BRAF V600E or V600K mutant type. Involvement of lymph nodes following complete resection. Used as adjuvant therapy. Medication is used in combination with Tafinlar (dabrafenib). Non-small Cell Lung Cancer (NSCLC): All of the following: diagnosis of non-small cell lung cancer AND disease is one of the following: metastatic, advanced, or recurrent AND cancer is BRAF V600E mutant type AND medication is used in combination with Tafinlar (dabrafenib). Anaplastic Thyroid Cancer (ATC): Diagnosis of ATC. One of the following: 1) Disease is one of the following: metastatic, locally advanced, or unresectable OR 2) Prescribed as adjuvant therapy following resection. Cancer is BRAF V600E mutant type. Medication is used in combination with Tafinlar (dabrafenib). Solid tumors: Presence of solid tumor. Disease is unresectable or metastatic. Patient has progressed on or following prior systemic treatment (e.g., carboplatin, 5-fluorouracil, paclitaxel). Cancer is BRAF V600E mutant type. Medication is used in combination with Tafinlar (dabrafenib). Low-grade glioma: Diagnosis of low-grade glioma. Cancer is BRAF V600E mutant type. Medication is used in combination with Tafinlar (dabrafenib). |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Plan year   |
| Other Criteria                     | Approve for continuation of prior therapy if within the past 120 days.  |

## **MEKTOVI**

#### **Products Affected**

#### • Mektovi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma. Patient is positive for BRAF V600 mutation. Used in combination with Braftovi (encorafenib). Non-Small Cell Lung Cancer (NSCLC): Diagnosis of metastatic NSCLC. Patient is positive for BRAF V600 mutation. Used in combination with Braftovi (encorafenib). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### **MEMANTINE**

#### **Products Affected**

- Memantine Hcl Titration Pak
- Memantine Hydrochloride SOLN 2MG/ML
- Memantine Hydrochloride TABS

- Memantine Hydrochloride Er
- Memantine/donepezil Hydrochloride Er

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Age 41 or older, or diagnosis of moderate to severe dementia of the Alzheimer's type. |
| Age Restrictions                    | No Prior Authorization if patient is age 41 or older.                                 |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                          |

# MENQUADFI

### **Products Affected**

### • Menquadfi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## MENVEO

### **Products Affected**

#### • Menveo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **M**IGLUSTAT

### **Products Affected**

• Miglustat

• Yargesa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Gaucher disease: Diagnosis of mild to moderate Type 1 Gaucher disease. Patient is unable to receive enzyme replacement therapy due to one of the following conditions: allergy or hypersensitivity to enzyme replacement therapy, poor venous access, or unavailability of enzyme replacement therapy (e.g. Cerezyme, VPRIV). Niemann-Pick disease type C (NPC) (off-label) (initial): Diagnosis of NPC. Requested drug will be used in combination with Miplyffa (arimoclomol). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Niemann-Pick disease type C (NPC) (off-label) (initial): Prescribed by or in consultation with a specialist knowledgeable in the treatment of Niemann-Pick disease type C.   |
| Coverage<br>Duration                | Gaucher disease: Plan year. NPC (initial): 6 months, (reauth): Plan year.  |
| Other Criteria                      | Niemann-Pick disease type C (NPC) (off-label) (reauth): Patient demonstrates positive clinical response to therapy. Requested drug will be used in combination with Miplyffa (arimoclomol).  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## MODAFINIL

### **Products Affected**

#### • Modafinil TABS

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Obstructive sleep apnea (OSA) (Initial): Diagnosis (dx) of OSA defined by one of the following: 15 or more obstructive respiratory events per hour of sleep confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible), or both of the following: 5 or more obstructive respiratory events per hour of sleep confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible), and 1 of the following symptoms: unintentional sleep episodes during wakefulness, daytime sleepiness, unrefreshing sleep, fatigue, insomnia, waking up breath holding/gasping/choking, loud snoring, or breathing interruptions during sleep. Shift-work disorder (SWD) (Initial):Dx of SWD confirmed by one of the following: 1) symptoms of excessive sleepiness or insomnia for at least 3 months, which is associated with a work period (usually night work) that occurs during the normal sleep period, OR 2) A sleep study demonstrating loss of a normal sleep-wake pattern (ie, disturbed chronobiologic rhythmicity). Confirmation that no other medical conditions or medications are causing the symptoms of excessive sleepiness or insomnia. Narcolepsy (initial): Dx of narcolepsy as confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible). MS Fatigue (initial): Dx of multiple sclerosis (MS). Patient is experiencing fatigue. Idiopathic Hypersomnia (initial): Dx of idiopathic hypersomnia as confirmed by a sleep study (unless prescriber provides justification confirming that a sleep study is not feasible). Depression (initial): Treatment-resistant depression defined as diagnosis of major depressive disorder (MDD) or bipolar depression, AND trial and failure, contraindication, or intolerance to at least two antidepressants from different classes (eg, SSRIs, SNRIs, bupropion). Used as adjunctive therapy. |
| Age Restrictions                   | N/A   |

| Prescriber<br>Restrictions          | N/A   |
|-------------------------------------|---|
| Coverage<br>Duration                | OSA,SWD,MS Fatigue,Hypersomnia,Depression:Initial,Reauth:6 mo.Narcolepsy:Initial,Reauth:Plan Yr   |
| Other Criteria                      | OSA, Narcolepsy, Idiopathic Hypersomnia (Reauth): Patient demonstrates positive clinical response to therapy. SWD (Reauth): Patient demonstrates positive clinical response to therapy. MS Fatigue (reauth): Patient is experiencing relief of fatigue with therapy. Depression (reauth): Patient demonstrates positive clinical response to therapy. Used as adjunctive therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **MODEYSO**

### **Products Affected**

• Modeyso

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## MOUNJARO

### **Products Affected**

#### • Mounjaro

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Excluded if used for weight loss only.   |
| Required<br>Medical<br>Information  | Diabetes Mellitus (DM): Submission of medical records (e.g., chart notes) confirming diagnosis of type 2 DM. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## **M**RESVIA

### **Products Affected**

#### • Mresvia

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## NAYZILAM

### **Products Affected**

### • Nayzilam

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Epilepsy: Diagnosis of epilepsy. Frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) that are distinct from a patient's usual seizure pattern. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## **NERLYNX**

### **Products Affected**

### • Nerlynx

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Early Stage Breast cancer: Diagnosis (dx) of early stage breast cancer. Disease is human epidermal growth factor receptor 2 (HER2)-positive. Patient has received adjuvant trastuzumab based therapy (e.g., Herceptin, Kanjinti, etc.). Advanced or Metastatic Breast Cancer: 1) All of the following: a) Dx of advanced or metastatic breast cancer, b) Disease is human epidermal growth factor receptor 2 (HER2)-positive, c) Patient has received two or more prior anti-HER2 based regimens (e.g., trastuzumab + pertuzumab + docetaxel, ado-trastuzumab emtansine, etc.), and d) Used in combination with capecitabine, OR 2) Both of the following: a) Diagnosis of Stage IV (M1) breast cancer, and b) One of the following: i) Hormone receptor-positive, (HER2)-negative disease in patients who have already received a CDK4/6 inhibitor therapy, or ii) Triple negative disease. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **NEXLETOL**

### **Products Affected**

#### • Nexletol

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Heterozygous familial hypercholesterolemia (HeFH), primary hyperlipidemia, established cardiovascular disease (CVD), or high risk for a CVD event but without established CVD (Initial): One of the following diagnoses: A) Heterozygous familial hypercholesterolemia (HeFH), B) Primary hyperlipidemia, C) Established CVD (e.g., coronary artery disease, symptomatic peripheral arterial disease, cerebrovascular atherosclerotic disease), OR D) At high risk for a CVD event but without established CVD (e.g., diabetes mellitus [type 1 or type 2] in females over 65 years of age or males over 60 years of age). One of the following: (1) used as adjunct to statin therapy, (2) patient has intolerance to a statin, OR (3) patient has a contraindication to all statins. One of the following LDL-C values within the last 120 days: (1) LDL-C greater than or equal to 55 mg/dL with ASCVD or (2) LDL-C greater than or equal to 70 mg/dL without ASCVD. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Initial, Reauth: Plan year  |
| Other Criteria                     | HeFH or primary hyperlipidemia, Established cardiovascular disease (CVD) or high risk for a CVD event but without established CVD (Reauth): Patient demonstrates positive clinical response to therapy. Patient continues to receive other lipid-lowering treatment (e.g., statins, ezetimibe) or patient has an inability to take other lipid-lowering therapy (e.g., statins, ezetimibe).   |

| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug. |
|-------------------------------------|--|
|-------------------------------------|--|

## NEXLIZET

### **Products Affected**

#### • Nexlizet

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Heterozygous familial hypercholesterolemia (HeFH), primary hyperlipidemia, established cardiovascular disease (CVD), or high risk for a CVD event but without established CVD (Initial): One of the following diagnoses: A) Heterozygous familial hypercholesterolemia (HeFH), B) Primary hyperlipidemia, C) Established CVD (e.g., coronary artery disease, symptomatic peripheral arterial disease, cerebrovascular atherosclerotic disease), OR D) At high risk for a CVD event but without established CVD (e.g., diabetes mellitus [type 1 or type 2] in females over 65 years of age or males over 60 years of age). One of the following: (1) used as adjunct to statin therapy, (2) patient has intolerance to a statin, OR (3) patient has a contraindication to all statins. One of the following LDL-C values within the last 120 days: (1) LDL-C greater than or equal to 55 mg/dL with ASCVD or (2) LDL-C greater than or equal to 70 mg/dL without ASCVD. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Initial, Reauth: Plan year  |
| Other Criteria                     | HeFH or primary hyperlipidemia, Established cardiovascular disease (CVD) or high risk for a CVD event but without established CVD (Reauth): Patient demonstrates positive clinical response to therapy. Patient continues to receive other lipid-lowering treatment (e.g., statins, ezetimibe) or patient has an inability to take other lipid-lowering therapy (e.g., statins, ezetimibe).   |

| Prerequisite Criteria DOES NOT require use of a prerequisite Part D drug.  Therapy Required |  |
|---|--|
|---|--|

## **NINLARO**

### **Products Affected**

#### • Ninlaro

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Multiple myeloma: Diagnosis of multiple myeloma.                       |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# NUBEQA

### **Products Affected**

• Nubeqa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## NUCALA

### **Products Affected**

#### • Nucala

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## NUEDEXTA

### **Products Affected**

#### • Nuedexta

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

### NUPLAZID

### **Products Affected**

• Nuplazid CAPS

• Nuplazid TABS 10MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Parkinson's disease psychosis: Diagnosis of Parkinson's disease. Patient has at least one of the following: hallucinations or delusions. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## NURTEC

### **Products Affected**

#### • Nurtec

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute Treatment of Migraine (initial): Diagnosis of migraine with or without aura. Will be used for the acute treatment of migraine. Trial and failure or intolerance to one triptan (e.g., eletriptan, rizatriptan, sumatriptan) or a contraindication to all triptans. Medication will not be used in combination with another CGRP inhibitor for the acute treatment of migraines. Preventive Treatment of Episodic Migraine (EM) (initial): Diagnosis of EM. Patient has greater than or equal to 4 migraine days per month. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | All Indications (initial, reauth): Plan year   |
| Other Criteria                      | Acute Treatment of Migraine (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another CGRP inhibitor for the acute treatment of migraines. Preventive Treatment of EM (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **O**CTREOTIDE

#### **Products Affected**

 Octreotide Acetate INJ 1000MCG/ML, 100MCG/ML, 200MCG/ML, 500MCG/ML, 50MCG/ML

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Acromegaly (initial): Diagnosis of acromegaly confirmed by one of the following: serum GH level greater than 1 ng/mL after a 2-hour oral glucose tolerance test at the time of diagnosis, or elevated serum IGF-1 levels (above the age and gender adjusted normal range as provided by the physician's lab) at the time of diagnosis. One of the following: A) Inadequate response to surgery, radiotherapy, or dopamine agonist (e.g., bromocriptine, cabergoline) therapy, or B) Not a candidate for any of the following: surgery, radiotherapy, dopamine agonist (e.g., bromocriptine, cabergoline) therapy. HIV/AIDS-Related Diarrhea (initial): Diagnosis of HIV/AIDS-related diarrhea. Carcinoid tumors, symptomatic treatment of diarrhea or flushing (initial): diagnosis of metastatic carcinoid tumor requiring symptomatic treatment of severe diarrhea or flushing episodes. Vasoactive Intestinal Peptide Tumors, symptomatic treatment of diarrhea (initial): Diagnosis of vasoactive intestinal peptide tumor requiring treatment of profuse watery diarrhea. Cancer Chemotherapy- and/or Radiation- Induced Diarrhea (initial): Diagnosis of complicated diarrhea due to concurrent cancer chemotherapy and/or radiation or uncomplicated diarrhea due to concurrent cancer chemotherapy and/or radiation. Carcinoid tumor: diagnosis of carcinoid tumor. Reauthorization (all except carcinoid tumor): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | All Indications: Plan Year  |

| Other Criteria                      | Uncomplicated diarrhea due to concurrent cancer chemotherapy and/or radiation (initial): Trial and failure, contraindication, or intolerance (TF/C/I) to standard therapy (e.g., loperamide). HIV/AIDS-related Diarrhea (initial): TF/C/I to standard therapy (e.g., loperamide, diphenoxylate with atropine). Carcinoid tumor: Approve for continuation of prior therapy if within the past 120 days. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **O**DOMZO

### **Products Affected**

#### • Odomzo

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Advanced basal cell carcinoma: Diagnosis of locally advanced basal cell carcinoma. One of the following: cancer that has recurred following surgery or radiation therapy, or patient is not a candidate for surgery or radiation therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **O**FEV

