ACTIMMUNE (S)

Products Affected

- Actimmune INJ 100MCG/0.5ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: 1) Chronic granulomatous disease (CGD), or 2) severe malignant osteopetrosis (SMO).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ADALIMUMAB-AATY (S)

Products Affected

- Adalimumab-aaty 1-pen Kit
- Adalimumab-aaty 2-pen Kit
- Adalimumab-aaty 2-syringe
- Adalimumab-aaty Cd/uc/hs Starter

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

ADALIMUMAB-ADBM (S)

Products Affected

- Adalimumab-adbm

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

ADALIMUMAB-FKJP (S)

Products Affected

- Adalimumab-fkjp

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

ADCIRCA (S)

Products Affected

- Alyq
- Tadalafil TABS 20MG

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

ADEMPAS (S)

Products Affected

- Adempas

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

AFINITOR (S)

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 Criteria	N/A
Required Medical Information	Subependymal Giant Cell Astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC): Diagnosis of SEGA associated with TSC that requires therapeutic intervention. Renal cell carcinoma: Diagnosis of advanced renal cell carcinoma AND trial and failure, contraindication, or intolerance to SUTENT (sunitinib) or NEXAVAR (sorafenib). Neuroendocrine tumors of pancreatic origin (pNET): Diagnosis of progressive pNET that are unresectable, locally advanced, or metastatic. Renal angiomyolipoma: Diagnosis of renal angiomyolipoma and TSC. Breast Cancer: Diagnosis of advanced hormone receptor-positive, HER2-negative breast cancer AND trial and failure, contraindication, or intolerance to FEMARA (letrozole) or ARIMIDEX (anastrozole). Neuroendocrine tumors of gastrointestinal (GI) or lung origin: Diagnosis of progressive, well-differentiated, non-functional NET of GI or lung origin AND patient has unresectable, locally advanced or metastatic disease.
Age Restrictions	SEGA associated with TSC: Patient is 1 year of age or older.
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	All Indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AFINITOR DISPERZ (S)

Products Affected

- Everolimus TBSO

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Subependymal Giant Cell Astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC): Diagnosis of SEGA associated with TSC that requires therapeutic intervention. TSC-associated partial-onset seizures: Diagnosis of TSC-associated partial-onset seizures.
Age Restrictions	SEGA associated with TSC: Patient is 1 year of age or older. TSC-associated partial-onset seizures: Patient is 2 years of age or older.
Prescriber Restrictions	TSC-associated partial-onset seizures: Prescribed by or in consultation with a neurologist.
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AIMOVIG (S)

Products Affected

- Aimovig

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

AKEEGA (S)

Products Affected

- Akeega

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of prostate cancer. Disease is both of the following: a) metastatic and b) deleterious or suspected deleterious BRCA-mutated (BRCAm). Used in combination with prednisone. One of the following: a) Used in combination with a gonadotropin-releasing hormone (GnRH) analog, or b) Patient has had a bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ALECENSA (S)

Products Affected

- Alecensa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALPHA-1 PROTEINASE INHIBITOR, PROLASTIN (S)

Products Affected

- Prolastin-c INJ 1000MG/20ML

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

ALUNBRIG (S)

Products Affected

- Alunbrig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AMPYRA (S)

Products Affected

- Dalfampridine Er

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple Sclerosis (MS) (initial): Diagnosis of MS. Physician confirmation that patient has difficulty walking (eg, timed 25 foot walk test). One of the following: expanded disability status scale (EDSS) score less than or equal to 7, or not restricted to using a wheelchair (if EDSS is not measured).
Age Restrictions	N/A
Prescriber Restrictions	MS (initial): Prescribed by or in consultation with a neurologist.
Coverage Duration	MS (Initial): 6 months. (Reauth): 12 months.
Other Criteria	MS (Reauth): Physician confirmation that the patient's walking improved with therapy. One of the following: EDSS score less than or equal to 7, or not restricted to using a wheelchair (if EDSS is not measured).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ARFORMOTEROL (S)

Products Affected

- Arformoterol Tartrate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic Obstructive Pulmonary Disease (COPD): Diagnosis of COPD. Used for maintenance treatment of bronchoconstriction in patients with COPD, including chronic bronchitis and emphysema.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	COPD: 12 months.
Other Criteria	Subject to Part B vs. Part D review.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ARIKAYCE (S)

Products Affected

- Arikayce

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	Prescribed by or in consultation with an infectious disease specialist or pulmonologist.
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AUGTYRO (S)

Products Affected

- Augtyro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: a) locally advanced, or b) metastatic. Disease is confirmed by the presence of ROS1 rearrangement positive tumor(s) as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Solid Tumors: Diagnosis of solid tumors. Disease is confirmed by the presence of neurotrophic tyrosine receptor kinase (NTRK) gene fusion (e.g., ETV6-NTRK3, TPM3-NTRK1, LMNA-NTRK1) as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). 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 previous treatment (e.g., radiation therapy, systemic therapy, tyrosine kinase inhibitor [TKI]), or b) Disease has no satisfactory alternative treatments.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All indications: 12 months.
Other Criteria	All indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AUSTEDO (S)

Products Affected

- Austedo
- Austedo Xr
- Austedo Xr Patient Titration Kit

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

AVMAPKI FAKZYNJA (S)

Products Affected

- Avmapki Fakzynja Co-pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of recurrent low-grade serous ovarian cancer (LGSOC). Presence of a KRAS-mutation as detected by a Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received prior systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AVTOZMA IV (S)

Products Affected

- Avtozma INJ 200MG/10ML, 400MG/20ML, 80MG/4ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Rheumatoid Arthritis (RA) (Initial): Diagnosis of moderately to severely active RA. One of the following: a) Trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: Enbrel (etanercept), one formulary adalimumab product, Rinvoq (upadacitinib), or tofacitinib, or b) attestation demonstrating a trial may be inappropriate, OR c) For continuation of prior therapy. Systemic Juvenile Idiopathic arthritis (SJIA) (Initial): Diagnosis of active SJIA. TF/C/I to one of the following conventional therapies at maximally tolerated doses: minimum duration of a one month trial of a nonsteroidal anti-inflammatory drug (NSAID) (eg, ibuprofen, naproxen), or 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: a) TF/C/I to two of the following, or attestation demonstrating a trial may be inappropriate: Enbrel (etanercept), one formulary adalimumab product, Rinvoq/LQ, or tofacitinib, OR b) for continuation of prior therapy. Giant Cell Arteritis (GCA) (Initial): Diagnosis of GCA. Cytokine Release Syndrome (CRS): Patient will receive or is receiving chimeric antigen receptor (CAR) T-cell immunotherapy [e.g., Kymriah (tisagenlecleuce), Yescarta (axicabtagene ciloleucel)]. Coronavirus disease 2019 (COVID-19): Diagnosis of COVID-19. Patient is hospitalized. Currently receiving systemic corticosteroids. Patient requires one of the following: supplemental oxygen, non-invasive mechanical ventilation, invasive mechanical ventilation, or extracorporeal membrane oxygenation (ECMO).
Age Restrictions	N/A
Prescriber Restrictions	RA, SJIA, PJIA, GCA (initial): Prescribed by or in consultation with a rheumatologist. CRS: Prescribed by or in consultation with an oncologist or hematologist.

Coverage Duration	RA, SJIA, PJIA, GCA (Init): 6 mo, (reauth): 12 mo. CRS: 2 mo. COVID-19: 14 days.
Other Criteria	RA, PJIA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline OR improvement in symptoms (eg, pain, stiffness, inflammation) from baseline. SJIA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline OR improvement in clinical features or symptoms (eg, pain, fever, inflammation, rash, lymphadenopathy, serositis) from baseline. GCA (Reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AVTOZMA SC (S)

Products Affected

- Avtozma INJ 162MG/0.9ML

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

AYVAKIT (S)

Products Affected

- Ayvakit

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Gastrointestinal stromal tumor (GIST): Diagnosis of GIST. Disease is one of the following: unresectable or metastatic. Presence of platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation, including PDGFRA D842V mutations, as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Advanced Systemic Mastocytosis (AdvSM): Diagnosis of AdvSM. Patient has one of the following: a) aggressive systemic mastocytosis (ASM), b) systemic mastocytosis with an associated hematological neoplasm (SM-AHN), or c) mast cell leukemia (MCL). Platelet count is greater than or equal to 50 x 10^9/L. Ayvakit 25 mg - Indolent Systemic Mastocytosis (ISM): Diagnosis of ISM. Platelet count is greater than or equal to 50 x 10^9/L.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months.
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BALVERSA (S)

Products Affected

- Balversa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Urothelial Carcinoma: Diagnosis of urothelial carcinoma (UC). Disease is one of the following: Locally advanced or Metastatic. Presence of susceptible fibroblast growth factor receptor (FGFR) 3 genetic alterations as detected by an U.S. Food and Drug Administration (FDA)-approved test (therascreen FGFR RGQ RT-PCR Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease has progressed on or after at least one line of prior systemic therapy (e.g., chemotherapy). One of the following: 1) Patient had been treated with prior PD-1 inhibitor (e.g., Opdivo [nivolumab], Keytruda [pembrolizumab]) or PD-L1 inhibitor therapy (e.g., Bavencio [avelumab]) or 2) Patient is not a candidate for PD-1 or PD-L1 inhibitor therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BENLYSTA (S)

Products Affected

- Benlysta INJ 200MG/ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Systemic lupus erythematosus (SLE) (init): 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 (init): 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).
Age Restrictions	SLE, Lupus Nephritis (init): Benlysta IV (vial), SC (Autoinjector): Patient is 5 years of age or older. SC (Prefilled Syringe): Patient is 18 years of age or older.
Prescriber Restrictions	SLE (init): Prescribed by or in consultation with a rheumatologist. Lupus Nephritis (init): Prescribed by or in consultation with a nephrologist or rheumatologist.
Coverage Duration	SLE, Lupus Nephritis (init, reauth): 6 months
Other Criteria	SLE, Lupus Nephritis (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BEQALZI (S)

Products Affected

- Beqalzi
- Beqalzi Starter Pack

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

BESREMI (S)

Products Affected

- Besremi
- Besremi Pen

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

BOSULIF (S)

Products Affected

- Bosulif CAPS
- Bosulif TABS 100MG, 500MG
- Bosutinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic myelogenous/myeloid leukemia (CML): Diagnosis of Philadelphia chromosome-positive (Ph+) CML.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BRAFTOVI (S)

Products Affected

- Braftovi CAPS 75MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma. Presence of BRAF V600E or V600K mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). 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. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Both of the following a) Used in combination with Erbitux (cetuximab) AND b) One of the following: i) Patient has received prior therapy OR ii) Used in combination with fluorouracil-based chemotherapy (e.g., mFOLFOX6, FOLFIRI). Non-Small Cell Lung Cancer (NSCLC): Diagnosis of metastatic NSCLC. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with Mektovi (binimetinib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BRIVIACT (S)

Products Affected

- Brivaracetam SOLN
- Brivaracetam TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Partial-onset seizures: Diagnosis of partial-onset seizures.
Age Restrictions	Patient is 1 month of age or older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BRONCHITOL (S)

Products Affected

- Bronchitol

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of cystic fibrosis (CF). Patient has passed the Bronchitol Tolerance Test (BTT).
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with a pulmonologist or specialist affiliated with a CF care center.
Coverage Duration	Initial: 6 months. Reauth: 12 months.
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BRUKINSA (S)

Products Affected

- Brukinsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Mantle Cell Lymphoma (MCL): Diagnosis of MCL. Paid claims or submission of medical records confirming patient has received at least one prior therapy for MCL [e.g., Calquence (acalabrutinib)]. Waldenstrom's Macroglobulinemia (WM)/Lymphoplasmacytic Lymphoma (LPL): Diagnosis of WM/LPL. Marginal Zone Lymphoma (MZL): Diagnosis of MZL. Disease is relapsed or refractory. Paid claims or submission of medical records confirming patient has received at least one prior anti-CD20-based regimen for MZL (e.g., rituximab, obinutuzumab). Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL): Diagnosis of ONE of the following: CLL or SLL. Submission of medical records confirming patient has a contraindication or intolerance to Calquence (acalabrutinib). Follicular Lymphoma (FL): Diagnosis of FL. Disease is relapsed or refractory. Used in combination with Gazyva (obinutuzumab). Paid claims or submission of medical records confirming patient has received at least two prior lines of systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months.
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BYLVAY (S)

Products Affected

- Bylvay
- Bylvay (pellets)

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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. Alagille syndrome (ALGS) (initial): Both of the following: 1) Diagnosis of ALGS, and 2) Molecular genetic testing confirms mutations in the JAG1 or NOTCH2 gene.
Age Restrictions	PFIC (initial): Patient is 3 months of age or older. ALGS (initial): Patient is 12 months of age or older.
Prescriber Restrictions	PFIC, ALGS (initial): Prescribed by or in consultation with a hepatologist or gastroenterologist
Coverage Duration	PFIC, ALGS (initial): 12 months. PFIC, ALGS (reauth): 12 months.
Other Criteria	PFIC (reauth): Patient demonstrates positive clinical response to therapy. ALGS (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CABLIVI (S)

Products Affected

- Cablivi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	Prescribed by or in consultation with a hematologist or oncologist.
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CABOMETYX (S)

Products Affected

- Cabometyx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Renal cell carcinoma (RCC): Diagnosis of RCC. Hepatocellular Carcinoma (HCC): Diagnosis of HCC. Trial and failure, contraindication, or intolerance to Nexavar (sorafenib tosylate). Differentiated Thyroid Cancer (DTC): Diagnosis of DTC. Disease has progressed following prior VEGFR-targeted therapy (e.g., Lenvima [lenvatinib], Nexavar [sorafenib]). Disease or patient is refractory to radioactive iodine treatment or ineligible. Neuroendocrine Tumors: Diagnosis of one of the following: 1) Pancreatic neuroendocrine tumors (pNET) or 2) Extra-pancreatic neuroendocrine tumors (epNET). Disease is one of the following: 1) Unresectable, 2) Locally advanced, or 3) Metastatic. Tumors are well-differentiated. Patient has been previously treated (e.g., octreotide, lanreotide, chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CALQUENCE (S)

Products Affected

- Calquence TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Mantle Cell Lymphoma: Diagnosis of mantle cell lymphoma (MCL). One of the following: A) All of the following: 1) Patient has received no prior therapy for MCL (e.g., bortezomib, rituximab), 2) Patient is ineligible for autologous hematopoietic stem cell transplantation (HSCT), and 3) Used in combination with bendamustine and rituximab, OR B) Patient has received at least one prior therapy for MCL (e.g., bortezomib, rituximab). Chronic Lymphocytic Leukemia (CLL) or Small Lymphocytic Lymphoma (SLL): Diagnosis of CLL or SLL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CAPLYTA (S)

Products Affected

- Caplyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Schizophrenia: Diagnosis of schizophrenia. Trial and failure, contraindication, or intolerance to two of the following oral generic formulary atypical antipsychotic agents: asenapine, aripiprazole, olanzapine, paliperidone, quetiapine (IR or ER), risperidone, ziprasidone. Bipolar disorder: Diagnosis of bipolar I or II disorder (bipolar depression). Patient has depressive episodes associated with bipolar disorder. Used as monotherapy or as adjunctive therapy with lithium or valproate. Trial and failure, contraindication, or intolerance to quetiapine (IR or ER) or olanzapine. Major depressive disorder (MDD): Diagnosis of MDD. Used as adjunctive therapy with an antidepressant (e.g., escitalopram, sertraline, venlafaxine/desvenlafaxine, etc.). Trial and failure, contraindication, or intolerance to both of the following oral generic formulary atypical antipsychotic agents: aripiprazole and quetiapine IR/ER.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CAPRELSA (S)

Products Affected

- Caprelsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Thyroid Cancer: Diagnosis of one of the following: a) metastatic medullary thyroid cancer (MTC) or b) unresectable locally advanced MTC. Patient has symptomatic disease or progressive disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CAVHANZA (S)

Products Affected

- Cavhanza

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

CAYSTON (S)

Products Affected

- Cayston

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cystic fibrosis (CF) (Initial, Reauth): Diagnosis of CF. Submission of medical records (e.g., chart notes, laboratory values) confirming Pseudomonas aeruginosa in the lungs.
Age Restrictions	CF (Initial): 7 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	CF (Initial, reauth): 12 months
Other Criteria	CF (Reauth): Patient is benefiting from treatment (i.e. improvement in lung function [forced expiratory volume in one second (FEV1)], decreased number of pulmonary exacerbations).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CHOLBAM (S)

Products Affected

- Cholbam

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 based on one of the following: a) an abnormal urinary bile acid analysis by mass spectrometry OR b) molecular genetic testing consistent with the diagnosis. Peroxisomal disorders (PD) (initial): Diagnosis of a peroxisomal disorder, including Zellweger spectrum disorders, based on one of the following: a) an abnormal urinary bile acid analysis by mass spectrometry, or b) molecular genetic testing consistent with the diagnosis. Patient exhibits at least one of the following: a) liver disease (eg, jaundice, elevated serum transaminases), b) steatorrhea, or c) complications from decreased fat-soluble vitamin absorption (eg, poor growth). Will be used as an adjunctive treatment.
Age Restrictions	N/A
Prescriber Restrictions	All uses (initial): Prescribed by a hepatologist, medical geneticist, gastroenterologist, or other specialist that treats inborn errors of metabolism.
Coverage Duration	All uses (initial): 4 months, (reauth) 12 months.
Other Criteria	All uses (reauth): Patient demonstrates positive clinical response to therapy as evidenced by improvement in liver function.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CIALIS (S)

Products Affected

- Tadalafil TABS 2.5MG, 5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Concurrent use of nitrates.
Required Medical Information	Diagnosis of benign prostatic hyperplasia (BPH). Trial and failure, contraindication, or intolerance to an alpha-blocker (e.g., doxazosin, prazosin, tamsulosin) or a 5-alpha reductase inhibitor (e.g., dutasteride, finasteride).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CICLOPIROX (S)

Products Affected

- Ciclodan SOLN
- Ciclopirox Nail Lacquer

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	All of the following: 1) Patient does not have lunula (matrix) involvement, 2) one of the following: a) Diagnosis of onychomycosis of the toenails, OR b) Diagnosis of onychomycosis of the fingernails, 3) Diagnosis of onychomycosis has been confirmed by one of the following: a) positive potassium hydroxide (KOH) preparation, OR b) culture, OR c) histology, 4) If toenail onychomycosis, patient has mild to moderate disease involving at least 1 target toenail, AND 5) One of the following: a) For onychomycosis of fingernails, Trial and failure (of a minimum 6-week supply), contraindication, or intolerance to oral terbinafine, or b) For onychomycosis of toenails, trial and failure (of a minimum 12-week supply), contraindication, or intolerance to oral terbinafine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	48 weeks.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CINRYZE (S)

Products Affected

- Cinryze

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

COBENFY (S)

Products Affected

- Cobenfy
- Cobenfy Starter Pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of schizophrenia. Trial and failure, contraindication, or intolerance to two of the following oral generic formulary atypical antipsychotic agents: a) aripiprazole, b) asenapine, c) olanzapine, d) paliperidone, e) quetiapine (IR or ER), f) risperidone, or g) ziprasidone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COMETRIQ (S)

Products Affected

- Cometriq

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Medullary thyroid cancer (MTC): Diagnosis of Metastatic MTC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

COPIKTRA (S)

Products Affected

- Copiktra

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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], Rituxan [rituximab], etc.).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COSENTYX (S)

Products Affected

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

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

COSENTYX IV (S)

Products Affected

- Cosentyx INJ 125MG/5ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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. One of the following: a) Trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: Cosentyx SC (secukinumab), Enbrel (etanercept), one formulary adalimumab product, Otezla (apremilast), Skyrizi (risankizumab-rzaa), Sotyktu (deucravacitinib), Tremfya (guselkumab), one formulary ustekinumab product, Rinvoq/LQ (upadacitinib), or tofacitinib, OR b) for continuation of prior therapy. Ankylosing Spondylitis (AS) (Initial): Diagnosis of active AS. One of the following: a) TF/C/I to two of the following: Cosentyx SC, Enbrel, one formulary adalimumab product, Rinvoq, tofacitinib, OR b) for continuation of prior therapy. Non-radiographic axial spondyloarthritis (nr-axSpA) (Initial): Dx of active nr-axSpA with objective signs of inflammation (eg, C-reactive protein [CRP] levels above the upper limit of normal and/or sacroiliitis on magnetic resonance imaging [MRI], indicative of inflammatory disease, but without definitive radiographic evidence of structural damage on sacroiliac joints). One of the following: a) TF/C/I to both of the following: Cosentyx SC and Rinvoq, OR b) for continuation of prior therapy.
Age Restrictions	N/A
Prescriber Restrictions	PsA (Initial): Prescribed by or in consultation with a rheumatologist or dermatologist. AS, nr-axSpA (Initial): Prescribed by or in consultation with a rheumatologist.
Coverage Duration	All uses (Initial): 6 months. All uses (Reauth): 12 months.

Other Criteria	PsA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline, improvement in symptoms (eg, pain, stiffness, pruritus, inflammation) from baseline, OR reduction in the body surface area (BSA) involvement from baseline. AS, nr-axSpA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by improvement from baseline for at least one of the following: disease activity (eg, pain, fatigue, inflammation, stiffness), lab values (erythrocyte sedimentation rate, C-reactive protein level), function, axial status (eg, lumbar spine motion, chest expansion), OR total active (swollen and tender) joint count.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COTELLIC (S)

Products Affected

- Cotellic

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable or metastatic melanoma. Presence of BRAF V600E or V600K mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with Zelboraf (vemurafenib). Histiocytic Neoplasm: Diagnosis of histiocytic neoplasm. Used as monotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CRESEMBA ORAL (S)

Products Affected

- Cresemba CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Fungal infection: Diagnosis of invasive aspergillosis or invasive mucormycosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CTEXLI (S)

Products Affected

- Ctexli

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of cerebrotendinous xanthomatosis (cholestanol storage disease). Disease is confirmed by the presence of pathogenic variant(s) in the CYP27A1 gene as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with one of the following: a) Neurologist, b) Geneticist, c) Metabolic disease specialist, or d) Endocrinologist.
Coverage Duration	Initial: 12 months. Reauth: 12 months.
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CYCLOBENZAPRINE (S)

Products Affected

- Cyclobenzaprine Hydrochloride TABS 10MG, 5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prescriber acknowledges anticholinergic risks (e.g., confusion, dry mouth, blurry vision, constipation, urinary retention) and will consider lowering the dose or discontinuing medication(s) that are no longer clinically warranted for the patient. Muscle spasm: Diagnosis of muscle spasm associated with acute, painful musculoskeletal conditions. Fibromyalgia (off-label): Used to treat fibromyalgia related symptoms (e.g., difficulty sleeping). Acute temporomandibular disorder (off-label): All of the following: a) Diagnosis of acute temporomandibular disorder, b) Patient has pain on palpitation of the lower jaw muscle, and c) Used in combination with a nonsteroidal anti-inflammatory drug (NSAID) (e.g., naproxen).
Age Restrictions	PA applies to patients 65 years or older.
Prescriber Restrictions	N/A
Coverage Duration	Muscle spasm, temporomandibular disorder: 4 weeks. Fibromyalgia: 12 months.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DALIRESP (S)

Products Affected

- Roflumilast

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	COPD (init, reauth): 12 months
Other Criteria	COPD (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DANZITEN (S)

Products Affected

- Danziten

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Philadelphia chromosome positive chronic myeloid leukemia (Ph+ CML)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DARAPRIM (S)

Products Affected

- Pyrimethamine TABS

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

DAURISMO (S)

Products Affected

- Daurismo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute myeloid leukemia (AML): Diagnosis of newly-diagnosed acute myeloid leukemia (AML) AND Used 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 comorbidities that preclude the use of intensive induction chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DEFERASIROX (S)

Products Affected

- Deferasirox

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic Iron Overload Due to Blood Transfusions (Initial): Diagnosis of chronic iron overload due to blood transfusions (transfusional hemosiderosis). Patient has a baseline ferritin level more than 1,000 mcg/L. Patient has required the transfusion of at least 100 mL/kg packed red blood cells. Myelodysplastic Syndrome (MDS) (Initial): Diagnosis of MDS. Patient has Low or Intermediate-1 disease or is a potential transplant patient. Patient has received more than 20 red blood cell transfusions. Chronic iron overload due to non-transfusion-dependent thalassemia (NTDT) (Initial): Diagnosis of chronic iron overload due to NTDT. Liver iron concentration (LIC) 5 milligrams of iron per gram of liver dry weight (mg Fe/g dw) or higher. Serum ferritin level greater than 300 mcg/L.
Age Restrictions	Iron Overload Due to Blood Transfusions (initial): 2 years of age or older. NTDT (initial): 10 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Iron Overload Due to Blood Transfusions, MDS (initial, reauth):12 mo. NTDT (initial, reauth): 6mo.
Other Criteria	Iron Overload Due to Blood Transfusions, MDS (Reauth): Patient experienced a reduction from baseline in serum ferritin level or LIC. NTDT (Reauth): Patient has LIC 3 mg Fe/g dw or higher. Patient experienced a reduction from baseline in serum ferritin level or LIC.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DEMSER (S)

Products Affected

- Metyrosine

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Preoperative preparation: Diagnosis of pheochromocytoma confirmed by one of the following biochemical tests: a) plasma free metanephrines OR b) urinary fractioned metanephrines. Medication is being used for preoperative preparation. Trial and failure, contraindication, or intolerance to both of the following: a) alpha-adrenergic blocker (e.g., phenoxybenzamine, doxazosin, terazosin) AND b) beta-adrenergic blocker (e.g., propranolol, metoprolol). Treatment (Tx) of pheochromocytoma (initial): Diagnosis of pheochromocytoma confirmed by one of the following biochemical tests: a) plasma free metanephrines OR b) urinary fractioned metanephrines. Patient with hormonally active (catecholamine excess) pheochromocytoma. One of the following: a) patient is not a candidate for surgery OR b) chronic treatment due to malignant pheochromocytoma. Patient has not reached normotension after treatment with a selective alpha-1-adrenergic blocker (e.g., doxazosin, terazosin) and beta-adrenergic blocker (e.g., propranolol, metoprolol). Medication will not be used to control essential hypertension.
Age Restrictions	N/A
Prescriber Restrictions	Preop prep: Prescribed by or in consultation with an endocrinologist OR Endocrine surgeon. Pheochromocytoma tx (initial): Prescribed by or in consultation with endocrinologist OR provider who specializes in the management of pheochromocytoma.
Coverage Duration	Preop prep: 4 wks. Pheochromocytoma tx (initial): 6 months, (reauth): 12 months.
Other Criteria	Pheochromocytoma tx (reauth): Patient demonstrates positive clinical response to therapy (e.g., decreased frequency and severity of hypertensive attacks).

Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DIACOMIT (S)

Products Affected

- Diacomit

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of seizures associated with Dravet syndrome (DS). Used in combination with clobazam. Patient weighs 7kg or more.
Age Restrictions	Patient is 6 months of age or older.
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DOPTELET (S)

Products Affected

- Doptelet

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

DULERA (S)

Products Affected

- Dulera

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Asthma (initial): Diagnosis of asthma. Trial and failure, contraindication (e.g., safety concerns, not indicated for patient's age), or intolerance to Breo Ellipta (fluticasone furoate and vilanterol trifenatate).
Age Restrictions	Initial: Patient is 5 years or older.
Prescriber Restrictions	N/A
Coverage Duration	Initial, reauth: 12 months
Other Criteria	Asthma (reauthorization): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DUPIXENT (S)

Products Affected

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

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

EBGLYSS (S)

Products Affected

- Ebglyss

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

EMGALITY (S)

Products Affected

- Emgality

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

EMPAVELI (S)

Products Affected

- Empaveli

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

ENBREL (S)

Products Affected

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

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

ENDARI (S)

Products Affected

- L-glutamine PACK

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

ENSACOVE (S)

Products Affected

- Ensacove

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-small cell lung cancer (NSCLC). Disease is one of the following: a) Locally advanced or b) Metastatic. Disease is anaplastic lymphoma kinase (ALK) rearrangement-positive as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has not previously received an ALK-inhibitor [e.g., Alecensa (alectinib), Alunbrig (brigatinib), Lorbrena (lorlatinib), Xalkori (crizotinib), Zykadia (ceritinib)].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ENTYVIO SC (S)

Products Affected

- Entyvio Pen

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

EPIDIOLEX (S)

Products Affected

- Epidiolex

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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	LGS, DS, TSC: Patient is 1 year of age or older.
Prescriber Restrictions	LGS, DS, TSC: Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

EPOETIN ALFA (S)

Products Affected

- Procrit

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

ERIVEDGE (S)

Products Affected

- Erivedge

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Basal cell carcinoma: One of the following: A) Diagnosis of metastatic basal cell carcinoma OR B) Both of the following: 1) Diagnosis of locally advanced basal cell carcinoma AND 2) One of the following: a) Disease recurred following surgery or b) Patient is not a candidate for surgery and radiation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ERLEADA (S)

Products Affected

- Erleada

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ESBRIET (S)

Products Affected

- Pirfenidone

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

ETCAMAH (S)

Products Affected

- Etcamah

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

EUCRISA (S)

Products Affected

- Eucrisa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Atopic dermatitis (initial): Diagnosis of mild to moderate atopic dermatitis. Trial and failure, contraindication, or intolerance to one prescription strength topical corticosteroid (e.g., triamcinolone acetonide, fluocinolone acetonide), unless the affected area is sensitive (i.e., face, axillae, groin).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Reauth: Patient demonstrates a positive clinical response to therapy (e.g., reduction in body surface area involvement, reduction in pruritus severity).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

EVRYSDI (S)

Products Affected

- Evrysdi SOLR

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

FABRAZYME (S)

Products Affected

- Fabrazyme

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Fabry Disease (init): Diagnosis of Fabry disease. One of the following: a) detection of pathogenic mutations in the GLA gene by molecular genetic testing, b) deficiency in α-galactosidase A (α-Gal A) enzyme activity in plasma, isolated leukocytes, or dried blood spots (DBS), or c) significant clinical manifestations (e.g., neuropathic pain, cardiomyopathy, renal insufficiency, angiokeratomas, cornea verticillata). Will not be used in combination with other drugs used for Fabry disease.
Age Restrictions	Fabry Disease (init): Patient is 2 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Fabry Disease (init, reauth): 12 months
Other Criteria	Fabry Disease (reauth): Patient demonstrates positive clinical response to therapy. Will not be used in combination with other drugs used for Fabry disease.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FILSPARI (S)

Products Affected

- Filspari

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

FINTEPLA (S)

Products Affected

- Fintepla

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Dravet Syndrome: Diagnosis of seizures associated with Dravet syndrome. Lennox-Gastaut Syndrome: Diagnosis of seizures associated with Lennox-Gastaut syndrome.
Age Restrictions	All Indications: Patient is 2 years of age or older.
Prescriber Restrictions	All Indications: Prescribed by or in consultation with a neurologist.
Coverage Duration	All Indications: 12 months
Other Criteria	All Indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FIRAZYR (S)

Products Affected

- Icatibant Acetate

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

FIRMAGON (S)

Products Affected

- Firmagon INJ 120MG/VIAL, 80MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of advanced or metastatic prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FOTIVDA (S)

Products Affected

- Fotivda

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

FRUZAQLA (S)

Products Affected

- Fruzaqla

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of metastatic colorectal cancer. Patient has been previously treated with both of the following: A) Fluoropyrimidine-, oxaliplatin-, irinotecan-based chemotherapy (e.g., FOLFOX, FOLFIRI, FOLFOXIRI), and B) Anti-VEGF biological therapy (e.g., Avastin [bevacizumab], Zaltrap [ziv-afibercept]). One of the following: A) Presence of RAS mutation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA), OR B) Both of the following: a) Patient has RAS wild-type tumors, AND b) Patient has been previously treated with both of the following: 1) An anti-EGFR biological therapy (e.g., Vectibix [panitumumab], Erbitux [cetuximab]), and 2) One of the following: i) Lonsurf [trifluridine/tipiracil] or ii) Stivarga [regorafenib].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GAMASTAN (S)

Products Affected

- Gamastan

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Immune globulin is being used intramuscularly. The immune globulin will be administered at the minimum effective dose and appropriate frequency for the prescribed diagnosis. Patient requires immunization for hepatitis A, measles, rubella, or varicella.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	14 days
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GAVRETO (S)

Products Affected

- Gavreto

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Presence of metastatic rearranged during transfection (RET) gene fusion-positive tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Thyroid Cancer: Diagnosis of thyroid cancer. Disease is one of the following: advanced or metastatic. Presence of rearranged during transfection (RET) gene fusion-positive tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease requires treatment with systemic therapy. One of the following: patient is radioactive iodine-refractory or radioactive iodine therapy is not appropriate.
Age Restrictions	Thyroid Cancer: Patient is 12 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GILENYA (S)

Products Affected

- Fingolimod Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Multiple Sclerosis (MS) (initial, reauth): Not used in combination with another disease-modifying therapy for MS.
Required Medical Information	MS (initial): Diagnosis of a relapsing form of MS (eg, clinically isolated syndrome, relapsing-remitting disease, secondary progressive disease, including active disease with new brain lesions).
Age Restrictions	N/A
Prescriber Restrictions	MS (initial, reauth): Prescribed by or in consultation with a neurologist
Coverage Duration	MS (initial, reauth): 12 months
Other Criteria	MS (reauth): Patient demonstrates positive clinical response to therapy (e.g., stability in radiologic disease activity, clinical relapses, disease progression).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GILOTRIF (S)

Products Affected

- Gilotrif

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): A) Diagnosis of metastatic NSCLC AND B) One of the following: 1) Both of the following: a) Disease is confirmed by non-resistant epidermal growth factor (EGFR) mutations as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA) AND b) GILOTRIF will be used as first-line treatment, OR 2) All of the following: a) disease progressed after platinum-based chemotherapy (e.g., cisplatin, carboplatin) and b) squamous NSCLC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GLATIRAMER ACETATE (S)