### **Products Affected**

• Ofev

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **O**GSIVEO

### **Products Affected**

### • Ogsiveo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of desmoid tumor. Patient requires systemic treatment. Disease is progressive. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.                   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                             |

# **O**JEMDA

### **Products Affected**

### • Ojemda

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of pediatric low-grade glioma. Disease is relapsed or refractory. Disease has a BRAF fusion or rearrangement, or BRAF V600 mutation. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **O**JJAARA

### **Products Affected**

• Ojjaara

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | One of the following: A) Diagnosis of symptomatic lower-risk myelofibrosis, OR B) All of the following: 1) Diagnosis of higher risk myelofibrosis, 2) Presence of symptomatic splenomegaly and/or constitutional symptoms, and 3) One of the following: a) Used as continued therapy near the start of conditioning therapy in a transplant candidate or b) Patient is not a transplant candidate or transplant not currently feasible, OR C) Diagnosis of myelofibrosis-associated anemia, OR D) Both of the following: 1) Diagnosis of accelerated/blast phase myeloproliferative neoplasm and 2) One of the following: a) Used for the improvement of splenomegaly or other disease-related symptoms or b) Continued treatment as a single agent near to the start of conditioning therapy in transplant candidates for the improvement of splenomegaly and other disease-related symptoms. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **O**NUREG

### **Products Affected**

#### • Onureg

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Acute Myeloid Leukemia (AML): Diagnosis of acute myeloid leukemia (AML). Patient has received previous treatment with an intensive induction chemotherapy regimen (e.g., cytarabine + daunorubicin, cytarabine + idarubicin). Patient has achieved one of the following: a) first complete remission (CR) or b) complete remission with incomplete blood count recovery (CRi). Patient is not a candidate for intensive curative therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **O**PIPZA

### **Products Affected**

• Opipza

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **ORENITRAM**

#### **Products Affected**

- Orenitram
- Orenitram Titration Kit Month 1
- Orenitram Titration Kit Month 2
- Orenitram Titration Kit Month 3

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH. PAH is symptomatic. One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PAH: Prescribed by or in consultation with a pulmonologist or cardiologist.   |
| Coverage<br>Duration                | PAH: plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **O**RGOVYX

### **Products Affected**

### • Orgovyx

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prostate Cancer: Diagnosis of prostate cancer.                         |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# **O**RKAMBI

### **Products Affected**

#### • Orkambi TABS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Cystic fibrosis (initial): Diagnosis of cystic fibrosis (CF). Submission of laboratory records confirming the patient is homozygous for the F508del mutation in the CFTR gene. |
| Age Restrictions                    | Patient is greater than or equal to 6 years of age   |
| Prescriber<br>Restrictions          | CF (initial, reauthorization): Prescribed by or in consultation with a specialist affiliated with a CF care center or pulmonologist  |
| Coverage<br>Duration                | CF (initial, reauth): plan year  |
| Other Criteria                      | CF (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# ORKAMBI GRANULES

### **Products Affected**

#### • Orkambi PACK

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Cystic fibrosis (initial): Diagnosis of cystic fibrosis (CF). Submission of laboratory records confirming the patient is homozygous for the F508del mutation in the CFTR gene. One of the following: A) Patient is 1 through 5 years of age, OR B) Both of the following: Patient is 6 years of age or greater AND Patient is unable to swallow oral tablets. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | CF (initial, reauthorization): Prescribed by or in consultation with a specialist affiliated with a CF care center or pulmonologist   |
| Coverage<br>Duration                | CF (initial, reauth): plan year   |
| Other Criteria                      | CF (Reauthorization): Patient demonstrates positive clinical response to therapy. One of the following: A) Patient is 1 through 5 years of age, OR B) Both of the following: Patient is 6 years of age or greater AND Patient is unable to swallow oral tablets.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **O**RSERDU

#### **Products Affected**

#### • Orserdu

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of breast cancer. Disease is advanced or metastatic. One of the following: a) Patient is male, b) Patient is a postmenopausal woman, or c) Patient is a premenopausal woman treated with ovarian ablation/suppression. Disease is estrogen receptor (ER)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Presence of estrogen receptor (ESR1) mutation(s). Disease has progressed following at least one line of endocrine therapy [e.g., Faslodex (fulvestrant)]. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **OSPHENA**

### **Products Affected**

### • Osphena

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Dyspareunia: Diagnosis of moderate to severe dyspareunia due to vulvar and vaginal atrophy associated with menopause. Vaginal dryness: Diagnosis of moderate to severe vaginal dryness due to vulvar and vaginal atrophy associated with menopause. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **O**TEZLA

### **Products Affected**

#### • Otezla

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Psoriatic arthritis (PsA, initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Plaque psoriasis (initial): Diagnosis of plaque psoriasis. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Oral ulcers associated with Behcet's Disease (initial): Diagnosis of Behcet's Disease. Patient has active oral ulcers. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PsA (init): Prescribed by or in consultation with one of the following specialists: dermatologist or rheumatologist. Plaque psoriasis (init): Prescribed by or in consultation with a dermatologist.  |
| Coverage<br>Duration                | All uses (initial, reauth): plan year.  |
| Other Criteria                      | All indications (Reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **OZEMPIC**

### **Products Affected**

• Ozempic INJ 2MG/3ML, 4MG/3ML, 8MG/3ML

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Excluded if used for weight loss only.   |
| Required<br>Medical<br>Information  | Diabetes Mellitus (DM): Submission of medical records (e.g., chart notes) confirming diagnosis of type 2 DM. Nonalcoholic steatohepatitis (NASH)/Metabolic dysfunction-associated steatohepatitis (MASH) (Initial): Diagnosis of MASH, formerly known as nonalcoholic steatohepatitis (NASH). Patient does not have cirrhosis (e.g., decompensated cirrhosis). Submission of medical records (e.g., chart notes) confirming disease is fibrosis stage F2 or F3 as confirmed by one of the following: 1) Both of the following: A) Serum biomarker [e.g., enhanced liver fibrosis (ELF) test, fibrosis-4 index (FIB-4)], and B) Imaging biormarker [e.g., FibroScan, magnetic resonance imaging-proton density fat fraction (MRI-PDFF)], or 2) One of the following: A) FibroScan aspartate aminotransferase (FAST), B) MRI-aspartate aminotransferase (MAST), C) Magnetic Resonance Elastography combined with fibrosis-4-index (MEFIB), or D) liver biopsy within the past 12 months. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | MASH (Initial): Prescribed by or in consultation with a gastroenterologist or hepatologist.  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | MASH (Reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **PANRETIN**

#### **Products Affected**

#### • Panretin

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Kaposi's sarcoma lesions: Diagnosis of cutaneous lesions in patients with AIDS-related Kaposi's sarcoma (KS). Not used when systemic anti-KS therapy is required (e.g., more than 10 new KS lesions in the prior month, symptomatic lymphedema, symptomatic pulmonary KS, or symptomatic visceral involvement). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# PEGFILGRASTIM PREFERRED

### **Products Affected**

• Neulasta

• Udenyca

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Neutropenia Associated with Dose Dense Chemotherapy (NDDC): Patient is receiving National Cancer Institute's Breast Intergroup, INT C9741 dose dense chemotherapy protocol for primary breast cancer, or a dose-dense chemotherapy regimen for which the incidence of febrile neutropenia is unknown. Chemotherapy-Induced Febrile Neutropenia (CFN): Patient is receiving a chemotherapy regimen associated with greater than 20% incidence of febrile neutropenia, or patient is receiving chemotherapy regimen associated with 10-20% incidence of febrile neutropenia and has 1 or more risk factors associated with chemotherapy-induced infection, febrile neutropenia or neutropenia. Secondary prophylaxis of FN: For patients who are receiving myelosuppressive anticancer drugs associated with neutropenia. Patient has a history of FN or dose-limiting event during a previous course of chemotherapy (secondary prophylaxis). Acute radiation syndrome (ARS): Patient was/will be acutely exposed to myelosuppressive doses of radiation (hematopoietic subsyndrome of ARS). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a hematologist/oncologist.  |
| Coverage<br>Duration                | ARS: 1 mo. CFN, NDDC, FN (prophylaxis): 3 mo or duration of tx.   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# PEGINTERFERON ALFA - 2A

### **Products Affected**

#### • Pegasys

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Chronic Hepatitis B: Diagnosis of chronic hepatitis B infection. Chronic Hepatitis C: Diagnosis of chronic hepatitis C infection. Patient has compensated liver disease. One of the following: a) Used in combination with one other hepatitis C virus (HCV) antiviral drug (e.g., Mavyret [glecaprevir-pibrentasvir], ribavirin) OR b) Both of the following: Used as monotherapy AND contraindication or intolerance to all other HCV antiviral drugs (e.g., Mavyret [glecaprevir-pibrentasvir], ribavirin). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | HepB, HepC: 48 wks.  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **PEMAZYRE**

### **Products Affected**

#### • Pemazyre

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## **PENBRAYA**

### **Products Affected**

#### • Penbraya

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Vaccine is being used for prevention of Neisseria meningitidis serogroups A, B, C, W, and Y. Patient has not received or will not receive more than two Penbraya vaccines in the previous 12 months. Age lesser than or equal to 25 years. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 1 year (2 vaccinations per year)   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **PENMENVY**

### **Products Affected**

#### • Penmenvy

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **PIQRAY**

### **Products Affected**

• Piqray 200mg Daily Dose

- Piqray 250mg Daily Dose Piqray 300mg Daily Dose

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Breast Cancer (BC): Diagnosis of advanced or metastatic BC. Disease is hormone receptor (HR)-positive, and human epidermal growth factor receptor 2 (HER2)-negative. Presence of one or more PIK3CA mutations. Used in combination with fulvestrant. Disease has progressed on or after an endocrine-based regimen. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

### **PIRFENIDONE**

#### **Products Affected**

• Pirfenidone CAPS

• Pirfenidone TABS 267MG, 801MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Idiopathic pulmonary fibrosis (IPF) (initial): Diagnosis of IPF, defined as exclusion of other known causes of interstitial lung disease and either the presence of usual interstitial pneumonia (UIP) pattern on high-resolution computed tomography (HRCT) revealing IPF or probable IPF in patients not subjected to lung biopsy, or HRCT and surgical lung biopsy pattern revealing IPF or probable IPF in patients subjected to a lung biopsy. IPF (reauth): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | (initial): Prescribed by or in consultation with a pulmonologist.   |
| Coverage<br>Duration                | (initial, reauth): plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **POMALYST**

### **Products Affected**

### • Pomalyst

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Multiple myeloma: Diagnosis of multiple myeloma. One of the following:  1) Trial and failure, contraindication or intolerance to one of the following: a) immunomodulatory agent [eg, Revlimid (lenalidomide)] or b) proteasome inhibitor [eg, Velcade (bortezomib)], OR 2) Induction therapy for the management of POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal protein, skin changes) syndrome. Kaposi sarcoma (KS): One of the following: 1) Both of the following: a) Diagnosis of AIDS-related KS and b) Patient is currently being treated with antiretroviral therapy (ART), OR 2) Both of the following: a) Diagnosis of KS and b) Patient is HIV-negative. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# POSACONAZOLE ORAL

#### **Products Affected**

#### • Posaconazole Dr

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prophylaxis of Invasive Fungal Infections (IFI): Used as prophylaxis of invasive fungal infections caused by Aspergillus or Candida for one of the following conditions: 1) Patient is at high risk of infections due to severe immunosuppression from hematopoietic stem cell transplant (HSCT) with graft-versus-host disease (GVHD) or hematologic malignancies with prolonged neutropenia from chemotherapy [eg, acute myeloid leukemia (AML), myelodysplastic syndrome (MDS)], OR 2) patient has a prior fungal infection requiring secondary prophylaxis. Treatment of IFI: Used as treatment of invasive fungal infections caused by Aspergillus. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Prophylaxis of IFI: plan year. Treatment of IFI: 3 months.   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# PREVYMIS ORAL

#### **Products Affected**

• Prevymis PACK

### • Prevymis TABS

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Cytomegalovirus (CMV) Prophylaxis in Hematopoietic Stem Cell Transplant (HSCT): Used for prophylaxis of CMV infection and disease AND patient is a CMV-seropositive recipient [R+] of an allogeneic HSCT. CMV Prophylaxis in Kidney Transplant: Used for prophylaxis CMV infection and disease. Patient is a CMV-seronegative recipient [R-]. Patient is receiving a kidney transplant from a CMV-seropositive donor [D+]. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | CMV Prophylaxis in HSCT: Prescribed by or in consultation with an oncologist, hematologist, physician experienced in the management of transplant patients, or infectious disease specialist. CMV Prophylaxis in Kidney Transplant: Prescribed by or in consultation with a nephrologist, physician experienced in the management of transplant patients, or infectious disease specialist.                                |
| Coverage<br>Duration                | CMV Prophylaxis in HSCT: 7 months. CMV Prophylaxis in Kidney Transplant: 7 months.   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **PROMACTA**