Products Affected

- Glatiramer Acetate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Multiple Sclerosis (MS) (initial, reauth): Not used in combination with another disease-modifying therapy for MS.
Required Medical Information	MS (initial): Diagnosis of a relapsing form of MS (eg, clinically isolated syndrome, relapsing-remitting disease, secondary progressive disease, including active disease with new brain lesions).
Age Restrictions	N/A
Prescriber Restrictions	MS (initial, reauth): Prescribed by or in consultation with a neurologist
Coverage Duration	MS (initial, reauth): 12 months
Other Criteria	MS (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GLEEVEC (S)

Products Affected

- Imatinib Mesylate TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	One of the following: A) Diagnosis of Philadelphia chromosome positive (Ph+)/BCR ABL-positive chronic myelogenous leukemia (CML) OR B) Ph+/BCR ABL+ acute lymphoblastic leukemia (ALL) OR C) Gastrointestinal stromal tumor (GIST) OR D) Dermatofibrosarcoma protuberans that is unresectable, recurrent, or metastatic OR E) Hypereosinophilic syndrome or chronic eosinophilic leukemia OR F) Myelodysplastic syndrome (MDS) or myeloproliferative disease OR G) Aggressive systemic mastocytosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	All uses: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GLYCOPYRROLATE TABLET (S)

Products Affected

- Glycopyrrolate TABS 1MG, 2MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	Initial, Reauth: Prescribed by or in consultation with a gastroenterologist.
Coverage 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 Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GOMEKLI (S)

Products Affected

- Gomekli

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of neurofibromatosis type 1. Patient has plexiform neurofibromas that are both of the following: inoperable and symptomatic.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GROWTH HORMONE, PREFERRED (S)

Products Affected

- Genotropin
- Genotropin Miniquick

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

HERNEXEOS (S)

Products Affected

- Hernexeos

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-squamous non-small cell lung cancer (NSCLC). Disease is one of the following: a) unresectable, b) metastatic. Presence of HER2 (ERBB2) tyrosine kinase domain activating mutations as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HYDROXYZINE ORAL (S)

Products Affected

- Hydroxyzine Hcl TABS 50MG
- Hydroxyzine Hydrochloride SYRP 10MG/5ML
- Hydroxyzine Hydrochloride TABS 10MG, 25MG
- Hydroxyzine Pamoate CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	One of the following: 1) Requested drug is FDA-approved for the condition being treated, or 2) If requested for an off-label indication, the off-label guideline approval criteria have been met. Prescriber acknowledges anticholinergic risks (e.g., confusion, dry mouth, blurry vision, constipation, urinary retention) and will consider lowering the dose or discontinuing medication(s) that are no longer clinically warranted for the patient.
Age Restrictions	Patient is 65 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HYRNUO (S)

Products Affected

- Hyrnuo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-squamous non-small cell lung cancer (NSCLC). Disease is locally advanced or metastatic. Presence of HER2 (ERBB2) tyrosine kinase domain activating mutations as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received prior systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IBRANCE (S)

Products Affected

- Ibrance

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of breast cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IBTROZI (S)

Products Affected

- Ibtrozi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-small cell lung cancer (NSCLC). Disease is one of the following: 1) Locally advanced or 2) Metastatic. Presence of ROS1 rearrangement-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ICLUSIG (S)

Products Affected

- Iclusig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic myelogenous leukemia: Diagnosis of chronic myelogenous leukemia. Acute Lymphoblastic Leukemia: Diagnosis of Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL). One of the following: a) Used in combination with chemotherapy up to 20 cycles OR b) Used as monotherapy in patients where one of the following applies: i) No other kinase inhibitors are indicated OR ii) Disease is T315I-positive Ph+ ALL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	All uses: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IDHIFA (S)

Products Affected

- Idhifa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute Myeloid Leukemia (AML): Diagnosis of AML. Disease is relapsed or refractory. Patient has an isocitrate dehydrogenase-2 (IDH2) mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test (e.g., Abbott RealTime IDH2 assay) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IGALMI (S)

Products Affected

- Igalmi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	One of the following diagnoses: Schizophrenia or Bipolar I or II disorder. For the treatment of acute agitation. Trial and failure, contraindication or intolerance to at least two products used in acute agitation (e.g., olanzapine, ziprasidone). Patient is currently being managed with maintenance medication for their underlying disorder (e.g., aripiprazole, olanzapine, quetiapine, lithium, valproic acid).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	14 days
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMBRUVICA (S)

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 Criteria	N/A
Required Medical Information	Chronic lymphocytic leukemia (CLL): Diagnosis of CLL. Waldenstrom's macroglobulinemia: Diagnosis of Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma. Small lymphocytic lymphoma (SLL): Diagnosis of SLL. Chronic graft versus host disease (cGVHD): Diagnosis of cGVHD AND trial and failure of one or more lines of systemic therapy (e.g., corticosteroids like prednisone or methylprednisolone, mycophenolate).
Age Restrictions	(cGVHD): Patient is 1 year of age or older.
Prescriber Restrictions	N/A
Coverage Duration	All Uses: 12 months
Other Criteria	All Uses: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMKELDI (S)

Products Affected

- Imkeldi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic Myelogenous/Myeloid Leukemia (CML): Diagnosis of Philadelphia chromosome/BCR ABL-positive (Ph+/BCR ABL+) CML. Acute Lymphoblastic Leukemia/Acute Lymphoblastic Lymphoma (ALL): Diagnosis of Ph+/BCR ABL+ ALL. Myelodysplastic Disease (MDS)/Myeloproliferative Disease (MPD): Diagnosis of MDS/MPD. Aggressive Systemic Mastocytosis (ASM): Diagnosis of ASM. 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. All indications: Patient is unable to swallow generic imatinib tablet due to one of the following: a) Age, b) Physical impairment (e.g., difficulties with motor or oral coordination), c) Dysphagia, or d) Patient is using a feeding tube or nasal gastric tube.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INBRIJA (S)

Products Affected

- Inbrija

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

INCRELEX (S)

Products Affected

- Increlex

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Insulin-like Growth Factor-1 (IGF-1) deficiency (initial): 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 (initial): Diagnosis of GH gene deletion in patients who have developed neutralizing antibodies to GH.
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with an endocrinologist
Coverage Duration	Initial, reauth: 12 months
Other Criteria	(Reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INLURIYO (S)

Products Affected

- Inluriyo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of breast cancer. Disease is advanced or metastatic. Disease is estrogen receptor (ER)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Presence of estrogen receptor-1 (ESR1) mutation(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease has progressed following at least one line of endocrine therapy [e.g., Faslodex (fulvestrant), Arimidex (anastrozole), Femara (letrozole), Aromasin (exemestane)].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INLYTA (S)

Products Affected

- Inlyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Renal cell carcinoma (RCC): Diagnosis of RCC. One of the following: (1) used as first-line treatment in combination with avelumab or pembrolizumab or (2) used after failure of one prior systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INQOVI (S)

Products Affected

- Inqovi

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

INREBIC (S)

Products Affected

- Inrebic

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Myelofibrosis: Diagnosis of one of the following: primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IRESSA (S)

Products Affected

- Gefitinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of metastatic NSCLC AND Disease is confirmed by known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutations as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ISTURISA (S)

Products Affected

- Isturisa TABS 1MG, 5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cushing's Syndrome (initial): Diagnosis of Cushing's syndrome. Used for treatment of endogenous hypercortisolemia. One of the following: a) Patient is not a candidate for surgery (e.g., adrenalectomy, transsphenoidal surgery), OR b) Surgery has not been curative for the patient.
Age Restrictions	N/A
Prescriber Restrictions	Cushing's Syndrome (initial): Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Cushing's Syndrome (initial, reauth): 12 months
Other Criteria	Cushing's Syndrome (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ITOVEBI (S)

Products Affected

- Itovebi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of breast cancer. Disease is one of the following: a) Locally advanced, or b) Metastatic. Disease is all of the following (as detected by a U.S. Food and Drug Administration [FDA]-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments [CLIA]): a) PIK3CA-mutated, b) Hormone receptor (HR)-positive, 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ITRACONAZOLE CAPSULE (S)

Products Affected

- Itraconazole CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Systemic Fungal Infection (SFI): Diagnosis of a systemic fungal infection (e.g., aspergillosis, histoplasmosis, blastomycosis). Fingernail Onychomycosis: Diagnosis of fingernail onychomycosis as confirmed by one of the following: i) positive potassium hydroxide (KOH) preparation, ii) fungal culture, OR iii) nail biopsy. Trial and failure (of a minimum 6-week supply), contraindication, or intolerance to oral terbinafine. Toenail Onychomycosis: Diagnosis of toenail onychomycosis as confirmed by one of the following: i) positive potassium hydroxide (KOH) preparation, ii) fungal culture, OR iii) nail biopsy. Trial and failure (of a minimum 12-week supply), contraindication, or intolerance to oral terbinafine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	SFI:6mo.Fingernail Onychomycosis:5wks.Toenail Onychomycosis:3mo.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IVERMECTIN (S)

Products Affected

- Ivermectin TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Strongyloidiasis: Diagnosis of intestinal (i.e., nondisseminated) strongyloidiasis due to the nematode parasite Strongyloides stercoralis. Onchocerciasis: Diagnosis of onchocerciasis due to the nematode parasite Onchocerca volvulus.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Strongyloidiasis: 3 weeks. Onchocerciasis: 6 months.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IVIG 2 (S)

Products Affected

- Bivigam INJ 5GM/50ML
- Privigen

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

IWILFIN (S)

Products Affected

- Iwilfin

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JAKAFI (S)

Products Affected

- Jakafi
- Jakafi Xr

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Myelofibrosis: Diagnosis of primary myelofibrosis, OR post-polycythemia vera myelofibrosis, OR post-essential thrombocythemia myelofibrosis. Polycythemia vera: Diagnosis of polycythemia vera, AND trial and failure, contraindication, or intolerance to hydroxyurea. Acute graft versus host disease (aGVHD): Diagnosis of aGVHD. Disease is steroid-refractory. Chronic graft versus host disease (cGVHD): Diagnosis of cGVHD. Trial and failure of at least one or more lines of systemic therapy (e.g., corticosteroids, mycophenolate, etc.).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months.
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JAYPIRCA (S)

Products Affected

- Jaypirca

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Mantle Cell Lymphoma (MCL): Diagnosis of MCL. Disease is one of the following: a) relapsed, or b) refractory. Patient has received at least two prior therapies 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. Disease is one of the following: a) Relapsed, or b) Refractory. Patient has previously been treated with a covalent BTK inhibitor [e.g., Calquence (acalabrutinib), Imbruvica (ibrutinib)].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JIDEYTRO (S)

Products Affected

- Jideytro

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

JYLAMVO (S)

Products Affected

- Jylamvo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Neoplastic diseases: Diagnosis of one of the following: A) acute lymphoblastic leukemia (ALL), B) mycosis fungoides (cutaneous T-cell lymphoma), or C) relapsed or refractory non-hodgkin lymphomas. Rheumatoid arthritis (RA): Diagnosis of RA. Psoriasis: Diagnosis of severe psoriasis. Polyarticular juvenile idiopathic arthritis (pJIA): Diagnosis of polyarticular juvenile idiopathic arthritis.
Age Restrictions	N/A
Prescriber Restrictions	RA: Prescribed by or in consultation with a rheumatologist. Psoriasis: Prescribed by or in consultation with a dermatologist. pJIA: Prescribed by or in consultation with a rheumatologist.
Coverage Duration	Neoplastic diseases, RA, Psoriasis, pJIA: 12 months.
Other Criteria	Subject to Part B vs D review. Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

JYNARQUE (S)

Products Affected

- Tolvaptan TABS

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

KALYDECO (S)

Products Affected

- Kalydeco

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

KERENDIA (S)

Products Affected

- Kerendia

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

KESIMPTA (S)

Products Affected

- Kesimpta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple Sclerosis (MS) (Initial): Diagnosis of a relapsing form of multiple sclerosis (MS) (e.g., clinically isolated syndrome, relapsing-remitting disease, secondary progressive disease, including active disease with new brain lesions). One of the following: 1) Trial and failure (of a minimum 4-week supply), contraindication, or intolerance to one disease-modifying therapy for MS [e.g., Avonex (interferon beta-1a), Betaseron (interferon beta-1b), Gilenya fingolimod)], OR 2) For continuation of prior therapy. Not used in combination with another disease-modifying therapy for MS. Not used in combination with another B-cell targeted therapy (e.g., rituximab [Rituxan], belimumab [Benlysta], ocrelizumab [Ocrevus]).
Age Restrictions	N/A
Prescriber Restrictions	MS (Initial, Reauth): Prescribed by or in consultation with a neurologist
Coverage Duration	MS (Initial, Reauth): 12 months
Other Criteria	MS (Reauth): Patient demonstrates positive clinical response to therapy. Not used in combination with another disease-modifying therapy for MS. Not used in combination with another B-cell targeted therapy (e.g., rituximab [Rituxan], belimumab [Benlysta], ocrelizumab [Ocrevus]).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KINERET (S)

Products Affected

- Kineret

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

KISQALI (S)

Products Affected

- Kisqali

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Breast cancer: Diagnosis of breast cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KOMZIFTI (S)

Products Affected

- Komzifti

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acute myeloid leukemia (AML). Disease is relapsed or refractory. Presence of a susceptible nucleophosmin 1 (NPM1) mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KORLYM (S)

Products Affected

- Mifepristone TABS 300MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cushing's syndrome (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.
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Initial, reauth: 6 months
Other Criteria	Reauth: Patient demonstrates one of the following: patient has improved glucose tolerance while on therapy or patient has stable glucose tolerance while on therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KOSELUGO (S)

Products Affected

- Koselugo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KRAZATI (S)

Products Affected

- Krazati

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: locally advanced or metastatic. Disease is KRAS G12C-mutated as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received at least one prior systemic therapy (e.g., chemotherapy, immunotherapy). Colorectal Cancer (CRC): Diagnosis of CRC. Disease is one of the following: locally advanced or metastatic. Tumor is KRAS G12C-mutated as detected by a U.S. Food and Drug Administration (FDA) -approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is used in combination with Erbitux (cetuximab). Patient has received prior treatment with ALL of the following: a) fluoropyrimidine-based chemotherapy, b) oxaliplatin-based chemotherapy, and c) irinotecan-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KUVAN (S)

Products Affected

- Sapropterin Dihydrochloride
- Zelvysia

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Phenylketonuria (PKU) (initial): Diagnosis of PKU. Patient will have blood Phe levels measured after 1 week of therapy (new starts to therapy only) and periodically for up to 2 months of therapy to determine response.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	PKU (Init): 2 months (Reauth): 12 months
Other Criteria	PKU (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KYGEVVI (S)

Products Affected

- Kygevvi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Submission of medical records (e.g., chart notes) confirming all of the following: a) Diagnosis of thymidine kinase 2 deficiency (TK2d), b) Presence of a mutation in the TK2 gene as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA), c) Patient experienced symptom onset on or before 12 years of age.
Age Restrictions	N/A
Prescriber Restrictions	Initial, Reauth: Prescribed by or in consultation with a clinical specialist knowledgeable in the treatment of TK2d (e.g., geneticist, neurologist).
Coverage Duration	Initial, Reauth: 12 months
Other Criteria	Reauth: Provider attests the patient would benefit from continued therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LAZCLUZE (S)

Products Affected

- Lazcluze

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-small cell lung cancer (NSCLC). Disease is locally advanced or metastatic. Used as first line treatment of NSCLC. Used in combination with Rybrevant (amivantamab). Presence of epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R substitution mutations as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LENVIMA (S)

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 Criteria	N/A
Required Medical Information	Differentiated thyroid cancer (DTC): Diagnosis of DTC. Renal Cell Carcinoma (RCC): Diagnosis of RCC. One of the following: 1) Both of the following: a) Used as first-line treatment and b) Used in combination with Keytruda (pembrolizumab), or 2) Both of the following: a) Treatment follows one prior anti-angiogenic therapy and b) Used in combination with everolimus. Hepatocellular Carcinoma (HCC): Diagnosis of HCC. Endometrial Carcinoma (EC): Diagnosis of advanced endometrial carcinoma that is not microsatellite instability-high (MSI-H), or mismatch repair deficient (dMMR) [i.e. disease is mismatch repair proficient (pMMR)], as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has disease progression following prior systemic therapy. Used in combination with Keytruda (pembrolizumab). Patient is not a candidate for curative surgery or radiation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.

Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LEQEMBI IQLIK (S)

Products Affected

- Leqembi Iqlik

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

LEUKINE (S)

Products Affected

- Leukine INJ 250MCG

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

LIDOCAINE TOPICAL (S)

Products Affected

- Lidocaine OINT 5%
- Premium Lidocaine

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The drug is being prescribed for an FDA-approved indication or compendia supported use.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIDOCAINE/PRILOCAINE (S)

Products Affected

- Lidocaine/prilocaine CREA

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The drug is being prescribed for an FDA-approved indication or compendia supported use.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIDODERM (S)

Products Affected

- Lidocaine PTCH 5%

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Post-herpetic neuralgia: Diagnosis of post-herpetic neuralgia.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIFYORLI (S)

Products Affected

- Lifyorli

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer. Disease is platinum resistant. Used in combination with nab-paclitaxel. Patient has received one prior bevacizumab-containing systemic treatment regimen. Patient has not received more than three prior systemic treatment regimens (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LONSURF (S)

Products Affected

- Lonsurf

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Metastatic Colorectal Cancer (mCRC): Diagnosis of mCRC. One of the following: A) Used as a single agent or B) Used in combination with bevacizumab. Patient has been previously treated with both of the following: A) fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy (e.g., FOLFOX, FOLFIRI, FOLFOXIRI) AND B) anti-VEGF therapy (e.g., Avastin [bevacizumab]). One of the following: A) Presence of RAS wild-type tumors AND patient has been previously treated with one anti-EGFR therapy (e.g., Vectibix [panitumumab], Erbitux [cetuximab]) OR B) Presence of RAS mutation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Metastatic Gastric/Gastroesophageal Junction Adenocarcinoma: Diagnosis of metastatic gastric cancer or metastatic gastroesophageal junction adenocarcinoma. Patient has been previously treated with two of the following: A) fluropyrimidine-based chemotherapy (e.g. fluorouracil), B) Platinum-based chemotherapy (e.g., carboplatin, cisplatin, oxaliplatin), C) Taxane (e.g., docetaxel, paclitaxel) or irinotecan-based chemotherapy, or D) HER2/neu-targeted therapy (e.g., trastuzumab) (if HER2 overexpression).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.

Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LORBRENA (S)

Products Affected

- Lorbrena

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of metastatic NSCLC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LOTRONEX (S)

Products Affected

- Alosetron Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Severe Diarrhea-Predominant Irritable Bowel Syndrome (IBS) in Women (initial): All of the following: 1) diagnosis of severe diarrhea-predominant IBS, 2) symptoms for at least 6 months, 3) female patient, AND 4) trial and failure, contraindication, or intolerance to an antidiarrheal agent [eg, loperamide].
Age Restrictions	Initial: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	IBS (initial): 12 weeks. IBS (reauth): 6 mo.
Other Criteria	IBS (reauth): Symptoms of IBS continue to persist. Patient demonstrates positive clinical response to therapy (e.g., relief of IBS abdominal pain and discomfort, improvement in stool consistency and frequency, improvement as measured by the Global Improvement Scale).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LUMAKRAS (S)

Products Affected

- Lumakras

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is one of the following: a) locally advanced or b) metastatic. Presence of KRAS G12C-mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received at least one prior systemic therapy (e.g., chemotherapy, immunotherapy). Metastatic Colorectal Cancer (mCRC): Diagnosis of mCRC. Presence of KRAS G12C-mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received prior therapy with fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy. Used in combination with Vectibix (panitumumab).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LUPRON (S)

Products Affected

- Leuprolide Acetate INJ 1MG/0.2ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prostate Cancer: Diagnosis of advanced or metastatic prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Prostate CA: 12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LUPRON DEPOT (S)

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 Criteria	N/A
Required Medical Information	Prostate Cancer (7.5 mg, 22.5 mg, 30 mg, 45 mg): Diagnosis of advanced or metastatic prostate cancer. Endometriosis (3.75 mg, 11.25 mg): Diagnosis of endometriosis. One of the following: Patient has had surgical ablation to prevent recurrence, or 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). Uterine Leiomyomata (UL) (3.75 mg, 11.25 mg): a) For use prior to surgery to reduce size of fibroids to facilitate a surgical procedure (eg, myomectomy, hysterectomy) OR b) all of the following: treatment of anemia, anemia is caused by uterine leiomyomata (fibroids), and for use prior to surgery.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Prostate CA: 12 mo. Endomet:6mo. UL (anemia):3 mo (fibroids):4 mo
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LUPRON DEPOT PED (S)

Products Affected

- Lupron Depot-ped (1-month)
- Lupron Depot-ped (3-month) INJ 30MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Central Precocious Puberty (CPP) (initial): Diagnosis of CPP (idiopathic or neurogenic). Early onset of secondary sexual characteristics in females less than age 8 or males less than age 9. Advanced bone age of at least one year compared with chronologic age. One of the following: a) patient has undergone gonadotropin-releasing hormone agonist (GnRHa) testing AND Peak luteinizing hormone (LH) level above pre-pubertal range, or b) patient has a random LH level in the pubertal range.
Age Restrictions	N/A
Prescriber Restrictions	CPP (initial, reauth): Prescribed by or in consultation with an endocrinologist.
Coverage Duration	CPP (initial, reauth): 12 months
Other Criteria	CPP (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LYNPARZA TABLET (S)

Products Affected

- Lynparza TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Epithelial ovarian, fallopian tube, or primary peritoneal cancer: Diagnosis of one of the following: epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer. Breast cancer: Diagnosis of breast cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Pancreatic adenocarcinoma: Diagnosis of pancreatic adenocarcinoma. Prostate cancer: Diagnosis of castration-resistant prostate cancer. All indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LYTGOBI (S)

Products Affected

- Lytgobi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of intrahepatic cholangiocarcinoma. Disease is one of the following: a) unresectable, b) locally advanced, or c) metastatic. Disease has presence of a fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangements. Patient has been previously treated (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MARINOL (S)

Products Affected

- Dronabinol CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Nausea and Vomiting Associated with Cancer Chemotherapy (CINV): Patient is receiving cancer chemotherapy. Trial and failure, contraindication, or intolerance to one 5HT-3 receptor antagonist (eg, Anzemet [dolasetron], Kytril [granisetron], or Zofran [ondansetron]). Trial and failure, contraindication, or intolerance to one of the following: Compazine (prochlorperazine), Decadron (dexamethasone), Haldol (haloperidol), Zyprexa (olanzapine). AIDS anorexia: Diagnosis of anorexia with weight loss in patients with AIDS. Patient is on antiretroviral therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	CINV: 6 months. AIDS anorexia: 12 months.
Other Criteria	Subject to Part B vs. Part D review. CINV: Approve for continuation of therapy for treatment covered under Part B when patient is receiving cancer chemotherapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MAVYRET (S)

Products Affected

- Mavyret

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Criteria will be applied consistent with current AASLD/IDSA guideline. All patients: 1) One of the following: a) Diagnosis of hepatitis C, or b) Patient was not infected with hepatitis C virus prior to receiving an organ transplant, and patient received a liver or non-liver organ transplant from a donor with a diagnosis of hepatitis C, 2) patient is without decompensated liver disease (defined as Child-Pugh Class B or C), and 3) not used in combination with another HCV direct acting antiviral agent [e.g., Harvoni, Zepatier].
Age Restrictions	N/A
Prescriber Restrictions	One of the following: 1) Non-transplant patient who is treatment-naïve without decompensation, or 2) Prescribed by or in consultation with one of the following: Hepatologist, Gastroenterologist, Infectious disease specialist, HIV specialist certified through the American Academy of HIV Medicine, Liver transplant specialist.
Coverage Duration	8 to 16 weeks. Criteria will be applied consistent with current AASLD/IDSA guideline.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MEKINIST (S)

Products Affected

- Mekinist

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable or metastatic melanoma AND cancer is BRAF V600E or V600K mutant type as detected by a U.S. Food and Drug Administration (FDA)-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Adjuvant Treatment for Melanoma: Diagnosis of melanoma. Cancer is BRAF V600E or V600K mutant type as detected by an FDA-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Involvement of lymph nodes following complete resection. Used as adjunctive therapy. Medication is used in combination with Tafinlar (dabrafenib).Non-small Cell Lung Cancer (NSCLC): All of the following: diagnosis of metastatic non-small cell lung cancer AND cancer is BRAF V600E mutant type as detected by an FDA-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA) AND medication is used in combination with Tafinlar (dabrafenib). Anaplastic Thyroid Cancer (ATC): Diagnosis of locally advanced or metastatic ATC. Cancer is BRAF V600E mutant type as detected by an FDA-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Cancer may not be treated with standard locoregional treatment options. Medication is used in combination with Tafinlar (dabrafenib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months

Other Criteria	Approve for continuation of prior therapy. Solid tumors: Diagnosis of solid tumors. Disease is unresectable or metastatic. Patient has progressed on or following prior treatment and have no satisfactory alternative treatment options. Cancer is BRAF V600E mutant type as detected by an FDA-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is used in combination with Tafinlar (dabrafenib). Low-grade glioma: Diagnosis of low-grade glioma. Patient requires systemic therapy. Cancer is BRAF V600E mutant type as detected by an FDA-approved test (THxID-BRAF Kit) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is used in combination with Tafinlar (dabrafenib).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MEKTOVI (S)

Products Affected

- Mektovi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma. Presence of BRAF V600E or V600K mutant type (MT) as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with Braftovi (encorafenib). Non-Small Cell Lung Cancer (NSCLC): Diagnosis of metastatic NSCLC. Presence of BRAF V600E mutant type (MT) as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with Braftovi (encorafenib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MIGRANAL (S)

Products Affected

- Dihydroergotamine Mesylate SOLN

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of migraine headaches with or without aura. Will be used for the acute treatment of migraine. One of the following: Trial and failure or intolerance to one triptan (e.g., eletriptan, rizatriptan, sumatriptan) or contraindication to all triptans.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial, Reauth: 12 months.
Other Criteria	Reauth: Patient has experienced a positive response to therapy (e.g., reduction in pain, photophobia, phonophobia, nausea).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MIMRYLO (S)

Products Affected

- Mimrylo

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

MODEYSO (S)

Products Affected

- Modeyso

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of diffuse midline glioma. Disease is confirmed by the presence of H3 K27M mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease has progressed following prior therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MOUNJARO (S)

Products Affected

- Mounjaro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: One of the following: a) For patients requiring ongoing treatment for type 2 diabetes mellitus (T2DM), submission of medical records (e.g., chart notes) confirming diagnosis of T2DM, OR b) Submission of medical records (e.g., chart notes) confirming diagnosis of T2DM as evidenced by one of the following laboratory values: i) A1c greater than or equal to 6.5%, ii) fasting plasma glucose (FPG) greater than or equal to 126 mg/dL, or iii) 2-hour plasma glucose (PG) greater than or equal to 200 mg/dL during OGTT (oral glucose tolerance test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MS INTERFERONS (PREFERRED) - BETASERON (S)

Products Affected

- Betaseron

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Multiple Sclerosis (MS) (initial, reauth): Not used in combination with another disease-modifying therapy for MS.
Required Medical Information	MS (initial): Diagnosis of a relapsing form of MS (eg, clinically isolated syndrome, relapsing-remitting disease, secondary progressive disease, including active disease with new brain lesions).
Age Restrictions	N/A
Prescriber Restrictions	MS (initial, reauth): Prescribed by or in consultation with a neurologist
Coverage Duration	MS (initial, reauth): 12 months
Other Criteria	MS (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NERLYNX (S)

Products Affected

- Nerlynx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Early Stage Breast cancer: Diagnosis (dx) of early stage breast cancer. Advanced or Metastatic Breast Cancer: Dx of advanced or metastatic breast cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NEULASTA (S)

Products Affected

- Neulasta
- Neulasta Onpro Kit

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

NEXAVAR (S)

Products Affected

- Sorafenib
- Sorafenib Tosylate TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Renal cell carcinoma (RCC): Diagnosis of advanced RCC. Hepatocellular carcinoma (HCC): Diagnosis of HCC. Differentiated thyroid carcinoma (DTC): Diagnosis of DTC. One of the following: locally recurrent disease, or metastatic disease. Patient has progressive disease. Disease is refractory to radioactive iodine (RAI) treatment.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NEXLETOL (S)

Products Affected

- Nexletol

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

NEXLIZET (S)

Products Affected

- Nexlizet

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

NINLARO (S)

Products Affected

- Ninlaro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple myeloma: Diagnosis of multiple myeloma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NORTHERA (S)

Products Affected

- Droxidopa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Neurogenic orthostatic hypotension (NOH) (init): 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. Trial and failure, contraindication, or intolerance to one of the following agents: fludrocortisone acetate, midodrine.
Age Restrictions	N/A
Prescriber Restrictions	NOH (init): Prescribed by or in consultation with a cardiologist, neurologist, or nephrologist
Coverage Duration	NOH (init): 1 month (reauth): 12 months
Other Criteria	NOH (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NOXAFIL SUSPENSION (S)

Products Affected

- Posaconazole SUSP

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prophylaxis of systemic fungal infections (SFI): 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 OR 2) patient has a prior fungal infection requiring secondary prophylaxis. Oropharyngeal Candidiasis (OPC): Diagnosis of OPC. One of the following: 1) Trial and failure, contraindication, or intolerance to fluconazole OR 2) Susceptibility results demonstrate resistance to fluconazole.
Age Restrictions	Prophylaxis of SFI, OPC: Patient is 13 years or older.
Prescriber Restrictions	N/A
Coverage Duration	Prophylaxis of SFI: 6 months. OPC: 1 month.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NUBEQA (S)

Products Affected

- Nubeqa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Castration-resistant prostate cancer (CRPC): Diagnosis of castration-resistant (chemical or surgical) prostate cancer. Metastatic Castration-sensitive prostate cancer (mCSPC): Diagnosis of metastatic castration-sensitive prostate cancer (mCSPC) [also known as metastatic hormone-sensitive prostate cancer (mHSPC)].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	CRPC, mCSPC: 12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NUCALA (S)

Products Affected

- Nucala

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

NUEDEXTA (S)

Products Affected

- Dextromethorphan Hydrobromide/quinidine Sulfate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Pseudobulbar affect (PBA) (initial): Diagnosis of PBA. Patient does not have any of the following contraindications: a) Concomitant use with other drugs containing quinidine, quinine, or mefloquine, b) History of Nuedexta, quinine, mefloquine or quinidine-induced thrombocytopenia, hepatitis, bone marrow depression, or lupus-like syndrome, c) Known hypersensitivity to dextromethorphan (e.g., rash, hives), d) Taking monoamine oxidase inhibitors (MAOIs) (e.g., phenelzine, selegiline, tranylcypromine) or have taken MAOIs within the preceding 14 days, e) Has prolonged QT interval, congenital long QT syndrome or a history suggestive of torsades de pointes, or has heart failure, f) Receiving drugs that both prolong QT interval and are metabolized by CYP2D6 (e.g., thioridazine, pimozide), g) Has complete atrioventricular (AV) block without implanted pacemakers, or at high risk of complete AV block.
Age Restrictions	N/A
Prescriber Restrictions	PBA (initial): Prescribed by or in consultation with one of the following specialists: neurologist, psychiatrist.
Coverage Duration	PBA (initial/reauth): 12 months
Other Criteria	PBA (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NUPLAZID (S)

Products Affected

- Nuplazid CAPS
- Nuplazid TABS 10MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NURTEC (S)

Products Affected

- Nurtec

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

NUVIGIL (S)

Products Affected

- Armodafinil

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

ODOMZO (S)

Products Affected

- Odomzo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of basal cell carcinoma. Disease is locally advanced. One of the following: 1) Disease has recurred following surgery or radiation therapy or 2) Patient is not a candidate for surgery or radiation therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OFEV BRAND (S)

Products Affected

- Ofev

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

OFEV GENERIC (S)

Products Affected

- Nintedanib Esylate

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

OGSIVEO (S)

Products Affected

- Ogsiveo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of desmoid tumor. Patient requires systemic treatment. Disease is progressive.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OJEMDA (S)

Products Affected

- Ojemda

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of pediatric low-grade glioma. Disease is relapsed or refractory. Presence of a BRAF fusion or rearrangement, or BRAF V600 mutation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OJJAARA (S)

Products Affected

- Ojjaara

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of ONE of the following: a) Primary myelofibrosis, b) Post-polycythemia vera myelofibrosis, OR c) Post-essential thrombocythemia myelofibrosis. Disease is intermediate or high risk. Patient has anemia.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ONPATTRO (S)

Products Affected

- Onpattro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Hereditary transthyretin-mediated amyloidosis (hATTR amyloidosis) (initial): Diagnosis of hATTR amyloidosis with polyneuropathy. Presence of a transthyretin (TTR) mutation (e.g., V30M) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Presence of clinical signs and symptoms of the disease (e.g., peripheral/autonomic neuropathy, walking ability, quality of life). Requested drug is not used in combination with a TTR silencer (e.g., Wainua) or a TTR stabilizer (e.g., Vyndamax).
Age Restrictions	N/A
Prescriber Restrictions	hATTR amyloidosis (initial): Prescribed by or in consultation with a neurologist.
Coverage Duration	hATTR amyloidosis (initial, reauth): 12 months.
Other Criteria	Subject to Part B vs D review. hATTR amyloidosis (reauth): Patient demonstrates postive clinical response to therapy. Requested drug is not used in combination with a TTR silencer (e.g., Wainua) or a TTR stabilizer (e.g., Vyndamax).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ONUREG (S)

Products Affected

- Onureg

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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, etc.). 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 able to complete intensive curative therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OPIPZA (S)

Products Affected

- Opipza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Schizophrenia: Diagnosis of Schizophrenia. Trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: a) aripiprazole (failure or contraindication are not required), b) olanzapine, c) quetiapine IR/ER, d) risperidone, e) clozapine, f) ziprasidone, g) paliperidone, or h) asenapine. Major Depressive Disorder (MDD): Diagnosis of MDD. Used as adjunctive therapy with an antidepressant. Both of the following: a) TF/C/I to quetiapine IR/ER and b) Trial of or intolerance to aripiprazole. Autism: Diagnosis of irritability associated with autistic disorder. Both of the following: a) Trial and failure, contraindication (e.g., age), or intolerance to risperidone and b) Trial of or intolerance to aripiprazole. Tourette's Syndrome: Diagnosis of Tourette's Syndrome. Trial of or intolerance to aripiprazole.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OPSUMIT (S)

Products Affected

- Macitentan

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

ORENCIA IV (S)

Products Affected

- Orencia INJ 250MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Rheumatoid Arthritis (RA) (Initial): Diagnosis of moderately to severely active RA. One of the following: a) Either a trial and failure, contraindication, or intolerance (TF/C/I) to two of the following: Enbrel (etanercept), one formulary adalimumab product, Rinvoq (upadacitinib), tofacitinib, or attestation demonstrating a trial may be inappropriate, OR b) For continuation of prior therapy. Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Diagnosis of moderately to severely active PJIA. One of the following: a) TF/C/I to two of the following, or attestation demonstrating a trial may be inappropriate: Enbrel, one formulary adalimumab product, Rinvoq/LQ, or tofacitinib, OR b) for continuation of prior therapy. 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. One of the following: a) Either a TF/C/I to two of the following: Cosentyx SC (secukinumab), Enbrel, one formulary adalimumab product, Otezla (apremilast), Skyrizi (risankizumab-rzaa), Sotyktu (deucravacitinib), Tremfya (guselkumab), one formulary ustekinumab product, Rinvoq/LQ, or tofacitinib, OR b) for continuation of prior therapy. Acute graft versus host disease (aGVHD): Used for prophylaxis of aGVHD. Patient will receive hematopoietic stem cell transplantation (HSCT) from a matched or 1 allele-mismatched unrelated donor. Recommended antiviral prophylactic treatment for Epstein-Barr Virus (EBV) reactivation (e.g., acyclovir) will be administered prior to Orencia and continued for six months after HSCT. Used in combination with both of the following: calcineurin inhibitor (e.g., cyclosporine, tacrolimus) and methotrexate.
Age Restrictions	N/A
Prescriber Restrictions	RA, JIA (initial): Prescribed by or in consultation with a rheumatologist. PsA (initial): Prescribed by or in consultation with a dermatologist or rheumatologist.