### **Products Affected**

### • Eltrombopag Olamine

| PA<br>Criteria                     | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label<br>Uses                  | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Idiopathic thrombocytopenic purpura (ITP) (initial): Diagnosis of one of the following: persistent ITP, chronic ITP or relapsed/refractory ITP. Baseline platelet count is less than 30,000/mcL. Patient's degree of thrombocytopenia and clinical condition increase the risk of bleeding. ITP (reauthorization): Patient demonstrates positive clinical response to therapy as evidenced by an increase in platelet count to a level sufficient to avoid clinically important bleeding. Chronic Hepatitis C-associated thrombocytopenia (initial): Diagnosis of chronic hepatitis C-associated thrombocytopenia. One of the following: planning to initiate and maintain interferon-based treatment or currently receiving interferon-based treatment. First-line for severe aplastic anemia (SAA): Diagnosis of SAA. Used for first-line treatment (i.e., patient has not received prior immunosuppressive therapy with any equine antithymocyte globulin plus cyclosporine, alemtuzumab, or high dose cyclophosphamide). Used in combination with standard immunosuppressive therapy (e.g., Atgam [antithymocyte globulin equine] and cyclosporine). Patient meets at least two of the following: 1) absolute neutrophil count less than 500/mcL, 2) platelet count less than 20,000/mcL, 3) absolute reticulocyte count less than 60,000/mcL. Refractory SAA (initial): Diagnosis of refractory SAA. Patient has a platelet count less than 30,000/mcL. SAA (reauthorization): Patient demonstrates positive clinical response to therapy. |
| Age<br>Restrictions                | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | 1stline SAA:6mo.HepC<br>(init):3mo.RefractSAA(init):16wk.ITP,HepC(reauth),RefractSAA(reauth):plan<br>yr   |

#### Other Criteria

ITP (initial): Trial and failure, contraindication, or intolerance to at least one of the following: corticosteroids (e.g., prednisone, methylprednisolone), immunoglobulins [e.g., Gammagard, immune globulin (human)], or splenectomy. Chronic Hepatitis C-associated thrombocytopenia (Reauthorization): One of the following criteria: For patients that started treatment with eltrombopag prior to initiation of treatment with interferon, eltrombopag will be approved when both of the following are met: currently on antiviral interferon treatment for treatment of chronic hepatitis C and documentation that patient reached threshold platelet count that allows initiation of antiviral interferon therapy with eltrombopag treatment by week 9. OR for patients that started treatment with eltrombopag while on concomitant treatment with interferon, eltrombopag will be approved based on the following criterion: currently on antiviral interferon therapy for treatment of chronic hepatitis C. Refractory SAA: Trial and failure, contraindication, or intolerance to at least one course of immunosuppressive therapy (eg, Atgam (antithymocyte globulin equine), Thymoglobulin (antithymocyte globulin rabbit), cyclosporine).

#### Prerequisite Therapy Required

Criteria DOES require use of a prerequisite Part D drug.

# **PURIXAN**

### **Products Affected**

### • Mercaptopurine SUSP

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | One of the following: History of contraindication or intolerance to generic mercaptopurine tablets OR patient is unable to swallow tablets. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **PYRUKYND**

### **Products Affected**

• Pyrukynd

• Pyrukynd Taper Pack

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Initial: Diagnosis of hemolytic anemia confirmed by the presence of chronic hemolysis (e.g., increased indirect bilirubin, elevated lactated dehydrogenase [LDH], decreased haptoglobin, increased reticulocyte count). Diagnosis of pyruvate kinase deficiency confirmed by molecular testing of ALL the following mutations on the PKLR gene: a) Presence of at least 2 variant alleles in the pyruvate kinase liver and red blood cell (PKLR) gene, of which at least 1 was a missense variant AND b) Patients is not homozygous for the c.1436G to A (p.R479H) variant AND c) Patient does not have 2 non-missense variants (without the presence of another missense variant) in the PKLR gene. Hemoglobin is less than or equal to 10g/dL. Patient has symptomatic anemia or is transfusion dependent. Exclusion of other causes of hemolytic anemias (e.g., infections, toxins, drugs). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Initial, Reauth: Prescribed by or in consultation with a hematologist.   |
| Coverage<br>Duration                | Initial: 6 months. Reauth: Plan Year.  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **Q**INLOCK

### **Products Affected**

#### • Qinlock

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Gastrointestinal Stromal Tumor (GIST): Diagnosis of gastrointestinal stromal tumor (GIST). One of the following: a) gross residual disease (R2 resection), b) tumor rupture, c) unresectable primary disease, or d) recurrent/metastatic. One of the following: a) Trial and failure, contraindication, or intolerance to all of the following: imatinib (Gleevec), sunitinib (Sutent), and regorafenib (Stivarga), b) All of the following: performance status 0-2, history of progression on imatinib (Gleevec), and history of intolerance to sunitinib (Sutent), or c) All of the following: PDGFRA exon 18 mutations that are insensitive to imatinib (Gleevec) (including PDGFRA D842V), history of progression on avapritinib (Ayvakit), and history of progression on dasatinib (Sprycel). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# QUININE

### **Products Affected**

• Quinine Sulfate CAPS 324MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis (dx) of uncomplicated malaria and one of the following: treatment in areas of chloroquine-sensitive malaria or treatment in areas of chloroquine-resistant malaria. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | 7 days  |
| Other Criteria                      | Chloroquine-sensitive malaria: Failure, contraindication or intolerance to chloroquine or hydroxychloroquine.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **Q**ULIPTA

### **Products Affected**

### • Qulipta

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Episodic Migraine (EM) (initial): Diagnosis of EM. Patient has greater than or equal to 4 migraine days per month. Chronic Migraines (CM) (initial): Diagnosis of CM. Patient has greater than or equal to 8 migraine days per month. All Indications (initial): Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | EM, CM (initial, reauth): Plan year  |
| Other Criteria                      | EM, CM (reauth): Patient demonstrates positive clinical response to therapy. Medication will not be used in combination with another CGRP inhibitor for the preventive treatment of migraines.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# REPATHA

### **Products Affected**

• Repatha

• Repatha Sureclick

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | HeFH/ASCVD/Primary HLD(init): One of the following diagnoses: A) Heterozygous familial hypercholesterolemia (HeFH), B) Atherosclerotic cardiovascular disease (ASCVD), OR C) Primary Hyperlipidemia (HLD). ONE of the following: A) One of the following LDL-C values while on maximally tolerated lipid lowering tx within the last 120 days: (1) Patient requires greater than or equal to 25% LDL-C reduction to achieve goal or (2) LDL-C greater than or equal to 55 mg/dL with ASCVD, or (3) LDL-C greater than or equal to 70 mg/dL without ASCVD. OR B) Both of the following: (1) Patient has been receiving PCSK9 therapy as adjunct to maximally tolerated lipid lowering therapy and (2) LDL-C values drawn within the past 12 months while on maximally tolerated lipid lowering therapy are within normal limits. AND One of the following: i) Pt has been receiving at least 12 weeks of one maximally-tolerated statin tx, ii) Pt is statin intolerant as evidenced by an inability to tolerate at least two statins, with at least one started at the lowest starting daily dose, due to intolerable symptoms or clinically significant biomarker changes of liver function or muscle function (e.g., creatine kinase), iii) patient has a contraindication to all statins. HoFH (initial): dx of homozygous familial hypercholesterolemia (HoFH) as confirmed by one of the following: (1) Gen confirmation of 2 mutations in LDL receptor, ApoB, PCSK9, or LDLRAP1 or ARH, or (2) either untreated LDL greater than 500 or treated LDL greater than 300, AND either xanthoma before 10 yo or evidence of HeFH in both parents. Pt is receiving other lipid-lowering tx (eg statin, ezetimibe) OR patient has inability to take other lipid lowering tx (eg., statin, ezetimibe). |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |

| Coverage<br>Duration                | Initial, Reauth: plan year   |
|-------------------------------------|--|
| Other Criteria                      | HeFH/ASCVD/Primary HLD/HoFH (reauth): Pt continues to receive other lipid-lowering therapy (eg statin, ezetimibe) at the maximally tolerated dose or pt has an inability to take other lipid-lowering therapy (e.g., statin, ezetimibe). Patient demonstrates positive clinical response to therapy. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **RETEVMO**

### **Products Affected**

#### • Retevmo TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-Small Cell Lung cancer: Diagnosis of non-small cell lung cancer (NSCLC). Disease is one of the following: a) recurrent, b) advanced, or c) metastatic. Presence of RET gene fusion-positive or RET rearrangement positive tumor(s). Medullary Thyroid Cancer (MTC): Diagnosis of medullary thyroid cancer (MTC). Disease is advanced or metastatic. Disease has presence of RET gene mutation. Disease requires treatment with systemic therapy. Thyroid Cancer: Diagnosis of thyroid cancer. Disease is advanced or metastatic. Disease is RET gene fusion-positive. Disease requires treatment with systemic therapy. Patient is radioactive iodine-refractory or radioactive iodine therapy is not appropriate. Solid Tumors: Presence of RET gene fusion-positive solid tumor. Disease is recurrent, advanced, or metastatic. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Non-Small Cell Lung Cancer, MTC, Thyroid Cancer, Solid Tumors: Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# REVCOVI

#### **Products Affected**

#### • Revcovi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of adenosine deaminase deficiency (ADA) with severe combined immunodeficiency (SCID). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                                    |

# **REVUFORJ**

### **Products Affected**

• Revuforj

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of acute leukemia. Disease is relapsed or refractory. Positive for lysine methyltransferase 2A gene (KMT2A) translocation. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# REZDIFFRA (I)

### **Products Affected**

#### • Rezdiffra

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# REZLIDHIA

### **Products Affected**

#### • Rezlidhia

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of acute myeloid leukemia (AML). Disease is relapsed or refractory. Positive for a susceptible isocitrate dehydrogenase-1 (IDH1) mutation (e.g., R132C, R132H, R132G, R132S, R132L). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# RINVOQ

### **Products Affected**

• Rinvoq

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# RINVOQ LQ

### **Products Affected**

### • Rinvoq Lq

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Polyarticular juvenile idiopathic arthritis (PJIA) (init): Diagnosis of active PJIA. Minimum duration of a 6-week trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies at maximally tolerated doses: methotrexate, leflunomide. Psoriatic arthritis (PsA) (init): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. PJIA, PsA (init): Patient has had an inadequate response or intolerance to one or more TNF inhibitors (eg, adalimumab, etanercept). PJIA, PsA (init, reauth): Not used in combination with other JAK inhibitors, biologic DMARDs, or potent immunosuppressants (eg, azathioprine, cyclosporine). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PJIA (init): Prescribed by or in consultation with a rheumatologist. PsA (init): Prescribed by or in consultation with a dermatologist or rheumatologist.   |
| Coverage<br>Duration                | PJIA, PsA (init, reauth): Plan year.  |
| Other Criteria                      | PJIA, PsA (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **ROFLUMILAST**

#### **Products Affected**

#### • Roflumilast

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Chronic Obstructive Pulmonary Disease (COPD) (initial): Diagnosis of COPD. History of COPD exacerbations which required the use of systemic corticosteroids, antibiotics, or hospital admission. Trial and failure, intolerance, or contraindication to two prior therapies for COPD (e.g., Combivent, Spiriva). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Initial, reauth: plan year   |
| Other Criteria                      | COPD (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **ROMVIMZA**

#### **Products Affected**

#### • Romvimza

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of tenosynovial giant cell tumor (TGCT). Surgical resection will potentially cause worsening functional limitation or severe morbidity. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# ROZLYTREK

#### **Products Affected**

### • Rozlytrek

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of metastatic non-small cell lung cancer (NSCLC). Patient has ROS1 rearrangement positive tumor(s). Solid Tumors: Patient has solid tumors with a neurotrophic tyrosine receptor kinase (NTRK) gene fusion (e.g., ETV6-NTRK3, TPM3-NTRK1, TPR-NTRK1, etc.). Disease is without a known acquired resistance mutation (e.g., TRKA G595R, TRKA G667C or TRKC G623R substitutions). Disease is one of the following: metastatic or unresectable (including cases where surgical resection is likely to result in severe morbidity). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# RUBRACA

#### **Products Affected**

#### • Rubraca

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Ovarian cancer: Diagnosis of epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer. Used for maintenance treatment in patients who are in a complete or partial response to platinum-based chemotherapy (e.g., cisplatin, carboplatin). Presence of deleterious BRCA-mutation. Prostate cancer: Diagnosis of metastatic castration-resistant prostate cancer. Presence of deleterious BRCA mutation. History of failure, contraindication, or intolerance to both of the following: 1) Androgen receptor-directed therapy [e.g., Erleada (apalutamide), Xtandi (enzalutamide), Zytiga (abiraterone)], AND 2) A taxane-based chemotherapy [e.g., docetaxel, Jevtana (cabazitaxel)]. One of the following: 1) Used in combination with a gonadotropin-releasing hormone (GnRH) analog [e.g., Lupron (leuprolide), Zoladex (goserelin)], OR 2) Patient received bilateral orchiectomy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **RYBELSUS**

### **Products Affected**

• Rybelsus TABS 14MG, 3MG, 7MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Excluded if used for weight loss only.   |
| Required<br>Medical<br>Information  | Diabetes Mellitus (DM): Submission of medical records (e.g., chart notes) confirming diagnosis of type 2 DM. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# RYDAPT

### **Products Affected**

• Rydapt

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute Myeloid Leukemia (AML): Diagnosis of acute myeloid leukemia (AML), AML is FMS-like tyrosine kinase 3 (FLT3) mutation-positive, Rydapt will be used in combination with standard induction and consolidation therapy. Aggressive Systemic Mastocytosis (ASM), Systemic Mastocytosis with Associated Hematological Neoplasm (SM-AHN), Mast Cell Leukemia (MCL): Diagnosis of one of the following: aggressive systemic mastocytosis (ASM), systemic mastocytosis with associated hematological neoplasm (SM-AHN), or mast cell leukemia (MCL). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

### **SCEMBLIX**

#### **Products Affected**

#### • Scemblix

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of chronic myelogenous/myeloid leukemia (CML). Disease is Philadelphia chromosome-positive (Ph+) or BCR::ABL1-positive. Disease is in chronic phase or disease is in accelerated phase. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **SHINGRIX**