Coverage Duration	RA, JIA, PsA (initial): 6 months, (reauth): 12 months. aGVHD: 2 months
Other Criteria	RA, PJIA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline OR improvement in symptoms (eg, pain, stiffness, inflammation) from baseline. PsA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline, improvement in symptoms (eg, pain, stiffness, pruritus, inflammation) from baseline, OR reduction in the BSA involvement from baseline.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ORENCIA SC (S)

Products Affected

- Orencia INJ 125MG/ML, 50MG/0.4ML, 87.5MG/0.7ML
- Orencia Clickject

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

ORENITRAM (S)

Products Affected

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

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

ORGOVYX (S)

Products Affected

- Orgovyx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prostate Cancer: Diagnosis of advanced or metastatic prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORKAMBI (S)

Products Affected

- Orkambi TABS

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

ORSERDU (S)

Products Affected

- Orserdu

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of breast cancer. Disease is advanced or metastatic. Disease is estrogen receptor (ER)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Presence of estrogen receptor (ESR1) mutation(s) as detected with a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease has progressed following at least one line of endocrine therapy [e.g., Faslodex (fulvestrant), Arimidex (anastrozole), Femara (letrozole), Aromasin (exemestane)].
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OSENVELT (S)

Products Affected

- Osenvelt

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Skeletal prevention in Multiple Myeloma (MM)/Bone Metastasis from Solid Tumors (BMST): One of the following: 1) Both of the following: a) Diagnosis of multiple myeloma and b) Trial and failure, contraindication (e.g., renal insufficiency), or intolerance to one bisphosphonate therapy, OR 2) Both of the following: a) Diagnosis of solid tumors (e.g., breast cancer, kidney cancer, lung cancer, prostate cancer, thyroid cancer) and b) Documented evidence of one or more metastatic bone lesions. Giant cell tumor of bone (GCTB): Diagnosis of giant cell tumor of bone. One of the following: 1) One of the following: a) tumor is unresectable or b) surgical resection is likely to result in severe morbidity, OR 2) Approve for continuation of prior therapy. Hypercalcemia of malignancy (HCM): Diagnosis of hypercalcemia of malignancy. Trial and failure, contraindication, or intolerance to one bisphosphonate therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	MM/BMST, GCTB, HCM: 12 months.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OSPHENA (S)

Products Affected

- Osphena

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Dyspareunia (initial): Diagnosis of moderate to severe dyspareunia due to vulvar and vaginal atrophy associated with menopause. Vaginal dryness (initial): Diagnosis of moderate to severe vaginal dryness due to vulvar and vaginal atrophy associated with menopause.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses (Initial, reauth): 12 months
Other Criteria	Dyspareunia, Vaginal dryness (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OTEZLA (S)

Products Affected

- Otezla

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

OTEZLA XR (S)

Products Affected

- Otezla Xr
- Otezla/otezla Xr 28 Day Treatment Initiation Pack

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

OZEMPIC (S)

Products Affected

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

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

OZEMPIC TABLET (S)

Products Affected

- Ozempic TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: One of the following: a) For patients requiring ongoing treatment for type 2 diabetes mellitus (T2DM), submission of medical records (e.g., chart notes) confirming diagnosis of T2DM, OR b) Submission of medical records (e.g., chart notes) confirming diagnosis of T2DM as evidenced by one of the following laboratory values: i) A1c greater than or equal to 6.5%, ii) fasting plasma glucose (FPG) greater than or equal to 126 mg/dL, or iii) 2-hour plasma glucose (PG) greater than or equal to 200 mg/dL during OGTT (oral glucose tolerance test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial, Reauth: 12 months.
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PEGASYS (S)

Products Affected

- Pegasys

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	Chronic Hepatitis C: Prescribed by or in consultation with one of the following: hepatologist, gastroenterologist, infectious disease specialist, HIV specialist certified through the American Academy of HIV Medicine
Coverage Duration	HepB, HepC: 48 wks.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PEMAZYRE (S)

Products Affected

- Pemazyre

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cholangiocarcinoma: Diagnosis of cholangiocarcinoma. Disease is one of the following: unresectable locally advanced or metastatic. Disease is confirmed by the presence of a fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has received at least one prior line of therapy (e.g., chemotherapy). Myeloid/lymphoid neoplasms: Diagnosis of myeloid/lymphoid neoplasms (MLNs). Disease is relapsed or refractory. Disease is confirmed by the presence of a fibroblast growth factor receptor 1 (FGFR1) rearrangement as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PENNSAID (S)

Products Affected

- Diclofenac Sodium TOPICAL SOLN 1.5%

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

PHESGO (S)

Products Affected

- Phesgo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PIQRAY (S)

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 Criteria	N/A
Required Medical Information	Diagnosis of advanced or metastatic breast cancer. Disease is hormone receptor (HR)-positive, and human epidermal growth factor receptor 2 (HER2)-negative. Disease is confirmed by the presence of PIK3CA-mutated as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with fulvestrant. Disease has progressed on or after an endocrine-based regimen.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

POMALYST BRAND (S)

Products Affected

- Pomalyst

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple Myeloma (MM): Diagnosis of MM. Kaposi sarcoma (KS): One of the following: 1) Diagnosis of AIDS-related KS, OR 2) Both of the following: a) Diagnosis of KS and b) Patient is HIV-negative. Trial or intolerance to pomalidomide.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

POMALYST GENERIC (S)

Products Affected

- Pomalidomide

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple Myeloma (MM): Diagnosis of MM. Kaposi sarcoma (KS): One of the following: 1) Diagnosis of AIDS-related KS, OR 2) Both of the following: a) Diagnosis of KS and b) Patient is HIV-negative.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

POSACONAZOLE TABLET (S)

Products Affected

- Posaconazole Dr

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prophylaxis of systemic fungal infections (SFI): 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 OR 2) patient has a prior fungal infection requiring secondary prophylaxis. Treatment (Tx) of SFI: Used as treatment of systemic fungal infections caused by Aspergillus. All indications: Patient weighs greater than 40 kg.
Age Restrictions	All indications: Patient is 2 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	All indications: 6 months.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PROMACTA (S)

Products Affected

- Eltrombopag Olamine

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

PROMETHAZINE (S)

Products Affected

- Promethazine Hcl SUPP 12.5MG
- Promethazine Hydrochloride SOLN 6.25MG/5ML
- Promethazine Hydrochloride TABS
- Promethazine Hydrochloride Plain

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	One of the following: 1) Requested drug is FDA-approved for the condition being treated, or 2) If requested for an off-label indication, the off-label guideline approval criteria have been met. Prescriber acknowledges anticholinergic risks (e.g., confusion, dry mouth, blurry vision, constipation, urinary retention) and will consider lowering the dose or discontinuing medication(s) that are no longer clinically warranted for the patient.
Age Restrictions	Patient is 65 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PROVIGIL (S)

Products Affected

- Modafinil TABS

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

PULMOZYME (S)

Products Affected

- Pulmozyme SOLN 2.5MG/2.5ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cystic Fibrosis (CF) (Initial, Reauth): Diagnosis of CF.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	CF (initial, reauth): 12 months
Other Criteria	Part B vs D determination applies. CF (reauth): Patient is benefiting from treatment (i.e. improvement in lung function [forced expiratory volume in one second (FEV1)], decreased number of pulmonary exacerbations).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PYRUKYND (S)

Products Affected

- Pyrukynd
- Pyrukynd Taper Pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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). Disease is confirmed by the presence of ALL the following mutations on the PKLR gene as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA): 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. Other causes of hemolytic anemias (e.g., infections, toxins, drugs) have been excluded.
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with a hematologist.
Coverage Duration	Initial: 6 months. Reauth: 12 months.
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

QINLOCK (S)

Products Affected

- Qinlock

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Gastrointestinal Stromal Tumor (GIST): Diagnosis of gastrointestinal stromal tumor (GIST). Disease is advanced. Patient has received prior treatment with three or more kinase inhibitors (e.g., sunitinib, regorafenib), one of which must include imatinib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

QUALAQUIN (S)

Products Affected

- Quinine Sulfate CAPS 324MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Excluded if used solely for the treatment or prevention of nocturnal leg cramps.
Required Medical Information	Malaria: Diagnosis of uncomplicated malaria. One of the following: 1) Treatment in areas of chloroquine-sensitive malaria, and trial and failure, contraindication, or intolerance to chloroquine or hydroxychloroquine, OR 2) Treatment in areas of chloroquine-resistant malaria.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	7 days
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RASONQUE (S)

Products Affected

- Rasonque

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

RELGAABI (S)

Products Affected

- Gabapentin CAPS 200MG

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

REMICADE (S)

Products Affected

- Infliximab

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's Disease (CD) and Fistulizing Crohn's Disease (FCD) (Initial): Diagnosis (Dx) of moderately to severely active CD or FCD. 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. Trial and failure, contraindication, or intolerance (TF/C/I) to one of the following conventional therapies: 6-mercaptopurine, azathioprine, corticosteroid (eg, prednisone), methotrexate. 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. TF/C/I to one of the following conventional therapies: 6-mercaptopurine, azathioprine, corticosteroid (eg, prednisone), aminosalicylate (eg, mesalamine, olsalazine, sulfasalazine). Rheumatoid arthritis (RA) (Initial): Dx of moderately to severely active RA. Used in combination with 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 (Initial): Dx of chronic severe (ie, extensive and/or disabling) 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. Minimum duration of a 4-week trial and failure, 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).
Age Restrictions	N/A

Prescriber Restrictions	CD, FCD, UC (initial): Prescribed by or in consultation with a gastroenterologist. RA, AS (initial): Prescribed by or in consultation with a rheumatologist. PsA (initial): Prescribed by or in consultation with rheumatologist or dermatologist. Plaque Psoriasis (initial): Prescribed by or in consultation with a dermatologist. Sarcoidosis (initial): Prescribed by or in consultation with a pulmonologist, dermatologist, or ophthalmologist.
Coverage Duration	All uses (initial): 6 months, (reauth): 12 months
Other Criteria	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. Sarcoidosis (Initial): Dx of sarcoidosis. TF/C/I to both of the following: one immunosuppressant (eg, methotrexate, cyclophosphamide, azathioprine) AND one corticosteroid (eg, prednisone). Plaque psoriasis (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by one of the following: reduction in the BSA involvement from baseline, OR improvement in symptoms (eg, pruritus, inflammation) from baseline. CD, UC (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: improvement in intestinal inflammation (eg, mucosal healing, improvement of lab values [platelet counts, erythrocyte sedimentation rate, C-reactive protein level]) from baseline OR reversal of high fecal output state. RA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline OR improvement in symptoms (eg, pain, stiffness, inflammation) from baseline. PsA (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by at least one of the following: reduction in the total active (swollen and tender) joint count from baseline, improvement in symptoms (eg, pain, stiffness, pruritus, inflammation) from baseline, OR reduction in the BSA involvement from baseline. AS (Reauth): Patient demonstrates positive clinical response to therapy as evidenced by improvement from baseline for at least one of the following: disease activity (eg, pain, fatigue, inflammation, stiffness), lab values (erythrocyte sedimentation rate, C-reactive protein level), function, axial status (eg, lumbar spine motion, chest expansion), OR total active (swollen and tender) joint count. Sarcoidosis (Reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

REPATHA (S)

Products Affected

- Repatha
- Repatha Sureclick

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

RETEVMO (S)

Products Affected

- Retevmo TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Disease is locally advanced or metastatic. Presence of rearranged during transfection (RET) gene fusion-positive tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medullary Thyroid Cancer (MTC): Diagnosis of MTC. Disease is advanced or metastatic. Presence of RET gene mutation tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease requires treatment with systemic therapy. Thyroid Cancer: Diagnosis of thyroid cancer. Disease is advanced or metastatic. Presence of RET gene fusion-positive tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease requires treatment with systemic therapy. Patient is radioactive iodine-refractory or radioactive iodine therapy is not appropriate. Solid Tumors: Diagnosis of solid tumors. Disease is locally advanced or metastatic. Presence of RET gene fusion-positive tumor(s) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). ONE of the following: a) Disease has progressed on or following prior systemic treatment (e.g., chemotherapy), OR b) There are no satisfactory alternative treatment options.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	NSCLC, MTC, Thyroid Cancer, Solid Tumors: 12 months
Other Criteria	Approve for continuation of prior therapy.

Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

REVATIO (S)

Products Affected

- Sildenafil Citrate TABS 20MG

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

REVCOVI (S)

Products Affected

- Revcovi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of adenosine deaminase deficiency (ADA) with severe combined immunodeficiency (SCID).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REVLIMID (S)

Products Affected

- Lenalidomide

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Multiple myeloma (MM): Diagnosis of MM. Myelodysplastic syndromes (MDS): Diagnosis of transfusion-dependent anemia due to low- or intermediate-1-risk MDS associated with a deletion 5q. Mantle cell lymphoma (MCL): Diagnosis of MCL. Follicular Lymphoma (FL): Diagnosis of FL. Marginal Zone Lymphoma (MZL): Diagnosis of MZL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REVUFORJ (S)

Products Affected

- Revuforj

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute leukemia: Diagnosis of acute leukemia. Disease is relapsed or refractory. Presence of lysine methyltransferase 2A gene (KMT2A) translocation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Acute myeloid leukemia (AML): Diagnosis of AML. Disease is relapsed or refractory. Presence of a susceptible nucleophosmin 1 (NPM1) mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZDIFFRA (S)

Products Affected

- Rezdiffra

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

REZLIDHIA (S)

Products Affected

- Rezlidhia

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acute myeloid leukemia (AML). Disease is relapsed or refractory. Presence of a susceptible isocitrate dehydrogenase-1(IDH1) mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test (e.g., Abbott RealTime IDH1 assay) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZUROCK (S)

Products Affected

- Rezurock

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic graft versus host disease (cGVHD) (initial): Diagnosis of cGVHD. Trial and failure of two or more lines of systemic therapy [e.g., corticosteroids (e.g., prednisone, methylprednisolone), mycophenolate, chemotherapy].
Age Restrictions	Initial: Patient is 12 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	cGVHD (initial, reauth): 12 months
Other Criteria	cGVHD (reauth): Patient does not show evidence of progressive disease while on therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RINVOQ (S)

Products Affected

- Rinvoq

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

RINVOQ LQ (S)

Products Affected

- Rinvoq Lq

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

RIVFLOZA (S)

Products Affected

- Rivfloza

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

ROMVIMZA (S)

Products Affected

- Romvimza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of tenosynovial giant cell tumor (TGCT). Patient is symptomatic. Surgical resection will potentially cause worsening functional limitation or severe morbidity.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ROZLYTREK (S)

Products Affected

- Rozlytrek

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of metastatic (NSCLC). Presence of ROS1 rearrangement as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Solid Tumors: Diagnosis of solid tumors. Presence of neurotrophic tyrosine receptor kinase (NTRK) gene fusion (e.g., ETV6-NTRK3, TPM3-NTRK1, TPR-NTRK1, etc.) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). No 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). One of the following: disease has progressed following previous treatment (e.g., surgery, radiation therapy, or systemic therapy) or disease has no satisfactory alternative treatments.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RUBRACA (S)

Products Affected

- Rubraca

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Ovarian cancer: Diagnosis of epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer. Prostate cancer: Diagnosis of castration-resistant prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RYBELSUS (S)

Products Affected

- Rybelsus TABS 14MG, 3MG, 7MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: One of the following: a) For patients requiring ongoing treatment for type 2 diabetes mellitus (T2DM), submission of medical records (e.g., chart notes) confirming diagnosis of T2DM, OR b) Submission of medical records (e.g., chart notes) confirming diagnosis of T2DM as evidenced by one of the following laboratory values: i) A1c greater than or equal to 6.5%, ii) fasting plasma glucose (FPG) greater than or equal to 126 mg/dL, or iii) 2-hour plasma glucose (PG) greater than or equal to 200 mg/dL during OGTT (oral glucose tolerance test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RYDAPT (S)

Products Affected

- Rydapt

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute Myeloid Leukemia (AML): Newly diagnosed acute myeloid leukemia (AML), Disease is confirmed by the presence of FMS-like tyrosine kinase 3 (FLT3) mutation-positive as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA), used in combination with standard cytarabine and daunorubicin induction and cytarabine consolidation. 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SABRIL (S)

Products Affected

- Vigabatrin
- Vigadrone

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Complex Partial Seizures (CPS): For use as adjunctive therapy. Failure, contraindication, or intolerance to two formulary anticonvulsants [eg, Lamictal (lamotrigine), Depakene (valproic acid), Dilantin (phenytoin)]. Infantile Spasms (IS): Diagnosis of infantile spasms.
Age Restrictions	IS: 1 month to 2 years of age. CPS: 2 years or older.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SANDOSTATIN (S)

Products Affected

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

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

SCEMBLIX (S)

Products Affected

- Scemblix

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of chronic myelogenous/myeloid leukemia (CML). Presence of a Philadelphia chromosome-positive (Ph+) as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease is in chronic phase.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SIGNIFOR (S)

Products Affected

- Signifor

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cushing's disease (initial): Diagnosis of Cushing's disease. One of the following: a) Pituitary surgery has not been curative for the patient or b) Patient is not a candidate for pituitary surgery.
Age Restrictions	N/A
Prescriber Restrictions	Cushing's disease (initial): Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Cushing's disease (initial, reauth): 12 months
Other Criteria	Cushing's disease (reauth): Patient demonstrates positive clinical response to therapy (e.g., a clinically meaningful reduction in 24-hour urinary free cortisol levels, improvement in signs or symptoms of the disease).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SKYCLARYS (S)

Products Affected

- Skyclarys

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

SKYRIZI (S)

Products Affected

- Skyrizi INJ 150MG/ML, 180MG/1.2ML, 360MG/2.4ML, 55MG/0.37ML
- Skyrizi Pen

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

SKYRIZI IV (S)

Products Affected

- Skyrizi INJ 600MG/10ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's disease (CD): Diagnosis 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. Will be administered as an intravenous induction dose. Ulcerative Colitis (UC): 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. Will be administered as an intravenous induction dose.
Age Restrictions	N/A
Prescriber Restrictions	UC, CD: Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	UC, CD: 3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SOMAVERT (S)

Products Affected

- Somavert

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

SPEVIGO (S)

Products Affected

- Spevigo INJ 150MG/ML, 300MG/2ML

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

SPRYCEL (S)

Products Affected

- Dasatinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Philadelphia chromosome positive (Ph+)/BCR ABL chronic myelogenous leukemia (CML): Diagnosis of Ph+/BCR ABL CML. Ph+/BCR ABL acute lymphoblastic leukemia (ALL): Diagnosis of Ph+/BCR ABL ALL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All Uses: 12 months
Other Criteria	All Uses: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

STARJEMZA (IV) (S)

Products Affected

- Starjemza INJ 130MG/26ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's Disease (CD): Diagnosis 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. Ulcerative Colitis (UC): 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. CD, UC: claims or submission of medical records confirming trial of either Steqeyma or Wezlana.
Age Restrictions	N/A
Prescriber Restrictions	All uses: Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	All uses: One time
Other Criteria	All uses: Will be administered as an intravenous induction dose. Induction dosing is in accordance with the United States Food and Drug Administration approved labeled dosing for Crohn's Disease/ulcerative colitis: 260 mg for patients weighing 55 kg or less, 390 mg for patients weighing more than 55 kg to 85 kg, or 520 mg for patients weighing more than 85 kg.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

STARJEMZA (S)

Products Affected

- Starjemza INJ 45MG/0.5ML, 90MG/ML

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

STEQEYMA (S)

Products Affected

- Steqeyma INJ 45MG/0.5ML, 90MG/ML

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

STEQEYMA IV (S)

Products Affected

- Steqeyma INJ 130MG/26ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's Disease (CD): Diagnosis 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. Ulcerative Colitis (UC): 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.
Age Restrictions	N/A
Prescriber Restrictions	All uses: Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	All uses: One time
Other Criteria	All uses: Will be administered as an intravenous induction dose. Induction dosing is in accordance with the United States Food and Drug Administration approved labeled dosing for Crohn's Disease/ulcerative colitis: 260 mg for patients weighing 55 kg or less, 390 mg for patients weighing more than 55 kg to 85 kg, or 520 mg for patients weighing more than 85 kg.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

STIVARGA (S)

Products Affected

- Stivarga

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Metastatic colorectal cancer (mCRC): Diagnosis of mCRC. Gastrointestinal stromal tumor (GIST): Diagnosis of locally advanced, unresectable or metastatic GIST. Hepatocellular Carcinoma (HCC): Diagnosis of HCC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

STRENSIQ (S)

Products Affected

- Strensiq

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Hypophosphatasia (initial): Diagnosis of perinatal/infantile or juvenile-onset hypophosphatasia. One of the following: 1) Both of the following: a) Patient has low level activity of serum alkaline phosphatase (ALP) evidenced by an ALP level below the age and gender-adjusted normal range AND b) Patient has an elevated level of tissue non-specific alkaline phosphatase (TNSALP) substrate (e.g., serum pyridoxal 5'-phosphate [PLP] level, serum or urine phosphoethanolamine [PEA] level, urinary inorganic pyrophosphate [PPi level]) OR 2) Confirmation of tissue-nonspecific alkaline phosphatase (TNSALP) gene mutation by ALPL genomic DNA testing. For patients requesting the 80 mg/0.8 mL vial only: Patient's weight is greater than or equal to 40 kg.
Age Restrictions	N/A
Prescriber Restrictions	Hypophosphatasia (initial, reauth): Prescribed by a specialist experienced in the treatment of inborn errors of metabolism or an endocrinologist
Coverage Duration	Hypophosphatasia (initial): 6 months, (reauth): 12 months
Other Criteria	Hypophosphatasia (reauth): Patient demonstrates positive clinical response to therapy. For patients requesting the 80 mg/0.8 mL vial only: Patient's weight is greater than or equal to 40 kg.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SUCRAID (S)

Products Affected

- Sucraid

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Congenital Sucrase-Isomaltase Deficiency (CSID) (initial): Diagnosis of sucrase deficiency (which is part of congenital sucrose-isomaltase deficiency [CSID]).
Age Restrictions	CSID (initial): Patient is 5 months of age or older.
Prescriber Restrictions	CSID (initial): Prescribed by or in consultation with a gastroenterologist or geneticist.
Coverage Duration	CSID (initial, reauth): 12 months.
Other Criteria	CSID (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SUTENT (S)

Products Affected

- Sunitinib Malate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Renal cell carcinoma: Diagnosis of advanced or metastatic renal cell carcinoma. Gastrointestinal stromal tumor (GIST): Diagnosis of GIST after disease progression on, or contraindication or intolerance to Gleevec (imatinib). Pancreatic neuroendocrine tumors: Diagnosis of progressive, well-differentiated pancreatic neuroendocrine tumor that is unresectable locally advanced or metastatic disease. Adjuvant treatment of renal cell carcinoma: Diagnosis of renal cell carcinoma (RCC). Used as adjuvant therapy. Patient is at high risk of recurrent RCC following nephrectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	All Indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SYPRINE (S)

Products Affected

- Trientine Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of Wilson's disease (i.e., hepatolenticular degeneration). Trial and failure, contraindication, or intolerance to a penicillamine product (e.g., Depen, Cuprimine)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TABRECTA (S)

Products Affected

- Tabrecta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-small cell lung cancer (NSCLC). Disease is metastatic. Presence of mesenchymal-epithelial transition (MET) exon 14 skipping positive tumors as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months.
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAFAMIDIS (S)

Products Affected

- Vyndamax

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Transthyretin-mediated amyloidosis with cardiomyopathy (ATTR-CM) (initial): Diagnosis of transthyretin-mediated amyloidosis with cardiomyopathy (ATTR-CM). One of the following: 1) Presence of a transthyretin (TTR) mutation (e.g., V122I) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA), 2) Cardiac or noncardiac tissue biopsy demonstrating histologic confirmation of TTR amyloid deposits, OR 3) Both of the following: i) echocardiogram or cardiac magnetic resonance imaging or scintigraphy scan suggestive of amyloidosis, and ii) absence of light-chain amyloidosis. Patient has New York Heart Association (NYHA) Functional Class I, II, or III heart failure. Requested drug is not used in combination with a TTR silencer (e.g., Amvuttra) or a TTR stabilizer (e.g., Diflunisal).
Age Restrictions	N/A
Prescriber Restrictions	ATTR-CM (initial, reauth): Prescribed by or in consultation with a cardiologist
Coverage Duration	ATTR-CM (initial, reauth): 12 months
Other Criteria	ATTR-CM (reauth): Patient continues to have New York Heart Association (NYHA) Functional Class I, II, or III heart failure. Requested drug is not used in combination with a TTR silencer (e.g., Amvuttra) or a TTR stabilizer (e.g., Diflunisal).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAFINLAR (S)

Products Affected

- Tafinlar

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable or metastatic melanoma. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA) OR both of the following: Presence of BRAF V600E or V600K mutation as detected by an FDA-approved test or a test performed at a facility approved by CLIA and medication is used in combination with Mekinist (trametinib). Adjuvant Treatment for Melanoma: Diagnosis of melanoma. Presence of BRAF V600E or V600K mutation as detected by an FDA-approved test or a test performed at a facility approved by CLIA. 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): Diagnosis of metastatic non-small cell lung cancer. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by CLIA. Medication is used in combination with Mekinist (trametinib). Anaplastic Thyroid Cancer (ATC): Diagnosis of locally advanced or metastatic anaplastic thyroid cancer. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by CLIA. Cancer may not be treated with standard locoregional treatment options. Medication is used in combination with Mekinist (trametinib) .
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months

Other Criteria	Approve for continuation of prior therapy. Solid tumors: Diagnosis of solid tumors. Disease is unresectable or metastatic. Patient has progressed on or following prior treatment and have no satisfactory alternative treatment options. Presence of BRAF V600E mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is used in combination with Mekinist (trametinib). Low-grade Glioma: Diagnosis of low-grade glioma. Patient requires systemic therapy. Presence of BRAF V600E mutation type as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is used in combination with Mekinist (trametinib).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TAGRISSO (S)

Products Affected

- Tagrisso

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): One of the following: A) All of the following: Diagnosis of metastatic NSCLC. One of the following: 1) Disease is confirmed by known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA), OR 2) Both of the following: a) Disease is confirmed by known active EGFR T790M mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA) and b) Patient has experienced disease progression on or after one of the following EGFR Tyrosine Kinase Inhibitors (TKIs): Gilotrif (afatinib), Iressa (gefitinib), Tarceva (erlotinib), or Vizimpro (dacomitinib). OR B) All of the following: Diagnosis of NSCLC. Disease is confirmed by known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Both of the following: 1) Patient is receiving as adjuvant therapy, and 2) Patient has had a complete surgical resection of the primary NSCLC tumor. OR C) All of the following: Diagnosis of NSCLC. Disease is locally advanced. Disease is confirmed by known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations as detected by an U.S. FDA-approved test or a test performed at a facility approved by CLIA. Used in combination with both of the following: a) Pemetrexed, and b) Platinum-based chemotherapy (e.g., cisplatin, carboplatin). Or D) Refer to Other Criteria element for additional indication and criteria.
Age Restrictions	N/A

Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	NSCLC cont: OR D) All of the following: Diagnosis of NSCLC. Disease is Locally advanced, Unresectable (Stage III). Disease is confirmed by the presence of known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations as detected by an U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Disease has not progressed during or following concurrent or sequential platinum-based chemoradiation therapy. All indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TALZENNA (S)

Products Affected

- Talzenna

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Breast cancer: Diagnosis of breast cancer. Prostate cancer: Diagnosis of metastatic castration-resistant prostate cancer (mCRPC). Presence of homologous recombination repair (HRR) gene-mutation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with Xtandi (enzalutamide).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TARCEVA (S)

Products Affected

- Erlotinib Hydrochloride TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of locally advanced or metastatic (Stage III or IV) NSCLC AND Disease is confirmed by known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Pancreatic Cancer: Diagnosis of locally advanced, unresectable, or metastatic pancreatic cancer AND erlotinib will be used in combination with gemcitabine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	All uses: 12 months
Other Criteria	All Indications: Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TARGRETIN (S)

Products Affected

- Bexarotene

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cutaneous T-Cell Lymphoma (CTCL): Diagnosis of CTCL. Trial and failure, contraindication, or intolerance to at least one prior therapy (including skin-directed therapies [eg, corticosteroids (ie, clobetasol, diflorasone, halobetasol, augmented betamethasone dipropionate)] or systemic therapies [eg, brentuximab vedotin, methotrexate]).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TARPEYO (S)

Products Affected

- Tarpeyo

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

TASIGNA (S)

Products Affected

- Nilotinib Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic myelogenous leukemia (CML): Diagnosis of Ph+/BCR ABL CML
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAVNEOS (S)

Products Affected

- Tavneos

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of one of the following types of severe active anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis: a) Granulomatosis with polyangiitis (GPA) OR b) Microscopic polyangiitis (MPA). Diagnosis is confirmed by one of the following: a) ANCA test positive for proteinase 3 (PR3) antigen, b) ANCA test positive for myeloperoxidase (MPO) antigen, OR c) Tissue biopsy. Patient is receiving concurrent immunosuppressant therapy with one of the following: a) cyclophosphamide OR b) rituximab. One of the following: a) Patient is concurrently on glucocorticoids (e.g., prednisone) OR b) History of contraindication or intolerance to glucocorticoids (e.g., prednisone).
Age Restrictions	N/A
Prescriber Restrictions	Initial, Reauth: Prescribed by or in consultation with a nephrologist, pulmonologist, or rheumatologist
Coverage Duration	Initial, Reauth: 12 months
Other Criteria	Reauth: Patient does not show evidence of progressive disease while on therapy. Patient is receiving concurrent immunosuppressant therapy (e.g., azathioprine, cyclophosphamide, methotrexate, rituximab).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TAZVERIK (S)

Products Affected

- Tazverik

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 follicular lymphoma. Disease is one of the following: relapsed or refractory.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TECFIDERA (S)

Products Affected

- Dimethyl Fumarate CPDR
- Dimethyl Fumarate Starterpack CDPK 0

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Multiple Sclerosis (MS) (initial, reauth): Not used in combination with another disease-modifying therapy for MS.
Required Medical Information	MS (initial): Diagnosis of a relapsing form of MS (eg, clinically isolated syndrome, relapsing-remitting disease, secondary progressive disease, including active disease with new brain lesions).
Age Restrictions	N/A
Prescriber Restrictions	MS (initial, reauth): Prescribed by or in consultation with a neurologist
Coverage Duration	MS (initial, reauth): 12 months
Other Criteria	MS (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TEPMETKO (S)

Products Affected

- Tepmetko

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is metastatic. Presence of mesenchymal-epithelial transition (MET) exon 14 skipping alterations as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TERIPARATIDE (S)

Products Affected

- Bonsity
- Teriparatide

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

TESTOSTERONE (S)

Products Affected

- Testosterone GEL 25MG/2.5GM, 50MG/5GM
- Testosterone Cypionate INJ 100MG/ML, 200MG/ML
- Testosterone Pump GEL 12.5MG/ACT, 20.25MG/ACT

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Hypogonadism (HG) (Initial): Diagnosis (dx) of HG AND male patient at birth AND one of the following: 1) Two pre-treatment serum total testosterone (T) levels less than 300 ng/dL (10.4 nmol/L) or less than the reference range for the lab, OR 2) Both of the following: a) Has a condition that may cause altered sex-hormone binding globulin (SHBG) (eg, thyroid disorder, HIV disease, liver disorder, diabetes, obesity), and b) one pre-treatment calculated free or bioavailable T level less than 5 ng/dL (0.17 nmol/L) or less than reference range for the lab, OR 3) History of bilateral orchiectomy, panhypopituitarism, or a genetic disorder known to cause HG (eg, congenital anorchia, Klinefelter's syndrome), OR 4) Both of the following: a) Patient is continuing testosterone therapy, and b) One of the following: i) Follow-up total serum T level or calculated free or bioavailable T level drawn within the past 12 months is within or below the normal limits of the reporting lab, or ii) follow-up total serum T level or calculated free or bioavailable T level drawn within the past 12 months is outside of upper limits of normal for the reporting lab and the dose is adjusted. Gender Dysphoria (GD)/Gender Incongruence (off-label): Dx of GD/Gender Incongruence. Using hormones to change characteristics to align with gender expression.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	HG(init): (New to T tx:6 mo. New to plan and cont T tx:12 mo), (reauth): 12 mo. GD: 12 mo.