### **Products Affected**

### • Shingrix

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Vaccine is being used for prevention of herpes zoster (shingles). One of the following: A) Age greater than or equal to 50 years OR B) Both of the following: 1) Age 18 to 49 years and 2) Patient is or will be at increased risk of herpes zoster due to immunodeficiency or immunosuppression caused by known disease or therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 6 months (2 injections per lifetime)   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **SIGNIFOR**

### **Products Affected**

### • Signifor

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Cushing's disease: Diagnosis of endogenous Cushing's disease (i.e, hypercortisolism is not a result of chronic administration of high dose glucocorticoids). Either pituitary surgery has not been curative for the patient OR patient is not a candidate for pituitary surgery. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# SILDENAFIL

#### **Products Affected**

#### • Sildenafil Citrate TABS 20MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH. PAH is symptomatic. One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PAH: Prescribed by or in consultation with a pulmonologist or cardiologist.   |
| Coverage<br>Duration                | PAH: plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **SIRTURO**

### **Products Affected**

#### • Sirturo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **SKYCLARYS**

### **Products Affected**

• Skyclarys

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of Friedreich's ataxia confirmed via genetic testing demonstrating mutation in the FXN gene. Patient has a B-type natriuretic peptide value less than or equal to 200 pg/mL. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with one of the following: Neurologist, Neurogeneticist, or Physiatrist (Physical Medicine and Rehabilitation Specialist).                            |
| Coverage<br>Duration                | Initial, Reauth: Plan Year.   |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# SKYRIZI (I)

### **Products Affected**

• Skyrizi INJ 150MG/ML, 180MG/1.2ML, 360MG/2.4ML

• Skyrizi Pen

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Plaque Psoriasis (initial): Diagnosis of chronic moderate to severe plaque psoriasis. One of the following: at least 3% body surface area involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Psoriatic arthritis (PsA) (initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Crohn's disease (CD) (Initial): Diagnosis of moderately to severely active CD. Will be used as a maintenance dose following the intravenous induction doses. Ulcerative Colitis (UC) (Initial): Diagnosis of moderately to severely active UC. Will be used as a maintenance dose following the intravenous induction doses. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | Plaque Psoriasis (initial): Prescribed by or in consultation with a dermatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. CD, UC (Initial): Prescribed by or in consultation with a gastroenterologist.   |
| Coverage<br>Duration               | All uses (initial, reauth): plan year.   |
| Other Criteria                     | All indications (Reauth): Patient demonstrates positive clinical response to therapy.  |

### **SOMAVERT**

#### **Products Affected**

#### • Somavert

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Acromegaly (Initial): Diagnosis of acromegaly by one of the following: serum growth hormone (GH) level greater than 1 ng/mL after a 2 hour oral glucose tolerance test (OGTT) at time of diagnosis, or elevated serum IGF-1 levels (above the age and gender adjusted normal range as provided by the physician's lab) at time of diagnosis. Inadequate response to one of the following: surgery, radiotherapy, or dopamine agonist (e.g., bromocriptine, cabergoline) therapy or not a candidate for surgery, radiotherapy or dopamine agonist (eg, bromocriptine, cabergoline) therapy. One of the following: 1) Inadequate response, contraindication, or intolerance to a somatostatin analog (e.g., octreotide, lanreotide) or 2) Clinical rationale provided for preferred treatment with pegvisomant (e.g., comorbid diabetes mellitus is present with acromegaly, IGF-1 levels greater than 900 ng/mL). Acromegaly (Reauth): Patient demonstrates positive clinical response to therapy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Acromegaly (Initial, Reauth): plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

### **SORAFENIB**

### **Products Affected**

• Sorafenib Tosylate TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Renal cell carcinoma (RCC): Diagnosis of RCC. Hepatocellular carcinoma (HCC): Diagnosis of HCC. One of the following: patient has metastatic disease, or patient has extensive liver tumor burden, or patient is inoperable by performance status or comorbidity (local disease or local disease with minimal extrahepatic disease only), or both of the following: a) patient is not a transplant candidate, and b) disease is unresectable. Differentiated thyroid carcinoma (DTC): Diagnosis of one of the following: a) follicular carcinoma, b) oncocytic carcinoma, or c) papillary carcinoma. One of the following: metastatic disease, unresectable recurrent disease, or persistent locoregional disease. One of the following: patient has symptomatic disease or patient has progressive disease. Disease is refractory to radioactive iodine (RAI) treatment. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **SOTYKTU**

### **Products Affected**

#### • Sotyktu

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Plaque Psoriasis (initial): Diagnosis of moderate to severe plaque psoriasis. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Not used in combination with other potent immunosuppressants (eg, azathioprine, cyclosporine, biologic disease-modifying antirheumatic drugs [DMARDs]). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Plaque Psoriasis (initial): Prescribed by or in consultation with a dermatologist  |
| Coverage<br>Duration                | Plaque Psoriasis (initial, reauth): plan year.   |
| Other Criteria                      | Plaque psoriasis (reauth): Patient demonstrates positive clinical response to therapy. Not used in combination with other potent immunosuppressants (eg, azathioprine, cyclosporine, biologic DMARDs).   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **SPRYCEL**

#### **Products Affected**

#### • Dasatinib

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Chronic Myeloid Leukemia (CML): Diagnosis of chronic myeloid leukemia (CML). Acute Lymphoblastic Leukemia (ALL): Diagnosis of acute lymphoblastic leukemia (ALL). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### **STELARA**

#### **Products Affected**

- Stelara INJ 45MG/0.5ML, 90MG/ML
- Ustekinumab INJ 45MG/0.5ML, 90MG/ML

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **STEQEYMA**

### **Products Affected**

• Steqeyma INJ 45MG/0.5ML, 90MG/ML

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Plaque psoriasis (Initial - 45mg/0.5mL): Diagnosis of moderate to severe plaque psoriasis. Plaque psoriasis (Initial - 90mg/1mL): Diagnosis of moderate to severe plaque psoriasis. Patient's weight is greater than 100 kg (220 lbs). Plaque psoriasis (Initial regardless of dose): One of the following: at least 3% body surface area involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Psoriatic arthritis (PsA) (Initial - 45mg/0.5mL): Diagnosis of active PsA. PsA (Initial - 90mg/1mL): Diagnosis of active PsA. Patient's weight is greater than 100 kg (220 lbs). Diagnosis of co-existent moderate to severe psoriasis. PsA (Initial regardless of dose): One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Crohn's disease (CD) (initial): Diagnosis of moderately to severely active Crohn's disease. Will be used as a maintenance dose following the intravenous induction dose. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | Plaque psoriasis (Initial): Prescribed by or in consultation with a dermatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. CD and UC (initial): Prescribed by or in consultation with a gastroenterologist.  |
| Coverage<br>Duration               | All indications (initial, reauth): plan year.  |

| Other Criteria                      | Ulcerative colitis (UC) (initial): Diagnosis of moderately to severely active UC. Will be used as a maintenance dose following the intravenous induction dose. All indications (Reauth): Patient demonstrates positive clinical response to therapy. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### **STIVARGA**

### **Products Affected**

• Stivarga

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Colorectal Cancer (CRC): Diagnosis of advanced or metastatic colorectal cancer. Trial and failure, contraindication, or intolerance to treatment with all the following: oxaliplatin-based chemotherapy, irinotecan-based chemotherapy, fluoropyrimidine-based chemotherapy, and anti-VEGF therapy-based chemotherapy. One of the following: 1) Tumor is RAS mutant-type OR 2) Tumor is RAS wild-type and trial and failure, contraindication, or intolerance to anti-EGFR therapy [e.g., Erbitux (cetuximab), Vectibix (panitumumab)]. Gastrointestinal stromal tumor (GIST): Diagnosis of gross residual (R2 resection), unresectable primary, tumor rupture, or recurrent/metastatic GIST. One of the following: 1) Patient has succinate dehydrogenase (SDH) deficient GIST or 2) Trial and failure, contraindication, or intolerance to imatinib mesylate and sunitinib malate. Hepatocellular Carcinoma (HCC): Diagnosis of HCC. Used as subsequent-line therapy for disease progression. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **SUNITINIB**

#### **Products Affected**

#### • Sunitinib Malate

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Gastrointestinal stromal tumor (GIST): Diagnosis of GIST. One of the following: 1) History of disease progression on, contraindication, or intolerance to Gleevec (imatinib), Stivarga (regorafenib), or standard dose Qinlock (ripretinib) or 2) Succinate dehydrogenase (SDH)-deficient GIST. Renal Cell Carcinoma (RCC): Diagnosis of RCC. One of the following: 1) Disease has relapsed, 2) Both of the following: used in adjuvant setting and patient has a high risk of recurrence following nephrectomy, or 3) Disease is advanced. Islet Cell Tumors/Pancreatic Neuroendocrine Tumors (pNET): Diagnosis of islet cell tumors/progressive pNET. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **SYMPAZAN**

#### **Products Affected**

#### • Sympazan

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Lennox-Gastaut syndrome: Diagnosis of Lennox-Gastaut syndrome. Used for the adjunctive treatment of seizures associated with Lennox-Gastaut syndrome. Dravet syndrome: Diagnosis of seizures associated with Dravet syndrome (DS). Used in combination with Diacomit. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TABLOID**

#### **Products Affected**

#### • Tabloid

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of acute myeloid leukemia.                                   |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

### **TABRECTA**

#### **Products Affected**

#### • Tabrecta

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of non-small cell lung cancer (NSCLC). One of the following: a) Presence of mesenchymal-epithelial transition (MET) exon 14 skipping positive tumors or b) High level MET amplification in lung cancer. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# TADALAFIL (BPH)

### **Products Affected**

• Tadalafil TABS 2.5MG, 5MG

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Excluded if used for the treatment of erectile dysfunction only.   |
| Required<br>Medical<br>Information  | Benign prostatic hyperplasia (BPH): Diagnosis of BPH. Male Gender.   |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | BPH: plan year   |
| Other Criteria                      | BPH: Trial and failure, contraindication, or intolerance to two formulary alpha blockers (e.g., tamsulosin, alfuzosin). 2.5mg strength: Patient has renal insufficiency. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# TADALAFIL (PAH)

### **Products Affected**

• Alyq

#### • Tadalafil TABS 20MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH. PAH is symptomatic. One of the following: A) Diagnosis of PAH was confirmed by right heart catheterization or B) Patient is currently on any therapy for the diagnosis of PAH. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | PAH: Prescribed by or in consultation with a pulmonologist or cardiologist.   |
| Coverage<br>Duration                | PAH: plan year  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TAFAMIDIS**

### **Products Affected**

• Vyndamax

• Vyndaqel

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# **TAFINLAR**

### **Products Affected**

#### • Tafinlar

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma AND cancer is BRAFV600 mutant type. Adjuvant Treatment for Melanoma: Diagnosis of melanoma. Cancer is BRAF V600E or V600K mutant type. Involvement of lymph nodes following complete resection. Used as adjunctive therapy. Medication is used in combination with Mekinist (trametinib). Non-small Cell Lung Cancer (NSCLC): All of the following: diagnosis of non-small cell lung cancer AND disease is one of the following: metastatic, advanced, or recurrent AND cancer is BRAF V600E mutant type AND medication is used in combination with Mekinist (trametinib). Anaplastic Thyroid Cancer (ATC): Diagnosis of anaplastic thyroid cancer. One of the following: 1) Disease is one of the following: metastatic, locally advanced, or unresectable OR 2) Prescribed as adjuvant therapy following resection. Cancer is BRAF V600E mutant type. Medication is used in combination with Mekinist (trametinib). Solid tumors: Presence of solid tumor. Disease is unresectable or metastatic. Patient has progressed on or following prior systemic treatment (e.g., carboplatin, 5-fluorouracil, paclitaxel). Cancer is BRAF V600E mutant type. Medication is used in combination with Mekinist (trametinib). Lowgrade glioma: Diagnosis of low-grade glioma. Cancer is BRAF V600E mutant type. Medication is used in combination with Mekinist (trametinib). |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan year  |
| Other Criteria                     | Approve for continuation of prior therapy if within the past 120 days.   |

# **TAGRISSO**

### **Products Affected**

### Tagrisso

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

### **TALZENNA**

#### **Products Affected**

#### • Talzenna

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Breast cancer: Diagnosis of breast cancer. Disease is one of the following: a) locally advanced or b) metastatic. Presence of deleterious or suspected deleterious germline BRCA-mutations as detected by the FDA-approved companion diagnostic for Talzenna. Prostate cancer: Diagnosis of prostate cancer. Disease is HRR gene-mutated. Disease is metastatic castration-resistant. Taken in combination with Xtandi (enzalutamide). Used in combination with a gonadotropin-releasing hormone (GnRH) analog [e.g., Lupron (leuprolide), Zoladex (goserelin), Trelstar (triptorelin), Vantas (histrelin), Firmagon (degarelix)] OR Patient has had bilateral orchiectomy. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TASIGNA**

### **Products Affected**

### • Nilotinib Hydrochloride

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of chronic myeloid leukemia.                                 |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

# TASIMELTEON CAPSULE

### **Products Affected**

#### • Tasimelteon

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Non-24-hour sleep-wake disorder: Diagnosis of non-24-hour sleep-wake disorder (also known as free-running disorder, free-running or non-entrained type circadian rhythm sleep disorder, or hypernychthemeral syndrome). Smith-Magenis Syndrome (SMS): Diagnosis of Smith-Magenis Syndrome (SMS). Patient is experiencing nighttime sleep disturbances (i.e., difficulty falling asleep, frequent nighttime waking and early waking). |
| Age Restrictions                    | SMS: 16 years of age or older  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Non-24, SMS: 6 months  |
| Other Criteria                      | Non-24: Patient is totally blind (has no light perception).  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **TAZAROTENE**