Other Criteria	HG (Reauth): 1) Follow-up total serum T level within or below the normal limits of the reporting lab, or 2) Follow-up total serum T level outside of upper limits of normal for the reporting lab and the dose is adjusted, OR 3) Has a condition that may cause altered SHBG (eg, thyroid disorder, HIV disease, liver disorder, diabetes, obesity), and one of the following: Follow-up calculated free or bioavailable T level within or below the normal limits of the reporting lab, or follow-up calculated free or bioavailable T level outside of upper limits of normal for the reporting lab and the dose is adjusted.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TESTOSTERONE ENANTHATE (S)

Products Affected

- Testosterone Enanthate INJ

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Hypogonadism (HG) (Initial): Diagnosis (dx) of HG AND male patient at birth AND one of the following: 1) Two pre-treatment serum total testosterone (T) levels less than 300 ng/dL (10.4 nmol/L) or less than the reference range for the lab, OR 2) Both of the following: a) Has a condition that may cause altered sex-hormone binding globulin (SHBG) (eg, thyroid disorder, HIV disease, liver disorder, diabetes, obesity), and b) one pre-treatment calculated free or bioavailable T level less than 5 ng/dL (0.17 nmol/L) or less than reference range for the lab, OR 3) History of bilateral orchiectomy, panhypopituitarism, or a genetic disorder known to cause HG (eg, congenital anorchia, Klinefelter's syndrome), OR 4) Both of the following: a) Patient is continuing testosterone therapy, and b) One of the following: i) Follow-up total serum T level or calculated free or bioavailable T level drawn within the past 12 months is within or below the normal limits of the reporting lab, or ii) follow-up total serum T level or calculated free or bioavailable T level drawn within the past 12 months is outside of upper limits of normal for the reporting lab and the dose is adjusted. Delayed puberty (DP): Dx of DP AND male patient at birth. Breast cancer (BC): Dx of inoperable BC AND used for palliative treatment AND female patient at birth. Gender Dysphoria (GD)/Gender Incongruence (off-label): Dx of GD/Gender Incongruence. Using hormones to change characteristics to align with gender expression.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	HG(init): (New to T tx:6 mo. Cont T tx:12 mo), (reauth): 12 mo. BC, GD: 12 mo. DP: 6 mo.

Other Criteria	HG (Reauth): 1) Follow-up total serum T level within or below the normal limits of the reporting lab, or 2) Follow-up total serum T level outside of upper limits of normal for the reporting lab and the dose is adjusted, OR 3) Has a condition that may cause altered SHBG (eg, thyroid disorder, HIV disease, liver disorder, diabetes, obesity), and one of the following: Follow-up calculated free or bioavailable T level within or below the normal limits of the reporting lab, or follow-up calculated free or bioavailable T level outside of upper limits of normal for the reporting lab and the dose is adjusted.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TEVIMBRA (S)

Products Affected

- Tevimbra

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Esophageal Squamous Cell Carcinoma: Diagnosis of esophageal squamous cell carcinoma. Disease is unresectable or metastatic. One of the following: 1) All of the following: a) Patient has received prior systemic chemotherapy, b) Used as a single agent, and c) Patient has not previously been treated with a PD-(L)1 inhibitor (e.g., Keytruda, Opdivo) OR 2) Both of the following: a) Used in combination with platinum-containing chemotherapy (e.g., carboplatin, cisplatin, oxaliplatin) and b) Tumor(s) express PD-L1 (greater than or equal to 1) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Gastric or Gastroesophageal Junction Adenocarcinoma: Diagnosis of gastric or gastroesophageal junction adenocarcinoma. Disease is unresectable or metastatic. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Tumor(s) express PD-L1 (greater than or equal to 1) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Used in combination with platinum (e.g., carboplatin, cisplatin, oxaliplatin) and fluoropyrimidine (e.g., fluorouracil)-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

THALOMID (S)

Products Affected

- Thalomid CAPS 100MG, 50MG

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

TIBSOVO (S)

Products Affected

- Tibsovo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Relapsed or refractory Acute Myeloid Leukemia (AML): Diagnosis of AML. Disease is relapsed or refractory. Newly-Diagnosed AML: Diagnosis of newly-diagnosed AML. One of the following: 1) patient is greater than or equal to 75 years old OR 2) patient has comorbidities that preclude use of intensive induction chemotherapy. Locally Advanced or Metastatic Cholangiocarcinoma: Diagnosis of cholangiocarcinoma. Disease is locally advanced or metastatic. Patient has been previously treated. Relapsed or Refractory Myelodysplastic Syndromes: Diagnosis of myelodysplastic syndromes (MDS). Disease is relapsed or refractory. All indications: Presence of an isocitrate dehydrogenase-1 (IDH1) mutation as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TOFACITINIB (S)

Products Affected

- Tofacitinib Citrate SOLN
- Tofacitinib Citrate TABS
- Tofacitinib Citrate Er

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

TOPICAL RETINOID (S)

Products Affected

- Tretinoin CREA 0.025%, 0.05%

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acne vulgaris: Diagnosis of acne vulgaris (i.e., acne).
Age Restrictions	PA applies to members 26 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRACLEER (S)

Products Affected

- Bosentan TABS

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

TRELSTAR (S)

Products Affected

- Trelstar Mixject INJ 11.25MG, 22.5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prostate Cancer: Diagnosis of advanced or metastatic prostate cancer. Trial and failure, contraindication, or intolerance to any brand Lupron formulation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TREMFYA INTRAVENOUS (S)

Products Affected

- Tremfya INJ 200MG/20ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's Disease (CD): Diagnosis 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, C-reactive protein [CRP]), OR CD Activity Index (CDAI) greater than 220. Will be administered as an intravenous induction dose. Ulcerative Colitis (UC): 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, erythrocyte sedimentation rate [ESR], CRP), OR dependent on, or refractory to, corticosteroids. Will be administered as an intravenous induction dose.
Age Restrictions	N/A
Prescriber Restrictions	CD, UC: Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	CD, UC: 3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TREMFYA SC (S)

Products Affected

- Tremfya INJ 100MG/ML, 200MG/2ML
- Tremfya Induction Pack For Crohns Disease/ulcerative Colitis
- Tremfya Pen

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

TRIKAFTA (S)

Products Affected

- Trikafta TBPK

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

TRULICITY (S)

Products Affected

- Trulicity

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: One of the following: a) For patients requiring ongoing treatment for type 2 diabetes mellitus (T2DM), submission of medical records (e.g., chart notes) confirming diagnosis of T2DM, OR b) Submission of medical records (e.g., chart notes) confirming diagnosis of T2DM as evidenced by one of the following laboratory values: i) A1c greater than or equal to 6.5%, ii) fasting plasma glucose (FPG) greater than or equal to 126 mg/dL, or iii) 2-hour plasma glucose (PG) greater than or equal to 200 mg/dL during OGTT (oral glucose tolerance test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRUQAP (S)

Products Affected

- Truqap

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

TRYNGOLZA (S)

Products Affected

- Tryngolza

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

TUKYSA (S)

Products Affected

- Tukysa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 received one or more prior anti-HER2 based regimens (e.g., trastuzumab, pertuzumab, ado-trastuzumab emtansine).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TURALIO (S)

Products Affected

- Turalio CAPS 125MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TYENNE SC (S)

Products Affected

- Tyenne INJ 162MG/0.9ML

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

TYKERB (S)

Products Affected

- Lapatinib Ditosylate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Metastatic Breast Cancer: Diagnosis of human epidermal growth factor receptor 2 (HER2)-positive breast cancer. One of the following: 1) All of the following: a) Disease is one of the following: Advanced OR Metastatic, b) All of the following: i) Patient has received prior therapy with an anthracycline, ii) Patient has received prior therapy with a taxane, AND iii) Patient has experienced disease progression while on trastuzumab therapy, AND c) Used in combination with Xeloda (capecitabine), or 2) Both of the following: a) Disease is BOTH of the following: Metastatic AND Hormone receptor positive (HR+), AND b) Used in combination with Femara (letrozole).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TYMLOS (S)

Products Affected

- Tymlos

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

UBRELVY (S)

Products Affected

- Ubrelvy

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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	Initial: 18 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Initial, Reauth: 12 months.
Other Criteria	Reauth: Patient has experienced a positive 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 Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

UDENYCA (S)

Products Affected

- Udenyca

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

UDENYCA ONBODY (S)

Products Affected

- Udenyca Onbody

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Febrile neutropenia (FN) prophylaxis: Patient will be receiving prophylaxis for FN due to one of the following: 1) Patient is receiving National Cancer Institute's Breast Intergroup, INT C9741 dose dense chemotherapy protocol for primary breast cancer, 2) patient is receiving a dose-dense chemotherapy regimen for which the incidence of FN is unknown, 3) patient is receiving chemotherapy regimen(s) associated with greater than 20% incidence of FN, 4) both of the following: a) patient is receiving chemotherapy regimen(s) associated with 10-20% incidence of FN, AND b) patient has one or more risk factors associated with chemotherapy-induced infection, FN, or neutropenia, OR 5) Both of the following: a) patient is receiving myelosuppressive anticancer drugs associated with neutropenia, AND b) patient has a history of FN or dose-limiting event during a previous course of chemotherapy (secondary prophylaxis). Treatment of FN (off-label): Patient has received or is receiving myelosuppressive anticancer drugs associated with neutropenia. Diagnosis of FN. Patient is at high risk for infection-associated complications.
Age Restrictions	N/A
Prescriber Restrictions	All uses: Prescribed by or in consultation with a hematologist/oncologist
Coverage Duration	FN (prophylaxis, treatment): 3 mo or duration of tx.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VALCHLOR (S)

Products Affected

- Valchlor

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Mycosis fungoides-type cutaneous T-cell lymphoma (MF-CTCL) (initial): All 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 skin-directed therapy (e.g., topical corticosteroids [e.g., clobetasol, fluocinonide], bexarotene topical gel [Targretin topical gel], etc.).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VANFLYTA (S)

Products Affected

- Vanflyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute Myeloid Leukemia (AML): Diagnosis of AML. Patient has a FMS-like tyrosine kinase 3 (FLT3) internal tandem duplication (FLT3-ITD) mutation as detected by a U.S. Food and Drug Administration (FDA)-approved test (e.g., LeukoStrat CDx FLT3 Mutation Assay) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Both of the following: a) Used in combination with standard cytarabine and anthracycline (e.g., daunorubicin, idarubicin) induction and cytarabine consolidation, and b) Used as maintenance monotherapy following consolidation chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VENCLEXTA (S)

Products Affected

- Venclexta
- Venclexta Starting Pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL): Diagnosis of CLL or SLL. Acute Myeloid Leukemia (AML): Diagnosis of newly diagnosed AML. Used in combination with azacitidine, or decitabine, or low-dose cytarabine. One of the following: 1) age 75 years or older OR 2) comorbidities that preclude use of intensive induction chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VEOPOZ (S)

Products Affected

- Veopoz

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of active CD55-deficient protein-losing enteropathy (PLE), also known as CHAPLE disease. Disease is confirmed by the presence of biallelic CD55 loss-of-function mutation genotype as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has hypoalbuminemia (serum albumin concentration of less than or equal to 3.2 g/dL). Patient has at least one of the following signs or symptoms within the last six months: abdominal pain, diarrhea, peripheral edema, or facial edema.
Age Restrictions	Initial: Patient is 1 year of age or older.
Prescriber Restrictions	Initial: Prescribed by or in consultation with an immunologist, geneticist, hematologist, or gastroenterologist.
Coverage Duration	Initial, Reauth: 12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VEOZAH (S)

Products Affected

- Veozah

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	Initial, Reauth: 6 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy. One of the following within the past 3 months: a) For patients with total bilirubin levels less than or equal to 2 times the ULN, transaminase elevations do not exceed 5 times the ULN, or b) For patients with total bilirubin levels greater than 2 times the ULN, transaminase elevations do not exceed 3 times the ULN.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VEPPANU (S)

Products Affected

- Veppanu

PA Criteria	Criteria Details
Indications	N/A
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	N/A
Other Criteria	N/A
Prerequisite Therapy Required	

VERQUVO (S)

Products Affected

- Verquvo

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

VERZENIO (S)

Products Affected

- Verzenio

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Breast Cancer: Diagnosis of breast cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VIGAFYDE (S)

Products Affected

- Vigafyde

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of infantile spasms. Trial or intolerance to generic vigabatrin.
Age Restrictions	Patient is 1 month to 2 years of age.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VITRAKVI (S)

Products Affected

- Vitrakvi

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

VIZIMPRO (S)

Products Affected

- Vizimpro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of NSCLC. Disease is metastatic. Presence of epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VONJO (S)

Products Affected

- Vonjo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of ONE of the following: a) Primary myelofibrosis, b) Post-polycythemia vera myelofibrosis, OR c) Post-essential thrombocythemia myelofibrosis. Disease is intermediate or high risk. Pre-treatment platelet count below 50 x 10^9/L.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VORANIGO (S)

Products Affected

- Voranigo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of astrocytoma or oligodendroglioma. Presence of a susceptible isocitrate dehydrogenase-1 (IDH1) or isocitrate dehydrogenase-2 (IDH2) mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). History of one of the following: a) Biopsy, b) Sub-total resection, or c) Gross total resection.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VORICONAZOLE INJECTION (S)

Products Affected

- Voriconazole INJ 200MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 Restrictions	N/A
Coverage Duration	12 weeks
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOSEVI (S)

Products Affected

- Vosevi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 not used in combination with another HCV direct acting antiviral agent [e.g., Harvoni, Zepatier].
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with one of the following: Hepatologist, Gastroenterologist, Infectious disease specialist, HIV specialist certified through the American Academy of HIV Medicine, Liver transplant specialist.
Coverage Duration	12 to 24 weeks. Criteria will be applied consistent with current AASLD/IDSA guideline.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VOTRIENT (S)

Products Affected

- Pazopanib Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Renal cell carcinoma (RCC): Diagnosis of advanced/metastatic RCC. Soft tissue sarcoma: Diagnosis of advanced soft tissue sarcoma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOWST (S)

Products Affected

- Vowst

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

VYJUVEK (S)

Products Affected

- Vyjuvek

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of dystrophic epidermolysis bullosa (DEB). Presence of collagen type VII alpha 1 chain (COL7A1) gene mutation as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Medication is being used for the treatment of wounds that require healing. One of the following: a) Medication will be applied by a healthcare professional or b) Patient or caregiver has received training for the application of Vyjuvek. Wound(s) being treated has no signs of infection. Medication is not being used concurrently with other FDA approved therapies (e.g., Filsuvez) for the treatment of epidermolysis bullosa.
Age Restrictions	N/A
Prescriber Restrictions	Initial, Reauth: Prescribed by or in consultation with a dermatologist or specialist with expertise in wound care.
Coverage Duration	Initial, Reauth: 12 months.
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy. Wound(s) being treated has no signs of infection. Medication is not being used concurrently with other FDA approved therapies (e.g., Filsuvez) for the treatment of epidermolysis bullosa.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VYVGART (S)

Products Affected

- Vyvgart Hytrulo

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

WEGOVY (S)

Products Affected

- Wegovy INJ

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

WEGOVY TABLETS (S)

Products Affected

- Wegovy TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Excluded if used for weight loss only.
Required Medical Information	Initial: Treatment is being requested to reduce the risk of major adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke). Submission of medical records (e.g., chart notes) documenting that patient has established cardiovascular disease as evidenced by one of the following: A) prior myocardial infarction (MI), B) prior stroke, or C) peripheral arterial disease. BMI greater than or equal to 27 kg/m2. Medication is not being co-administered with another GLP-1 receptor agonist (e.g., Trulicity, Mounjaro, Ozempic, Rybelsus).
Age Restrictions	Initial: Patient is 18 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Initial, Reauth: 12 months.
Other Criteria	Reauth: Medication is not being co-administered with another GLP-1 receptor agonist (e.g., Trulicity, Mounjaro, Ozempic, Rybelsus).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

WELIREG (S)

Products Affected

- Welireg

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

WINREVAIR (S)

Products Affected

- Winrevair

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

WYOST (S)

Products Affected

- Wyost

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Skeletal prevention in Multiple Myeloma (MM)/Bone Metastasis from Solid Tumors (BMST): One of the following: 1) Both of the following: a) Diagnosis of multiple myeloma and b) Trial and failure, contraindication (e.g., renal insufficiency), or intolerance to one bisphosphonate therapy, OR 2) Both of the following: a) Diagnosis of solid tumors (e.g., breast cancer, kidney cancer, lung cancer, prostate cancer, thyroid cancer) and b) Documented evidence of one or more metastatic bone lesions. Giant cell tumor of bone (GCTB): Diagnosis of giant cell tumor of bone. One of the following: 1) One of the following: a) tumor is unresectable or b) surgical resection is likely to result in severe morbidity, OR 2) Approve for continuation of prior therapy. Hypercalcemia of malignancy (HCM): Diagnosis of hypercalcemia of malignancy. Trial and failure, contraindication, or intolerance to one bisphosphonate therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	MM/BMST, GCTB, HCM: 12 months.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XALKORI (S)