### **Products Affected**

#### • Tazarotene CREA 0.1%

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | All indications: Excluded if treatment for cosmetic purposes.                              |
| Required<br>Medical<br>Information  | Acne vulgaris: Diagnosis of acne vulgaris (i.e., acne). Psoriasis: Diagnosis of psoriasis. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.                               |

# **TAZVERIK**

#### **Products Affected**

#### • Tazverik

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Epithelioid sarcoma: Diagnosis of epithelioid sarcoma. Disease is one of the following: metastatic or locally advanced. Patient is not eligible for complete resection. Follicular lymphoma: Diagnosis of relapsed or refractory follicular lymphoma. One of the following: a) Subsequent therapy in EZH2 mutation positive disease after two prior therapies, b) Second-line therapy irrespective of EZH2 mutation status for older or infirm patients with indications for treatment (i.e., other therapy options are not expected to be tolerable), or c) Third-line and/or subsequent therapy (if not previously given) irrespective of EZH2 mutation status in patients with indications for treatment. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## ТЕРМЕТКО

### **Products Affected**

### • Tepmetko

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: recurrent, advanced, or metastatic. Tumor is MET exon 14 skipping mutation positive. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TERIPARATIDE**

### **Products Affected**

• Bonsity

• Teriparatide

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Postmenopausal osteoporosis or osteopenia or men with primary or hypogonadal osteoporosis or osteopenia (initial): Diagnosis of one of the following: a) postmenopausal osteoporosis or osteopenia or b) primary or hypogonadal osteoporosis or osteopenia. One of the following: Set I) Both of the following: A) Bone mineral density (BMD) T-score of -2.5 or lower in the lumbar spine, femoral neck, total hip, or radius (one-third radius site) AND B) One of the following: 1) history of low-trauma fracture of the hip, spine, proximal humerus, pelvis, or distal forearm, or 2) trial and failure, contraindication, or intolerance (TF/C/I) to one osteoporosis treatment (e.g., alendronate, risedronate, zoledronic acid, Prolia [denosumab]), or Set II) Both of the following: A) BMD T-score between -1.0 and -2.5 in the lumbar spine, femoral neck, total hip, or radius (one-third radius site) AND B) One of the following: 1) history of low-trauma fracture of the hip, spine, proximal humerus, pelvis, or distal forearm, or 2) both of the following: i) TF/C/I to one osteoporosis treatment (e.g., alendronate, risedronate, zoledronic acid, Prolia [denosumab]) and ii) One of the following FRAX 10-year probabilities: a) Major osteoporotic fracture at 20% or more in the U.S., or the country-specific threshold in other countries or regions, or b) Hip fracture at 3% or more in the U.S., or the country-specific threshold in other countries or regions. Glucocorticoid-Induced Osteoporosis: See Other Criteria section. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | All indications (initial, reauth): plan year.   |

#### Other Criteria Glucocorticoid-Induced Osteoporosis (initial): Diagnosis of glucocorticoid-induced osteoporosis. History of prednisone or its equivalent at a dose greater than or equal to 5mg/day for greater than or equal to 3 months. One of the following: 1) BMD T-score less than or equal to -2.5 based on BMD measurements from lumbar spine, femoral neck, total hip, or radius (one-third radius site), or 2) One of the following FRAX 10-year probabilities: a) Major osteoporotic fracture at 20% or more in the U.S., or the country-specific threshold in other countries or regions, or b) Hip fracture at 3% or more in the U.S., or the countryspecific threshold in other countries or regions, 3) History of one of the following fractures resulting from minimal trauma: vertebral compression fx, fx of the hip, fx of the distal radius, fx of the pelvis, or fx of the proximal humerus, or 4) One of the following: a) glucocorticoid dosing of at least 30 mg per day, or b) cumulative glucocorticoid dosing of at least 5 grams per year. TF/C/I to one bisphosphonate (e.g., alendronate). All uses (initial, reauth): One of the following: 1) Treatment duration of parathyroid hormones (e.g., teriparatide, Tymlos [abaloparatide]) has not

#### Prerequisite Therapy Required

Criteria DOES require use of a prerequisite Part D drug.

[abaloparatide]).

exceeded a total of 24 months during the patient's lifetime, or 2) Patient remains at or has returned to having a high risk for fracture despite a total of 24 months of use of parathyroid hormones (e.g., teriparatide, Tymlos

# **TETRABENAZINE**

#### **Products Affected**

#### • Tetrabenazine

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Huntington's Disease: Diagnosis of chorea in patients with Huntington's disease. Tardive dyskinesia: Diagnosis of tardive dyskinesia. One of the following: 1) Patient has persistent symptoms of tardive dyskinesia despite a trial of dose reduction, tapering, or discontinuation of the offending medication or 2) Patient is not a candidate for a trial of dose reduction, tapering, or discontinuation of the offending medication. Tourette's syndrome: Patient has tics associated with Tourette's syndrome. Trial and failure, contraindication, or intolerance to Haldol (haloperidol). |
| Age Restrictions                    | Tardive dyskinesia: Age greater than or equal to 18 years.   |
| Prescriber<br>Restrictions          | Huntington's: Prescribed by or in consultation with a neurologist. Tardive dyskinesia, Tourette's: Prescribed by or in consultation with a neurologist or psychiatrist.  |
| Coverage<br>Duration                | Plan year.   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **THALOMID**

### **Products Affected**

• Thalomid CAPS 100MG, 50MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Erythema Nodosum Leprosum (ENL): Diagnosis (Dx) of moderate to severe ENL. One of the following: used for acute treatment OR used as maintenance therapy for prevention & suppression of cutaneous manifestations of ENL recurrence. Multiple Myeloma (MM): Dx of multiple myeloma. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TIBSOVO**

### **Products Affected**

#### • Tibsovo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Relapsed or refractory Acute Myeloid Leukemia (AML): Diagnosis of AML. Disease is relapsed or refractory. AML is isocitrate dehydrogenase-1 (IDH1) mutation-positive. Newly-Diagnosed AML: Diagnosis of newly-diagnosed AML. AML is isocitrate dehydrogenase-1 (IDH1) mutation-positive. One of the following: 1) Patient is greater than or equal to 75 years old OR 2) Patient has comorbidities that preclude the use of intensive induction chemotherapy OR 3) Patient is greater than or equal to 60 years old and not a candidate for or declines intensive induction therapy OR 4) Patient is greater than or equal to 60 years old and receiving post-induction therapy following response to previous lower intensity therapy. Cholangiocarcinoma: Diagnosis of cholangiocarcinoma. Disease is locally advanced, unresectable, or metastatic. Cholangiocarcinoma is IDH1 mutation-positive. Disease has progressed on or after systemic treatment. Myelodysplastic Syndromes (MDS): Diagnosis of MDS. Disease is one of the following: relapsed or refractory. MDS is IDH1 mutation-positive. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# TOBI PODHALER

### **Products Affected**

#### • Tobi Podhaler

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                          |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | N/A  |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug. |

# **TOLVAPTAN (SAMSCA)**

### **Products Affected**

#### • Tolvaptan TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of significant hyponatremia (euvolemic or hypervolemic). Treatment has been initiated or re-initiated in a hospital setting prior to discharge within the past 30 days. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Length of authorization: 30 days  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## TOPICAL RETINOIDS

#### **Products Affected**

• Tretinoin CREA

- Tretinoin GEL 0.01%, 0.025% Tretinoin Microsphere GEL 0.1%

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.                           |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | All indications: Excluded if treatment for cosmetic purposes. |
| Required<br>Medical<br>Information  | Acne vulgaris: Diagnosis of acne vulgaris (i.e., acne).       |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TORPENZ**

### **Products Affected**

### • Torpenz

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Advanced Neuroendocrine Tumors (NET): Diagnosis (Dx) of neuroendocrine tumors of pancreatic origin, gastrointestinal origin, lung origin, or thymic origin. Disease is progressive. Disease is unresectable, locally advanced or metastatic. Advanced Renal Cell Carcinoma/Kidney Cancer: Dx of advanced renal cell cancer/kidney cancer. Disease is one of the following: (1) relapsed or (2) stage IV disease. Renal angiomyolipoma with tuberous sclerosis complex (TSC): Dx of renal angiomyolipoma and TSC, not requiring immediate surgery. Subependymal Giant Cell Astrocytoma (SEGA) with tuberous sclerosis (TS): Dx of SEGA associated with TS. Patient is not a candidate for curative surgical resection. Breast Cancer: Dx of recurrent or metastatic breast cancer. One of the following: Disease is hormone receptor positive (HR+) [i.e., estrogen-receptor-positive (ER+) or progesterone-receptor-positive (PR+)] OR both of the following: disease is HR- and disease has clinical characteristics that predict a HR+ tumor. Disease is HER2-negative. One of the following: Patient is a postmenopausal woman, patient is a premenopausal woman being treated with ovarian ablation/suppression, or patient is male. One of the following: A) Both of the following: a) one of the following: 1) Disease progressed while on or within 12 months of non-steroidal aromatase inhibitor [e.g., Arimidex (anastrozole), Femara (letrozole)] therapy or 2) patient was treated with tamoxifen at any time AND b) Used in combination with Aromasin (exemestane) OR B) Used in combination with Fulvestrant or Tamoxifen. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan year  |

| Other Criteria                      | All uses: Approve for continuation of prior therapy if within the past 120 days. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.                         |

# TREMFYA (I)

#### **Products Affected**

• Tremfya INJ 100MG/ML, 200MG/2ML

• Tremfya Induction Pack For Crohns Disease/ulcerative Colitis

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Plaque Psoriasis (initial): Diagnosis of chronic moderate-to-severe plaque psoriasis. One of the following: at least 3% body surface area (BSA) involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Psoriatic Arthritis (PsA) (initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Ulcerative Colitis (UC) (Initial): Diagnosis of moderately to severely active UC. Will be used as a maintenance dose following the intravenous induction doses. Crohn's Disease (CD) (initial) in Other Criteria element. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | Plaque Psoriasis (initial): Prescribed by or in consultation with a dermatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. CD, UC (Initial): Prescribed by or in consultation with a gastroenterologist.  |
| Coverage<br>Duration               | All indications (initial, reauth): plan year  |

| Other Criteria                      | Crohn's Disease (CD) (initial): Diagnosis of moderately to severely active CD. One of the following: 1) Will be used as a maintenance dose following the intravenous induction doses, OR 2) Both of the following: a) Will be used as induction dosing, and b) One of the following: frequent diarrhea and abdominal pain, at least 10% weight loss, complications (eg, obstruction, fever, abdominal mass), abnormal lab values (eg, CRP), or CD Activity Index (CDAI) greater than 220. All indications (Reauth): Patient demonstrates positive clinical response to therapy. |
|-------------------------------------|---|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **TRIENTINE**

### **Products Affected**

# • Trientine Hydrochloride

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of Wilson's disease (i.e., hepatolenticular degeneration). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.         |

# TRIKAFTA

### **Products Affected**

#### • Trikafta

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of cystic fibrosis. Submission of laboratory results documenting that the patient has at least one of the following mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene: F508del mutation OR A mutation in the CFTR gene that is responsive based on clinical and/or in vitro data. |
| Age Restrictions                    | Initial: For granule packets: patient is at least 2 to less than 6 years of age. For tablets: patient is 6 years of age or older  |
| Prescriber<br>Restrictions          | Initial/Reauth: Prescribed by or in consultation with a pulmonologist or specialist affiliated with a CF care center  |
| Coverage<br>Duration                | Initial, Reauth: Plan Year.   |
| Other Criteria                      | Reauthorization: Patient demonstrates positive clinical response to therapy.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **TRULICITY**

## **Products Affected**

### • Trulicity

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | Excluded if used for weight loss only.   |
| Required<br>Medical<br>Information  | Diabetes Mellitus (DM): Submission of medical records (e.g., chart notes) confirming diagnosis of type 2 DM. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# TRUMENBA

## **Products Affected**

#### • Trumenba

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# TRUQAP

### **Products Affected**

### • Truqap TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of breast cancer. Disease is one of the following: locally advanced or metastatic. Will be taken in combination with fulvestrant. Disease is hormone receptor (HR)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Patient has one or more PIK3CA/AKT1/PTEN-alterations. One of the following: A) Following progression on at least one endocrine-based regimen in the metastatic setting (e.g., anastrozole, letrozole, exemestane, tamoxifen, etc.) OR B) Recurrence on or within 12 months of completing adjuvant therapy (e.g., chemotherapy). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan Year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **TUKYSA**

## **Products Affected**

• Tukysa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Breast cancer: Diagnosis of breast cancer. Disease is one of the following: a) advanced unresectable or b) metastatic. Disease is human epidermal growth factor receptor 2 (HER2)-positive. Used in combination with trastuzumab and capecitabine. Patient has been previously treated with an anti-HER2-based regimen (e.g., trastuzumab, pertuzumab, adotrastuzumab emtansine) in the metastatic setting. Colorectal cancer: Diagnosis of colorectal cancer (HER2-amplified and RAS and BRAF wild-type). Disease is HER2-positive. Disease is one of the following: a) advanced, b) unresectable, c) metastatic. One of the following: a) patient has previously been treated with one of the following regimens: i) fluoropyrimidine-based chemotherapy, ii) oxaliplatin-based chemotherapy, iii) irinotecan-based chemotherapy or b) patient is not appropriate for intensive therapy. Used in combination with trastuzumab. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **TURALIO**

## **Products Affected**

#### • Turalio CAPS 125MG

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Tenosynovial Giant Cell Tumor (TGCT): Diagnosis of TGCT. Patient is symptomatic. Patient is not a candidate for surgery due to worsening functional limitation or severe morbidity with surgical removal. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# TYENNE SC

### **Products Affected**

• Tyenne INJ 162MG/0.9ML

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Rheumatoid Arthritis (RA) (Initial): Diagnosis of moderately to severely active RA. One of the following: 1) Trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: Enbrel (etanercept), one formulary adalimumab product, Rinvoq (upadacitinib), or Xeljanz/Xeljanz XR (tofacitinib), OR 2) for continuation of prior therapy if within the past 120 days. Systemic Juvenile Idiopathic Arthritis (SJIA) (Initial): Diagnosis of active SJIA. TF/C/I to one of the following conventional therapies at maximally tolerated doses: a) minimum duration of a one month trial of a nonsteroidal anti-inflammatory drug (NSAID) (eg, ibuprofen, naproxen), or b) minimum duration of a 2-week trial of a systemic glucocorticoid (eg, prednisone). Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Diagnosis of active PJIA. One of the following: 1) TF/C/I to two of the following: Enbrel (etanercept), one formulary adalimumab product, Rinvoq/Rinvoq LQ, Xeljanz (tofacitinib), OR 2) for continuation of prior therapy if within the past 120 days. Giant Cell Arteritis (GCA) (Initial): Diagnosis of GCA. TF/C/I to a glucocorticoid (eg, prednisone). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | RA, SJIA, PJIA, GCA (Initial): Prescribed by or in consultation with a rheumatologist.  |
| Coverage<br>Duration                | RA, SJIA, PJIA, GCA (Initial, Reauth): Plan year  |
| Other Criteria                      | RA, SJIA, PJIA, GCA (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **TYMLOS**

### **Products Affected**

#### • Tymlos

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | One of the following diagnoses: 1) Postmenopausal osteoporosis or osteopenia, OR 2) Primary or hypogonadal osteoporosis or osteopenia. One of the following: Set I) Both of the following: A) Bone mineral density (BMD) T-score of -2.5 or lower in the lumbar spine, femoral neck, total hip, or radius (one-third radius site) AND B) One of the following: 1) history of low-trauma fracture of the hip, spine, proximal humerus, pelvis, or distal forearm, or 2) trial and failure, contraindication, or intolerance (TF/C/I) to one osteoporosis treatment (e.g., alendronate, risedronate, zoledronic acid, Prolia [denosumab]), or Set II) Both of the following: A) BMD T-score between -1.0 and -2.5 in the lumbar spine, femoral neck, total hip, or radius (one-third radius site) AND B) One of the following: 1) history of low-trauma fracture of the hip, spine, proximal humerus, pelvis, or distal forearm, or 2) both of the following: i) TF/C/I to one osteoporosis treatment (e.g., alendronate, risedronate, zoledronic acid, Prolia [denosumab]) and ii) one of the following FRAX (Fracture Risk Assessment Tool) 10-year probabilities: a) major osteoporotic fracture at 20% or more in the U.S., or the country-specific threshold in other countries or regions, or b) hip fracture at 3% or more in the U.S., or the country-specific threshold in other countries or regions. Treatment duration of parathyroid hormones (e.g., teriparatide, Tymlos [abaloparatide]) has not exceeded a total of 24 months during the patient's lifetime. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan year (up to 24 months per lifetime)   |
| Other Criteria                     | N/A  |

| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug. |
|-------------------------------------|--|
|-------------------------------------|--|

# **UBRELVY**

## **Products Affected**

• Ubrelvy

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of migraine with or without aura. Will be used for the acute treatment of migraine. Trial and failure or intolerance to one triptan (e.g., eletriptan, rizatriptan, sumatriptan) or a contraindication to all triptans. Medication will not be used in combination with another CGRP inhibitor for the acute treatment of migraines. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Initial, Reauth: Plan year  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy. Will not be used for preventive treatment of migraine. Medication will not be used in combination with another CGRP inhibitor for the acute treatment of migraines.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **VALCHLOR**

## **Products Affected**

#### • Valchlor

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Mycosis fungoides-type cutaneous T-cell lymphoma (MF-CTCL): Both of the following: 1) diagnosis of Stage IA MF-CTCL OR diagnosis of Stage IB MF-CTCL AND 2) patient has received at least one prior skindirected therapy (e.g., topical corticosteroids, bexarotene topical gel [Targretin topical gel], etc.). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## VALTOCO

#### **Products Affected**

- Valtoco 10 Mg Dose
- Valtoco 15 Mg Dose

- Valtoco 20 Mg Dose Valtoco 5 Mg Dose

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Epilepsy: Diagnosis of epilepsy. Frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) that are distinct from a patient's usual seizure pattern. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## VANDETANIB

### **Products Affected**

• Caprelsa

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Thyroid Cancer: Diagnosis of medullary thyroid cancer.                 |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

## VANFLYTA

## **Products Affected**

### • Vanflyta

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute Myeloid Leukemia (AML): Diagnosis of AML. Disease is FLT3 internal tandem duplication (ITD) positive. Vanflyta will be used in combination with standard cytarabine and anthracycline induction and cytarabine consolidation, and as maintenance monotherapy following consolidation chemotherapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# VAXCHORA

#### **Products Affected**

#### • Vaxchora

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Vaccine is being used for active immunization against disease caused by Vibrio cholerae serogroup O1. Patient is traveling to a cholera-affected area. Age 2 through 64 years. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 3 months   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# VENCLEXTA

### **Products Affected**

• Venclexta

• Venclexta Starting Pack

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL): Diagnosis of CLL or SLL. Acute Myeloid Leukemia (AML): One of the following: 1) Diagnosis of newly diagnosed AML. Used in combination with azacitidine, or decitabine, or low-dose cytarabine. One of the following: a) used as treatment induction in candidates for intensive induction therapy, b) used as treatment induction in candidates for lower-intensity induction therapy, c) used as follow-up after induction therapy following response to previous lower intensity therapy with the same regimen, or d) used as consolidation therapy as continuation of lower-intensity regimen used for induction. 2) Diagnosis of relapsed/refractory acute myeloid leukemia (AML). Relapse is greater than or equal to 12 months from most recent disease remission. Venclexta therapy to be given in combination with the patients previous initial successful induction regimen (e.g., azacitidine, decitabine, low-dose cytarabine, etc.) |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | Plan year   |
| Other Criteria                     | AML (cont.): 3) Diagnosis of blastic plasmacytoid dendritic cell neoplasm (BPDCN) - acute myeloid leukemia (AML). Considered systemic disease and therapy is given as palliative intent. Patient has low performance and/or nutritional status (i.e., serum albumin less than 3.2 g/dL, not a candidate for intensitve remission therapy of Elzonris). Venclexta therapy will be given in combination with azactidine, decitabine, or low-dose cytarabine. Approve for continuation of prior therapy if within the past 120 days.   |

| Therapy<br>Required |  | Criteria DOES NOT require use of a prerequisite Part D drug. | 1 0 |
|---------------------|--|--|-----|
|---------------------|--|--|-----|

# VEOZAH

#### **Products Affected**

#### • Veozah

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of moderate to severe vasomotor symptoms due to menopause. Prescriber attests that baseline serum alanine aminotransferase (ALT), serum aspartate aminotransferase (AST) and total bilirubin levels are less than 2 times the upper limit of normal (ULN) prior to initiating Veozah.    |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Initial, Reauth: 6 months   |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy. Both of the following within the past 3 months: a) Transaminase elevations are less than 5 times the ULN, and b) Both transaminase elevations are less than 3 times the ULN and the total bilirubin level is less than 2 times the ULN. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# VERQUVO

### **Products Affected**

#### • Verquvo

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Chronic Heart Failure (CHF) (initial): Diagnosis of CHF. Patient has an ejection fraction less than 45 percent. Patient has New York Heart Association (NYHA) Class II, III, or IV symptoms. One of the following: A) Patient was hospitalized for heart failure within the last 6 months, or B) Patient used outpatient intravenous diuretics (e.g., bumetanide, furosemide) for heart failure within the last 3 months. Trial and failure, contraindication, or intolerance to two of the following at a maximally tolerated dose: A) One of the following: 1) Angiotensin converting enzyme (ACE) inhibitor (e.g., captopril, enalapril), 2) Angiotensin II receptor blocker (ARB) (e.g., candesartan, valsartan), or 3) Angiotensin receptor-neprilysin inhibitor (ARNI) [e.g., Entresto (sacubitril and valsartan)], B) One of the following: 1) bisoprolol, 2) carvedilol, or 3) metoprolol succinate extended release, C) Sodium-glucose co-transporter 2 (SGLT2) inhibitor [e.g., Jardiance (empagliflozin), Farxiga (dapagliflozin), Xigduo XR (dapagliflozin and metformin)], or D) Mineralocorticoid receptor antagonist (MRA) [e.g., eplerenone, spironolactone]. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | CHF (initial): Prescribed by or in consultation with a cardiologist.  |
| Coverage<br>Duration                | CHF (initial, reauth): plan year  |
| Other Criteria                      | CHF (reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## VERZENIO

#### **Products Affected**

#### • Verzenio

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Advanced, Recurrent, or Metastatic Breast Cancer: Diagnosis of advanced, recurrent, or metastatic breast cancer. Disease is hormone receptor (HR)-positive and human epidermal growth factor receptor 2 (HER2)-negative. One of the following: a) used in combination with an aromatase inhibitor (e.g., anastrozole, letrozole, exemestane), OR b) used in combination with Faslodex (fulvestrant) OR c) used as monotherapy and disease has progressed following endocrine therapy and patient has already received at least one prior chemotherapy regimen. Early Breast Cancer: Diagnosis of early breast cancer at high risk of recurrence. Disease is hormone receptor (HR)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Used in combination with one of the following endocrine therapies: 1) tamoxifen or 2) aromatase inhibitor (e.g., anastrozole, letrozole, exemestane). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

### VIGABATRIN

### **Products Affected**

• Vigabatrin

• Vigpoder

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Complex Partial Seizures (CPS): For use as adjunctive therapy. Infantile Spasms (IS): Diagnosis of infantile spasms.  |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. CPS: Trial and failure, contraindication, or intolerance (TF/C/I) to two formulary anticonvulsants [eg, Lamictal (lamotrigine), Depakene (valproic acid), Dilantin (phenytoin)]. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## VIGAFYDE

### **Products Affected**

• Vigafyde

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

### VITRAKVI

#### **Products Affected**

#### • Vitrakvi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Presence of solid tumors. Disease is positive for neurotrophic receptor tyrosine kinase (NTRK) gene fusion (e.g. ETV6-NTRK3, TPM3-NTRK1, LMNA-NTRK1, etc.). Disease is without a known acquired resistance mutation [e.g., TRKA G595R, G623R, G696A, F617L]. Disease is one of the following: metastatic or unresectable. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

### **VIZIMPRO**

#### **Products Affected**

### • Vizimpro

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is recurrent, advanced, or metastatic. Disease is positive for one of the following epidermal growth factor receptor (EGFR) mutations: a) exon 19 deletion, b) exon 21 L858R substitution, c) S768I, d) L861Q, or e) G719X. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## Vonjo

### **Products Affected**

#### • Vonjo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Patient has been diagnosed with one of the following: a) Primary myelofibrosis, b) Post-polycythemia vera myelofibrosis, c) Post-essential thrombocythemia myelofibrosis, OR d) Accelerated/blast phase myeloproliferative neoplasm. 1) One of the following: A) Both of the following: i) Patient has symptomatic lower-risk myelofibrosis AND ii) Patient has a platelet count below 50 x 10^9/L, or B) All of the following: i) Patient has higher-risk myelofibrosis and ii) Patient is not a transplant candidate or transplant not currently feasible and iii) One of the following: a) Patient has a platelet count less than 50 x 10^9/L or b) Both of the following: i) Patient has symptomatic splenomegaly and/or constitutional symptoms and ii) Patient has a platelet count greater than or equal to 50 x 10^9/L, or C) Used for treatment of myelofibrosis-associated anemia, or D) Used for splenomegaly and other disease-related symptoms in one of the following: i) Continued near the start of conditioning therapy of transplant candidates, or ii) Palliation in combination with hypomethylating agents (e.g., azacitidine or decitabine) as bridging therapy prior to transplant, or if not a candidate for transplant. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# VOQUEZNA

### **Products Affected**

• Voquezna

- Voquezna Dual Pak
- Voquezna Triple Pak

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Helicobacter pylori (H. pylori) (Voquezna Dual Pak, Voquezna Triple Pak): Diagnosis of H. pylori infection. Trial and failure, contraindication, or intolerance to bismuth quadruple therapy (e.g., bismuth and metronidazole and tetracycline and proton pump inhibitor [PPI]). H. pylori (Voquezna): Diagnosis of H. pylori infection. One of the following: a) Used in combination with amoxicillin and clarithromycin for the treatment of H. pylori infection, or b) Used in combination with amoxicillin for the treatment of H. pylori infecion. Trial and failure, contraindication, or intolerance to bismuth quadruple therapy (e.g., bismuth and metronidazole and tetracycline and proton pump inhibitor [PPI]). Healing and Relief of Heartburn associated with Erosive Esophagitis (HRH) (Voquezna): Diagnosis of erosive esophagitis. Used for healing of all grades of erosive esophagitis and relief of heartburn associated with erosive esophagitis. Trial and failure, contraindication, or intolerance to TWO of the following: a) generic omeprazole, b) generic oral pantoprazole tablet, or c) generic esomeprazole granules. Maintenance of Healing and Relief of Heartburn associated with Erosive Esophagitis (MHRH) (Voquezna): Used to maintain healing and relief of heartburn associated with erosive esophagitis. Trial and failure, contraindication, or intolerance to TWO of the following: a) generic omeprazole, b) generic oral pantoprazole tablet, or c) generic esomeprazole granules. |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | N/A   |
| Coverage<br>Duration               | H. pylori, NERD: 1 mo. HRH: 8 wks. MHRH: 6 mos.   |

| Other Criteria                      | Relief of Heartburn associated with non-erosive Gastroesophageal Reflux Disease (NERD) (Voquezna): Diagnosis of non-erosive Gastroesophageal Reflux Disease. Both of the following: a) Patient has history of heartburn for at least 6 months and b) Heartburn symptoms are present for at least 4 days during any consecutive 7-day period. Trial and failure, contraindication, or intolerance to TWO of the following: a) generic omeprazole, b) generic oral pantoprazole tablet, or c) generic esomeprazole granules. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## Voranigo