Products Affected

- Xalkori

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small cell lung cancer (NSCLC): Diagnosis of metastatic NSCLC. One of the following: a) Patient has an anaplastic lymphoma kinase (ALK)-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA) or b) Patient has ROS1 rearrangement-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Anaplastic Large Cell Lymphoma (ALCL): Diagnosis of systemic ALCL. Disease is relapsed or refractory. Patient has an anaplastic lymphoma kinase (ALK)-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Inflammatory Myofibroblastic Tumor (IMT): Diagnosis of IMT. Disease is one of the following: a) unresectable, b) recurrent, or c) refractory. Patient has an anaplastic lymphoma kinase (ALK)-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	IMT, ALCL: Patient is 1 year of age or older.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XATMEP (S)

Products Affected

- Xatmep

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute lymphoblastic leukemia (ALL): Diagnosis of acute lymphoblastic leukemia (ALL). Polyarticular juvenile idiopathic arthritis (pJIA): Diagnosis of polyarticular juvenile idiopathic arthritis.
Age Restrictions	N/A
Prescriber Restrictions	pJIA: Prescribed by or in consultation with a rheumatologist.
Coverage Duration	ALL, pJIA: 12 months.
Other Criteria	Subject to Part B vs D review. Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XCOPRI (S)

Products Affected

- Xcopri

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of partial onset seizures.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XENAZINE (S)

Products Affected

- Tetrabenazine

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

XERMELO (S)

Products Affected

- Xermelo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical 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 [Sandostatin, Sandostatin LAR], lanreotide [Somatuline Depot]) for at least 3 months AND used in combination with SSA therapy.
Age Restrictions	N/A
Prescriber Restrictions	Initial: Prescribed by or in consultation with an oncologist, endocrinologist, or gastroenterologist
Coverage Duration	Initial: 6 months. Reauth: 12 months
Other Criteria	Carcinoid syndrome diarrhea (Reauthorization): Patient demonstrates positive clinical response to therapy AND drug will continue to be used in combination with SSA therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XIFAXAN (S)

Products Affected

- Xifaxan

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

XOLAIR (S)

Products Affected

- Xolair

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Asthma (init): Diagnosis of moderate to severe persistent allergic asthma. Positive skin test or in vitro reactivity to a perennial aeroallergen. One of the following: A) All of the following: a) Patient is 6 years of age or older but less than 12 years of age, b) Pretreatment serum immunoglobulin (Ig)E level between 30 to 1300 IU/mL, c) Patient is currently being treated with one of the following unless there is a contraindication or intolerance to these medications: i) Both of the following: 1) Medium-dose inhaled corticosteroid [ICS] (eg, greater than 100-200 mcg fluticasone propionate equivalent/day), and 2) 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 B) All of the following: a) Patient is 12 years of age or older, b) Treatment serum immunoglobulin (Ig)E level between 30 to 700 IU/mL, c) Patient is currently being treated with one of the following unless there is a contraindication or intolerance to these medications: i) Both of the following: 1) High-dose ICS [eg, greater than 500 mcg fluticasone propionate equivalent/day], and 2) Additional asthma controller medication (eg, LTRA [eg, montelukast], LABA [eg, salmeterol], LAMA [eg, tiotropium]), OR ii) One maximally-dosed combination ICS/LABA product [eg, Wixela Inhub (fluticasone propionate 500mcg/salmeterol 50mcg), budesonide 160mcg/ formoterol 4.5mcg, Breo Ellipta (fluticasone 200mcg/vilanterol 25mcg)].
Age Restrictions	IgE-Mediated Food Allergy (init): Patient is 1 year of age or older.

Prescriber Restrictions	Asthma (init): Prescribed by or in consultation with an allergist/immunologist, or pulmonologist. CSU (init): Prescribed by or in consultation with an allergist/immunologist, or dermatologist. CRSwNP (init/reauth): Prescribed by or in consultation with an allergist/immunologist, otolaryngologist, or pulmonologist. IgE-Mediated Food Allergy (Init/Reauth): Prescribed by or in consultation with an allergist/immunologist.
Coverage Duration	Asthma, CSU, Allergy (init, reauth): 12 months. CRSwNP: 12 months.
Other Criteria	Asthma (reauth): Patient demonstrates positive clinical response to therapy. Chronic Spontaneous Urticaria (CSU) (init): Diagnosis of CSU. Persistent symptoms (itching and hives) with a second generation H1 antihistamine (eg, cetirizine, fexofenadine), unless there is a contraindication or intolerance to H1 antihistamines. Used concurrently with an H1 antihistamine, unless there is a contraindication or intolerance to H1 antihistamines. CSU (reauth): Patient's disease status has been re-evaluated since the last authorization to confirm the patient's condition warrants continued treatment. Patient has experienced one or both of the following: Reduction in itching severity from baseline or Reduction in the number of hives from baseline. Chronic Rhinosinusitis with Nasal polyps (CRSwNP) (init): Diagnosis of CRSwNP. Unless contraindicated, the patient has had an inadequate response to an intranasal corticosteroid (eg, fluticasone, mometasone). Used in combination with another agent for chronic rhinosinusitis with nasal polyps. CRSwNP (reauth): Patient demonstrates positive clinical response to therapy. Used in combination with another agent for chronic rhinosinusitis with nasal polyps. IgE-Mediated Food Allergy (Initial): One of the following: A) Both of the following: 1) Diagnosis of IgE Mediated Food Allergy as evidenced by one of the following: a) Positive skin prick test (defined as greater than or equal to 4 mm wheal greater than saline control) to food, b) Positive food specific IgE (greater than or equal to 6 kUA/L), c) Positive oral food challenge, defined as experiencing dose-limiting symptoms at a single dose of less than or equal to 300 mg of food protein, AND 2) Clinical history of IgE Mediated Food Allergy, OR B) Provider attestation that patient has a history of severe allergic response, including anaphylaxis, following exposure to one or more foods. Used in conjunction with food allergen avoidance. Baseline (pre-Xolair tx) serum total IgE level is greater than or equal to 30 IU/mL and less than or equal to 1850 IU/mL. Dosing is according to serum total IgE levels and body weight. IgE-Mediated Food Allergy (Reauth): Patient demonstrates positive clinical response to therapy. Used in conjunction with food allergen avoidance. Dosing will continue to be based on body weight and pretreatment total IgE serum levels.

Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XOLREMDI (S)

Products Affected

- Xolremdi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of WHIM (warts, hypogammaglobulinemia, infections and myelokathexis) syndrome. Presence of genotype confirmed variant of CXCR4 as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Patient has an absolute neutrophil count (ANC) less than 500 cells/µL.
Age Restrictions	Initial: Patient is 12 years of age or older.
Prescriber Restrictions	Initial: Prescribed by or in consultation with one of the following: immunologist, hematologist, geneticist, dermatologist, or allergist.
Coverage Duration	Initial: 6 months, Reauth: 12 months
Other Criteria	Reauth: Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XOSPATA (S)

Products Affected

- Xospata

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acute myeloid leukemia (AML). Disease is relapsed or refractory. Patient has a FMS-like tyrosine kinase (FLT3) mutation as determined by a U.S. Food and Drug Administration (FDA)-approved test (e.g., LeukoStrat CDx FLT3 Mutation Assay) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XPOVIO (S)

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 Criteria	N/A
Required Medical Information	Multiple Myeloma (MM), Diffuse large B-cell lymphoma (DLBCL): Diagnosis of one of the following: 1) DLBCL OR 2) Multiple Myeloma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XTANDI (S)

Products Affected

- Xtandi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Castration-resistant prostate cancer (CRPC): Diagnosis of castration-resistant (chemical or surgical) prostate cancer. Metastatic castration-sensitive prostate cancer (M-CSPC): Diagnosis of metastatic castration-sensitive prostate cancer. Non-metastatic castrion-sensitive prostate cancer (nm-CSPC): Diagnosis of non-metastatic castration-sensitive prostate cancer (nmCSPC). Patient has high-risk biochemical recurrence (BCR) defined by a PSA doubling time less than or equal to 9 months and one of the following: A) PSA values greater than or equal to 1 ng/mL if the patient had prior prostatectomy (with or without radiotherapy) OR B) PSA values at least 2 ng/mL above the nadir if the patient had prior radiotherapy only.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XYREM (S)

Products Affected

- Sodium Oxybate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Narcolepsy with cataplexy (Narcolepsy Type 1)(initial): Diagnosis of narcolepsy as confirmed by sleep study (unless the prescriber provides justification confirming that a sleep study would not be feasible), AND symptoms of cataplexy are present, AND symptoms of excessive daytime sleepiness (eg, irrepressible need to sleep or daytime lapses into sleep) are present. Narcolepsy without cataplexy (Narcolepsy Type 2)(initial): Diagnosis of narcolepsy as confirmed by sleep study (unless the prescriber provides justification confirming that a sleep study would not be feasible), AND symptoms of cataplexy are absent, AND symptoms of excessive daytime sleepiness (eg, irrepressible need to sleep or daytime lapses into sleep) are present, AND One of the following: a) patient is under 18 years of age or b) trial and failure, contraindication or intolerance to one of the following: 1) amphetamine-based stimulant (eg, amphetamine, dextroamphetamine), OR 2) methylphenidate-based stimulant.
Age Restrictions	All Indications (initial): Patient is 7 years of age or older.
Prescriber Restrictions	All uses (initial): Prescribed by or in consultation with one of the following: neurologist, psychiatrist, or sleep medicine specialist.
Coverage Duration	All uses (initial): 6 months. All uses (reauth): 12 months
Other Criteria	All uses (Reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

YESINTEK IV (S)

Products Affected

- Yesintek INJ 130MG/26ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Crohn's Disease (CD): Diagnosis 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. Ulcerative Colitis (UC): 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. CD, UC: claims or submission of medical records confirming trial of either Steqeyma or Wezlana.
Age Restrictions	N/A
Prescriber Restrictions	All uses: Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	All uses: One time
Other Criteria	All uses: Will be administered as an intravenous induction dose. Induction dosing is in accordance with the United States Food and Drug Administration approved labeled dosing for Crohn's Disease/ulcerative colitis: 260 mg for patients weighing 55 kg or less, 390 mg for patients weighing more than 55 kg to 85 kg, or 520 mg for patients weighing more than 85 kg.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

YESINTEK SC (S)

Products Affected

- Yesintek INJ 45MG/0.5ML, 90MG/ML

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

YONSA (S)

Products Affected

- Yonsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Prostate Cancer: Diagnosis of castration-resistant (chemical or surgical) prostate cancer. Trial and failure or intolerance to Xtandi (enzalutamide).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

YULITHIRA (S)

Products Affected

- Yulithira

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

ZAVESCA (S)

Products Affected

- Miglustat
- Yargesa

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

ZEJULA (S)

Products Affected

- Zejula TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	First-Line Maintenance Treatment of Homologous Recombination Deficiency (HRD)-Positive Advanced Ovarian Cancer: Diagnosis of one of the following: a) Advanced epithelial ovarian cancer, b) Advanced fallopian tube cancer, or c) Advanced primary peritoneal cancer. Used for maintenance treatment in patients who are in a complete or partial response to first-line platinum-based chemotherapy (e.g., cisplatin, carboplatin). Cancer is associated with HRD-positive status defined by one of the following: a) Presence of deleterious or suspected deleterious BRCA mutation or b) Genomic instability. Maintenance Treatment of Recurrent Germline BRCA-Mutated Ovarian Cancer: Diagnosis of one of the following: a) Recurrent epithelial ovarian cancer, b) Recurrent fallopian tube cancer, or c) Recurrent 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 or suspected deleterious germline BRCA-mutations (gBRCAmut) as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZELBORAF (S)

Products Affected

- Zelboraf

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Melanoma: Diagnosis of unresectable melanoma or metastatic melanoma. Cancer is BRAFV600 mutant type (MT) as detected by a U.S. Food and Drug Administration (FDA)-approved test (eg, cobas 4600 BRAFV600 Mutation Test) or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA). Erdheim-Chester Disease: Diagnosis of Erdheim-Chester disease AND Disease is BRAFV600 mutant type (MT).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	All indications: Approve for continuation of therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZENBEXUS (S)

Products Affected

- Zenbexus

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

ZEPOSIA (S)

Products Affected

- Zeposia
- Zeposia 7-day Starter Pack
- Zeposia Starter Kit

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

ZOKINVY (S)

Products Affected

- Zokinvy

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	One of the following: 1) Diagnosis of Hutchinson-Gilford Progeria Syndrome, OR 2) For treatment of processing-deficient Progeroid Laminopathies with presence of one of the following mutations as detected by a FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA): i) Heterozygous LMNA mutation with progerin-like protein accumulation OR ii) Homozygous or compound heterozygous ZMPSTE24 mutations. Patient has a body surface area of 0.39 m^2 and above.
Age Restrictions	Patient is 12 months of age or older.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZOLINZA (S)

Products Affected

- Zolinza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cutaneous T-cell lymphoma (CTCL): Diagnosis of CTCL. Progressive, persistent or recurrent disease on or following two systemic therapies (e.g., bexarotene, romidepsin, etc.).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZTALMY (S)

Products Affected

- Ztalmy

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

ZURZUVAE (S)

Products Affected

- Zurzuvae

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Postpartum Depression (PPD): Diagnosis of PPD. Onset of symptoms in the third trimester or within 4 weeks of delivery. Prescriber attests that the patient has been counseled and has agreed to adhere to the following: Will follow instructions to not drive or operate machinery until at least 12 hours after taking each dose of Zurzuvae for the duration of the 14-day treatment course and that patients are informed that they may not be able to assess their own driving competence or the degree of driving impairment caused by Zurzuvae.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	14 days
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZYDELIG (S)

Products Affected

- Zydelig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic lymphocytic leukemia (CLL): Diagnosis of CLL. Used in combination with Rituxan (rituximab). The patient has relapsed on at least one prior therapy (eg, purine analogues [fludarabine, pentostatin, cladribine], alkylating agents [chlorambucil, cyclophosphamide], or monoclonal antibodies [rituximab]). Patient is a candidate for Rituxan (rituximab) monotherapy due to presence of other comorbidities (eg, coronary artery disease, peripheral vascular disease, diabetes mellitus, pulmonary disease [COPD]).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZYKADIA (S)

Products Affected

- Zykadia TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-small cell lung cancer (NSCLC) that is metastatic or recurrent. Disease is confirmed by the presence of an anaplastic lymphoma kinase (ALK)-positive tumor as detected by an FDA-approved test or a test performed at a facility approved by Clinical Laboratory Improvement Amendments (CLIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZYMFENTRA (S)

Products Affected

- Zymfentra 1-pen
- Zymfentra 2-pen
- Zymfentra 2-syringe

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

ZYTIGA (PREFERRED) (S)

Products Affected

- Abiraterone Acetate TABS 250MG, 500MG
- Abirtega

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Castration-Resistant Prostate Cancer (CRPC): Diagnosis of castration-resistant (chemical or surgical) or recurrent prostate cancer. Castration-Sensitive Prostate Cancer (CSPC): Diagnosis of castration-sensitive prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	CRPC, CSPC: 12 months
Other Criteria	Approve for continuation of prior therapy
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PART B VERSUS PART D

Products Affected

- Acyclovir Sodium INJ 50MG/ML
- Albuterol Sulfate NEBU 0.083%, 2.5MG/0.5ML
- Aminosyn II INJ 107.6MEQ/L; 1490MG/100ML; 1527MG/100ML; 1050MG/100ML; 1107MG/100ML; 750MG/100ML; 450MG/100ML; 990MG/100ML; 1500MG/100ML; 1575MG/100ML; 258MG/100ML; 405MG/100ML; 447MG/100ML; 1083MG/100ML; 795MG/100ML; 50MEQ/L; 600MG/100ML; 300MG/100ML; 750MG/100ML
- Amphotericin B INJ
- Amphotericin B Liposome
- Aprepitant
- Azathioprine TABS 50MG
- Budesonide SUSP
- Cromolyn Sodium NEBU
- Cyclophosphamide CAPS
- Cyclosporine CAPS
- Cyclosporine Modified
- Engerix-b
- Envarsus Xr
- Everolimus TABS 0.25MG, 0.5MG, 0.75MG, 1MG
- Ganciclovir INJ 500MG, 500MG/10ML
- Gengraf CAPS 100MG, 25MG
- Heplisav-b
- Hyperhep B INJ 110UNIT/0.5ML
- Imovax Rabies (h.d.c.v.)
- Ipratropium Bromide INHALATION SOLN 0.02%
- Ipratropium Bromide/albuterol Sulfate
- Mycophenolate Mofetil CAPS
- Mycophenolate Mofetil SUSR
- Mycophenolate Mofetil TABS
- Mycophenolic Acid Dr
- Nutrilipid
- Ondansetron Hcl SOLN 4MG/5ML
- Ondansetron Hydrochloride TABS
- Ondansetron Odt TBDP 4MG, 8MG
- 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
- Prograf PACK
- Rabavert
- Recombivax Hb
- Sirolimus SOLN
- Sirolimus TABS
- Tacrolimus CAPS
- Tacrolimus Er
- Tobramycin NEBU 300MG/5ML

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.

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