#### **Products Affected**

### • Voranigo

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of Astrocytoma or Oligodendroglioma. Presence of an isocitrate dehydrogenase-1 (IDH1) or isocitrate dehydrogenase-2 (IDH2) mutation. History of one of the following: a) Biopsy, b) Sub-total resection, or c) Gross total resection. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# **VORICONAZOLE INJECTION**

#### **Products Affected**

#### • Voriconazole INJ

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Invasive aspergillosis: Diagnosis of invasive aspergillosis (IA). Candidemia: Diagnosis of candidemia. One of the following: (1) patient is non-neutropenic or (2) infection is located in skin, abdomen, kidney, bladder wall, or wounds. Esophageal Candidiasis: Diagnosis of esophageal candidiasis. Mycosis: Diagnosis of fungal infection caused by Scedosporium apiospermum (asexual form of Pseudallescheria boydii) or Fusarium spp. including Fusarium solani. For fusariosis: Patient is intolerant of, or refractory to, other therapy (e.g., liposomal amphotericin B, amphotericin B lipid complex). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | 12 weeks  |
| Other Criteria                      | N/A   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## Vosevi

#### **Products Affected**

#### • Vosevi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Criteria will be applied consistent with current AASLD/IDSA guideline. All patients: Diagnosis of chronic hepatitis C, patient is without decompensated liver disease (defined as Child-Pugh Class B or C), and patient is not receiving Vosevi in combination with another HCV direct acting antiviral agent [e.g., Harvoni, Zepatier]. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | 12 to 24 weeks. Criteria will be applied consistent with current AASLD/IDSA guideline.   |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## **VOTRIENT**

### **Products Affected**

### • Pazopanib Hydrochloride

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Renal Cell Carcinoma (RCC): Diagnosis of RCC. Soft tissue sarcoma (STS): Diagnosis of advanced STS. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.                              |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## Vowst

#### **Products Affected**

#### • Vowst

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of recurrent clostridioides difficile infection (CDI) as defined by both of the following: 1) Presence of diarrhea defined as a passage of 3 or more loose bowel movements within a 24-hour period for two consecutive days, and 2) A positive stool test for C.difficile toxin or toxigenic C.difficile. Patient has a history of two or more recurrent episodes of CDI within 12 months. All of the following: 1) Patient has completed at least 10 consecutive days of one of the following antibiotic therapies 2-4 days prior to initiating Vowst: oral vancomycin or Dificid (fidaxomicin), 2) Patient has completed the recommended course of magnesium citrate the day before and at least 8 hours prior to initiating Vowst, and 3) Previous episode of CDI is under control (e.g., less than 3 unformed/loose [i.e., Bristol Stool Scale type 6-7] stools/day for 2 consecutive days). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a gastroenterologist or infectious disease specialist.   |
| Coverage<br>Duration                | 14 days  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## WELIREG

### **Products Affected**

### • Welireg

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | von Hippel-Lindau (VHL) disease: Diagnosis of von Hippel-Lindau (VHL) disease. Patient requires therapy for one of the following: a) renal cell carcinoma (RCC), b) central nervous system (CNS) hemangioblastoma, or c) pancreatic neuroendocrine tumor (pNET). Patient does not require immediate surgery. Advanced Renal Cell Carcinoma (RCC): Diagnosis of advanced renal cell carcinoma. Disease has progressed after treatment with both of the following: a) Programmed death receptor 1 (PD-1) or programmed death ligand 1 (PD-L1) checkpoint inhibitor [e.g., Keytruda (pembrolizumab), Opdivo (nivolumab)] and b) Vascular endothelial growth factor tyrosine kinase inhibitor (VEGF-TKI) [e.g., Inlyta (axitinib), Lenvima (lenvatinib), Cabometyx (cabozantinib]. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## WINREVAIR

#### **Products Affected**

#### • Winrevair

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Pulmonary arterial hypertension (PAH): Diagnosis of PAH. PAH is symptomatic. Patient is currently on at least two therapies indicated for the treatment of PAH from the following different mechanisms of action, unless there is a contraindication or intolerance: a) Endothelin receptor antagonists (i.e., Bosentan, ambrisentan or macitentan) and b) Phosphodiesterase 5 inhibitors (i.e., Tadalafil or sildenafil). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a pulmonologist or cardiologist.   |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | N/A  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## XALKORI

### **Products Affected**

#### • Xalkori

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Non-small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Anaplastic Large Cell Lymphoma (ALCL): Diagnosis of systemic ALCL. Disease is relapsed or refractory. Tumor is anaplastic lymphoma kinase (ALK)-positive. Inflammatory Myofibroblastic Tumor (IMT): Diagnosis of IMT. Tumor is anaplastic lymphoma kinase (ALK)-positive. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **X**ATMEP

### **Products Affected**

### • Xatmep

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Acute lymphoblastic leukemia (ALL): Diagnosis of acute lymphoblastic leukemia (ALL). Polyarticular juvenile idiopathic arthritis (pJIA) (initial): Diagnosis of active polyarticular juvenile idiopathic arthritis. Trial and failure, contraindication, or intolerance to one nonsteroidal anti-inflammatory drug (NSAID) (e.g., diclofenac, ibuprofen, meloxicam, naproxen). |
| Age Restrictions                    | ALL: Patient is 18 years of age or younger. pJIA (initial): Patient is 18 years of age or younger.   |
| Prescriber<br>Restrictions          | pJIA (initial): Prescribed by or in consultation with a rheumatologist.  |
| Coverage<br>Duration                | ALL: plan year. pJIA (initial, reauth): plan year  |
| Other Criteria                      | Subject to Part B vs. Part D review. ALL: Approve for continuation of prior therapy if within the past 120 days. pJIA (reauth): Patient demonstrates positive clinical response to therapy   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# **X**COPRI

### **Products Affected**

#### • Xcopri

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.                                    |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of partial onset seizures.                                   |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days. |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.           |

## XELJANZ

### **Products Affected**

• Xeljanz

• Xeljanz Xr

| PA Criteria                        | Criteria Details  |
|------------------------------------|---|
| Indications                        | All Medically-accepted Indications.   |
| Off-Label Uses                     | N/A   |
| Exclusion<br>Criteria              | N/A   |
| Required<br>Medical<br>Information | Xeljanz tab/Xeljanz XR tab: Rheumatoid arthritis (RA) (Initial): Diagnosis of moderately to severely active RA. Minimum duration of a 3-month trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies at maximally tolerated doses: methotrexate, leflunomide, sulfasalazine. Xeljanz tab/Xeljanz XR tab: Psoriatic arthritis (PsA) (Initial): Diagnosis of active PsA. One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Xeljanz tab/Xeljanz XR tab: Ankylosing spondylitis (AS) (Initial): Diagnosis of active AS. Minimum duration of a one-month TF/C/I to one nonsteroidal anti-inflammatory drug (NSAID) (eg, ibuprofen, naproxen) at maximally tolerated doses. RA, PsA, AS (Initial): Patient has had an inadequate response or intolerance to one or more TNF inhibitors (eg, adalimumab, etanercept). Xeljanz tab/Xeljanz XR tab: Ulcerative colitis (UC) (Initial): Diagnosis of moderately to severely active UC. One of the following: greater than 6 stools per day, frequent blood in the stools, frequent urgency, presence of ulcers, abnormal lab values (eg, hemoglobin, ESR, CRP), OR dependent on, or refractory to, corticosteroids. Patient has had an inadequate response or intolerance to one or more TNF inhibitors (eg, adalimumab). Not used in combination with other Janus kinase (JAK) inhibitors, biological therapies for UC, or potent immunosuppressants (eg, azathioprine, cyclosporine). |
| Age Restrictions                   | N/A   |
| Prescriber<br>Restrictions         | RA, PJIA, AS (initial): Prescribed by or in consultation with a rheumatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. UC (initial): Prescribed by or in consultation with a gastroenterologist.   |

| Coverage<br>Duration                | RA, PJIA, PsA, AS, UC (initial, reauth): plan year.   |
|-------------------------------------|---|
| Other Criteria                      | Xeljanz: Polyarticular course juvenile idiopathic arthritis (PJIA) (Initial): Diagnosis of active polyarticular course juvenile idiopathic arthritis. Minimum duration of a 6-week TF/C/I to one of the following conventional therapies at maximally tolerated doses: leflunomide or methotrexate. Patient has had an inadequate response or intolerance to one or more TNF inhibitors (eg, adalimumab, etanercept). RA, PsA, AS, PJIA (Initial, Reauth): Not used in combination with other JAK inhibitors, biologic disease-modifying antirheumatic drugs (DMARDs), or potent immunosuppressants (eg, azathioprine, cyclosporine). All indications (Reauth): Patient demonstrates positive clinical response to therapy. UC (Reauth): Not used in combination with other JAK inhibitors, biological therapies for UC, or potent immunosuppressants (e.g., azathioprine, cyclosporine). |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## XERMELO

### **Products Affected**

#### • Xermelo

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Carcinoid syndrome diarrhea (Initial): Diagnosis of carcinoid syndrome diarrhea AND diarrhea is inadequately controlled by a stable dose of somatostatin analog (SSA) therapy (e.g., octreotide, lanreotide) for at least 3 months AND used in combination with SSA therapy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with an oncologist, endocrinologist, or gastroenterologist   |
| Coverage<br>Duration                | Initial: 6 months, Reauth: plan year   |
| Other Criteria                      | Carcinoid syndrome diarrhea (Reauthorization): Patient demonstrates positive clinical response to therapy. Will continue to be used in combination with SSA therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## **XIFAXAN**

### **Products Affected**

#### • Xifaxan

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Traveler's Diarrhea (TD): Diagnosis of traveler's diarrhea. Trial and failure, contraindication, or intolerance to one of the following: Cipro (ciprofloxacin), Levaquin (levofloxacin), ofloxacin, Zithromax (azithromycin) OR resistance to all of the following: Cipro (ciprofloxacin), Levaquin (levofloxacin), ofloxacin, Zithromax (azithromycin). Prophylaxis of Hepatic Encephalopathy (HE): Used for the prophylaxis of hepatic encephalopathy recurrence. One of the following: 1) Trial and failure, contraindication, or intolerance to lactulose OR 2) Add-on treatment to lactulose. Treatment of HE: Diagnosis of HE. Used for the treatment of HE. One of the following: 1) Trial and failure, contraindiation, or intolerance to lactulose OR 2) Add-on treatment to lactulose. Irritable Bowel Syndrome with Diarrhea (Initial): Diagnosis of irritable bowel syndrome with diarrhea (IBS-D). Trial and failure, contraindication or intolerance to an antidiarrheal agent (e.g., loperamide). |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | TD: 14 days (one treatment course). HE (Prophylaxis, Tx): plan year. IBS-D (initial/reauth): 2 wks   |
| Other Criteria                      | IBS-D Reauthorization: Patient experiences IBS-D symptom recurrence.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

# XOLAIR

### **Products Affected**

• Xolair

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

# XOLREMDI

### **Products Affected**

#### • Xolremdi

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of WHIM (warts, hypogammaglobulinemia, infections and myelokathexis) syndrome. Patient has genotype confirmed variant of CXCR4. Patient has an absolute neutrophil count (ANC) less than or equal to $500 \text{ cells/}\mu\text{L}$ . |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with one of the following: dermatologist, immunologist, hematologist, geneticist, or allergist.   |
| Coverage<br>Duration                | Initial: 6 months, Reauth: Plan Year  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

# XOSPATA

### **Products Affected**

#### • Xospata

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of acute myeloid leukemia (AML). AML is FMS-like tyrosine kinase 3 (FLT3) mutation-positive. One of the following: a) Used as low-intensity treatment induction when not a candidate for intensive induction therapy, b) Follow-up after induction therapy with response to previous lower intensity therapy with the same regimen, c) Post-allogeneic hematopoietic cell transplantation and in remission, d) Disease is relapsed or refractory, or e) Consolidation therapy as continuation of low-intensity regimen used for induction in patients with poor-risk AML. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## **XPOVIO**

### **Products Affected**

• Xpovio

- Xpovio 60 Mg Twice Weekly Xpovio 80 Mg Twice Weekly

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Multiple Myeloma (MM): Diagnosis of multiple myeloma. Patient has received at least one prior therapy (e.g., lenalidomide, bortezomib, daratumumab, pomalidomide). Used in combination with one of the following: bortezomib and dexamethasone, daratumumab and dexamethasone, or carfilzomib and dexamethasone. Relapsed/Refractory Multiple Myeloma (RRMM): Diagnosis of relapsed or refractory multiple myeloma (RRMM). Patient has received at least four prior therapies (e.g., lenalidomide, bortezomib, daratumumab, pomalidomide). Disease is refractory to all of the following: 1) Two proteasome inhibitors (e.g., bortezomib, carfilzomib), 2) Two immunomodulatory agents (e.g., lenalidomide, thalidomide), and 3) An anti-CD38 monoclonal antibody (e.g. daratumumab). Used in combination with dexamethasone. Diffuse large B-cell lymphoma (DLBCL): Diagnosis of one of the following: 1) Relapsed or refractory DLBCL not otherwise specified OR 2) Relapsed or refractory DLBCL arising from follicular lymphoma. Patient has previously received at least two lines of systemic therapy (e.g., CHOP: cyclophosphamide, doxorubicin, vincristine, and prednisone plus rituximab). |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | N/A  |
| Coverage<br>Duration               | Plan Year  |
| Other Criteria                     | Approve for continuation of prior therapy if within the past 120 days.   |

# XTANDI

### **Products Affected**

#### • Xtandi

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Prostate cancer (PC): Diagnosis of prostate cancer. One of the following:  1) Disease is castration-resistant, 2) Disease is both of the following: a)  Metastatic, and b) Castration-sensitive, or 3) Disease is all of the following: a) Non-metastatic, b) Castration-sensitive, c) Recurrent, and d)  High risk of metastasis. One of the following: 1) Used in combination with a gonadotropin-releasing hormone (GnRH) analog [e.g. Lupron (leuprolide), Zoladex (goserelin), Trelstar (triptorelin), Vantas (histrelin), Firmagon (degarelix)] or 2) Patient has had bilateral orchiectomy. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## YESINTEK

### **Products Affected**

• Yesintek INJ 45MG/0.5ML, 90MG/ML

| PA Criteria                        | Criteria Details   |
|------------------------------------|--|
| Indications                        | All Medically-accepted Indications.  |
| Off-Label Uses                     | N/A  |
| Exclusion<br>Criteria              | N/A  |
| Required<br>Medical<br>Information | Plaque psoriasis (Initial - 45mg/0.5mL): Diagnosis of moderate to severe plaque psoriasis. Plaque psoriasis (Initial - 90mg/1mL): Diagnosis of moderate to severe plaque psoriasis. Patient's weight is greater than 100 kg (220 lbs). Plaque psoriasis (Initial regardless of dose): One of the following: at least 3% body surface area involvement, severe scalp psoriasis, OR palmoplantar (ie, palms, soles), facial, or genital involvement. Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, or calcineurin inhibitors (eg, tacrolimus, pimecrolimus). Psoriatic arthritis (PsA) (Initial - 45mg/0.5mL): Diagnosis of active PsA. PsA (Initial - 90mg/1mL): Diagnosis of active PsA. Patient's weight is greater than 100 kg (220 lbs). Diagnosis of co-existent moderate to severe psoriasis. PsA (Initial regardless of dose): One of the following: actively inflamed joints, dactylitis, enthesitis, axial disease, or active skin and/or nail involvement. Crohn's disease (CD) (initial): Diagnosis of moderately to severely active Crohn's disease. Will be used as a maintenance dose following the intravenous induction dose. |
| Age Restrictions                   | N/A  |
| Prescriber<br>Restrictions         | Plaque psoriasis (Initial): Prescribed by or in consultation with a dermatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist. CD and UC (initial): Prescribed by or in consultation with a gastroenterologist.  |
| Coverage<br>Duration               | All indications (initial, reauth): plan year.  |

| Other Criteria                      | Ulcerative colitis (UC) (initial): Diagnosis of moderately to severely active UC. Will be used as a maintenance dose following the intravenous induction dose. All indications (Reauth): Patient demonstrates positive clinical response to therapy. |
|-------------------------------------|--|
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.   |

## ZEJULA

### **Products Affected**

### • Zejula TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Ovarian cancer: Diagnosis of one of the following: epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer. One of the following: A) Used for maintenance treatment in patients who are in a complete or partial response to platinum-based chemotherapy (e.g., cisplatin, carboplatin), or B) Used for recurrence therapy for platinum-sensitive disease in combination with bevacizumab. Disease is stage II-IV. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **Z**ELBORAF

### **Products Affected**

#### • Zelboraf

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Melanoma: Diagnosis of unresectable or metastatic melanoma. Patient is positive for BRAF V600 mutation. Erdheim-Chester Disease: Diagnosis of Erdheim-Chester disease AND Disease is BRAFV600 mutant type (MT). |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

## ZOLINZA

#### **Products Affected**

#### • Zolinza

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Cutaneous T-cell lymphoma (CTCL): Diagnosis of CTCL. Patient has progressive, persistent, or recurrent disease on or following two systemic therapies [e.g., Adcetris (brentuximab vedotin), bexarotene, interferon gamma-1b, methotrexate, Poteligeo (mogamulizumab), romidepsin]. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **Z**ORYVE

### **Products Affected**

### • Zoryve CREA

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Zoryve 0.3% Plaque psoriasis (PsO) (Initial): Diagnosis of plaque psoriasis. One of the following: a) Minimum duration of a 4 week trial and failure, contraindication, or intolerance (TF/C/I) to one of the following topical therapies: corticosteroids (eg, betamethasone, clobetasol), vitamin D analogs (eg, calcitriol, calcipotriene), tazarotene, calcineurin inhibitors (eg, tacrolimus, pimecrolimus), or a topical combination therapy [e.g., vitamin D analog/corticosteroid (eg, Enstilar, Taclonex), or Duobrii (halobetasol/tazarotene)]. Zoryve 0.15% Atopic dermatitis (AD) (Initial): Diagnosis of mild to moderate AD. One of the following: a) Greater than or equal to 3% body surface area (BSA) involvement, or b) Involvement of sensitive body areas (e.g., face, hands, feet, scalp, groin). Trial and failure of a minimum 30-day supply (14-day supply for topical corticosteroids), contraindication, or intolerance to both of the following: a) Medium or higher potency topical corticosteroid, b) Elidel (pimecrolimus) cream or Tacrolimus ointment. |
| Age Restrictions                    | AD (initial): Patient is 6 years of age or older.   |
| Prescriber<br>Restrictions          | PsO (Initial): Prescribed by or in consultation with a dermatologist. AD (Initial): Prescribed by or in consultation with a dermatologist or allergist/immunologist.  |
| Coverage<br>Duration                | PsO, AD (Initial): 6 months, Reauth: Plan Year  |
| Other Criteria                      | PsO, AD (Reauth): Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

# **ZORYVE FOAM**

#### **Products Affected**

### • Zoryve FOAM

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Initial: Diagnosis of seborrheic dermatitis. Minimum duration of a 4-week trial and failure, contraindication, or intolerance to one of the following generic topical therapies: corticosteroids (eg, betamethasone, clobetasol), antifungals (eg, ciclopirox, ketoconazole), or calcineurin inhibitors (eg, tacrolimus). |
| Age Restrictions                    | Initial: Patient is 9 years of age or older.  |
| Prescriber<br>Restrictions          | Initial: Prescribed by or in consultation with a dermatologist.   |
| Coverage<br>Duration                | Initial: 6 months, Reauth: Plan Year  |
| Other Criteria                      | Reauth: Patient demonstrates positive clinical response to therapy.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES require use of a prerequisite Part D drug.  |

## **Z**TALMY

### **Products Affected**

#### • Ztalmy

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Diagnosis of cyclin-dependent kinase-like 5 (CDKL5) deficiency disorder (CDD). Patient has a mutation in the CDKL5 gene. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | Prescribed by or in consultation with a neurologist.   |
| Coverage<br>Duration                | Plan Year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

# **Z**URZUVAE

#### **Products Affected**

#### • Zurzuvae

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--------------------|
| Indications                         | Pending CMS Review |
| Off-Label Uses                      | Pending CMS Review |
| Exclusion<br>Criteria               | Pending CMS Review |
| Required<br>Medical<br>Information  | Pending CMS Review |
| Age Restrictions                    | Pending CMS Review |
| Prescriber<br>Restrictions          | Pending CMS Review |
| Coverage<br>Duration                | Pending CMS Review |
| Other Criteria                      | Pending CMS Review |
| Prerequisite<br>Therapy<br>Required | Pending CMS Review |

## ZYDELIG

### **Products Affected**

### • Zydelig

| PA Criteria                         | Criteria Details   |
|-------------------------------------|--|
| Indications                         | All Medically-accepted Indications.  |
| Off-Label Uses                      | N/A  |
| Exclusion<br>Criteria               | N/A  |
| Required<br>Medical<br>Information  | Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL. Disease has relapsed or is refractory. |
| Age Restrictions                    | N/A  |
| Prescriber<br>Restrictions          | N/A  |
| Coverage<br>Duration                | Plan year  |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.   |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.   |

## ZYKADIA

### **Products Affected**

### • Zykadia TABS

| PA Criteria                         | Criteria Details  |
|-------------------------------------|---|
| Indications                         | All Medically-accepted Indications.   |
| Off-Label Uses                      | N/A   |
| Exclusion<br>Criteria               | N/A   |
| Required<br>Medical<br>Information  | Diagnosis of metastatic, recurrent, or advanced non-small cell lung cancer (NSCLC), tumor is anaplastic lymphoma kinase (ALK)-positive. |
| Age Restrictions                    | N/A   |
| Prescriber<br>Restrictions          | N/A   |
| Coverage<br>Duration                | Plan year   |
| Other Criteria                      | Approve for continuation of prior therapy if within the past 120 days.  |
| Prerequisite<br>Therapy<br>Required | Criteria DOES NOT require use of a prerequisite Part D drug.  |

#### PART B VERSUS PART D

#### **Products Affected**

- Acetylcysteine INHALATION SOLN
- Acyclovir Sodium INJ 50MG/ML
- Albuterol Sulfate NEBU 0.083%, 0.63MG/3ML, 1.25MG/3ML, 2.5MG/0.5ML
- Amphotericin B INJ
- Amphotericin B Liposome
- Arformoterol Tartrate
- Astagraf XL
- Azathioprine TABS 50MG
- Budesonide SUSP
- Calcitriol CAPS
- Calcitriol ORAL SOLN
- Cinacalcet Hydrochloride
- Clinolipid
- Cromolyn Sodium NEBU
- Cyclophosphamide CAPS
- Cyclophosphamide TABS
- Cyclosporine CAPS
- Cyclosporine Modified
- Doxercalciferol CAPS
- Engerix-b
- Everolimus TABS 0.25MG, 0.5MG, 0.75MG, 1MG
- Gengraf CAPS 100MG, 25MG
- Granisetron Hydrochloride TABS
- Heparin Sodium INJ 10000UNIT/ML, 1000UNIT/ML, 20000UNIT/ML, 5000UNIT/ML
- Heplisav-b
- Imovax Rabies (h.d.c.v.)
- Intralipid INJ 20GM/100ML, 30GM/100ML
- Ipratropium Bromide INHALATION SOLN 0.02%
- Ipratropium Bromide/albuterol Sulfate
- Levalbuterol NEBU
- Levalbuterol Hcl NEBU 0.31MG/3ML, 1.25MG/3ML

- Levalbuterol Hydrochloride NEBU 0.63MG/3ML
- Mycophenolate Mofetil CAPS
- Mycophenolate Mofetil SUSR
- Mycophenolate Mofetil TABS
- Mycophenolic Acid Dr
- Nutrilipid
- Ondansetron Hcl SOLN
- Ondansetron Hydrochloride TABS
- Ondansetron Odt TBDP 4MG, 8MG
- Paricalcitol CAPS
- Pentamidine Isethionate INHALATION SOLR
- Plenamine INJ 147.4MEQ/L;
  2.17GM/100ML; 1.47GM/100ML;
  434MG/100ML; 749MG/100ML;
  1.04GM/100ML; 894MG/100ML;
  749MG/100ML; 1.04GM/100ML;
  1.18GM/100ML; 749MG/100ML;
  1.04GM/100ML; 894MG/100ML;
  592MG/100ML; 749MG/100ML;
  250MG/100ML; 39MG/100ML;
  960MG/100ML
- Premasol INJ 52MEQ/L; 1760MG/100ML; 880MG/100ML; 34MEQ/L; 1760MG/100ML; 372MG/100ML; 406MG/100ML; 526MG/100ML; 492MG/100ML; 492MG/100ML; 526MG/100ML; 356MG/100ML; 356MG/100ML; 390MG/100ML; 34MG/100ML; 152MG/100ML
- Prograf PACK
- Prosol
- Pulmozyme SOLN 2.5MG/2.5ML
- Rabavert
- Recombivax Hb
- Sirolimus SOLN
- Sirolimus TABS
- Tacrolimus CAPS
- Tobramycin NEBU 300MG/5ML

```
Travasol INJ 52MEQ/L;
1760MG/100ML; 880MG/100ML;
34MEQ/L; 1760MG/100ML;
372MG/100ML; 406MG/100ML;
526MG/100ML; 492MG/100ML;
492MG/100ML; 526MG/100ML;
356MG/100ML; 500MG/100ML;
356MG/100ML; 390MG/100ML;
34MG/100ML; 152MG/100ML
```

Trophamine INJ 0.54GM/100ML;
1.2GM/100ML; 0.32GM/100ML; 0; 0;
0.5GM/100ML; 0.36GM/100ML;
0.48GM/100ML; 0.82GM/100ML;
1.4GM/100ML; 1.2GM/100ML;
0.34GM/100ML; 0.48GM/100ML;
0.68GM/100ML; 0.38GM/100ML;
5MEQ/L; 0.025GM/100ML;
0.42GM/100ML; 0.2GM/100ML;
0.24GM/100ML; 0.78GM/100ML

#### **Details**

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.

Plans are insured through UnitedHealthcare Insurance Company or one of its affiliated companies, a Medicare Advantage organization with a Medicare contract and a Medicare-approved Part D sponsor. Enrollment in the plan depends on the plan's contract renewal with Medicare.

[<OVEX3386715\_000>] Formulary ID# 00026002 Y0066 130404 093713 CMS Approved