TUFTS Health Plan

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Tufts Health Plan Senior Care Options (HMO-SNP)

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2026 Prior Authorization Medical Necessity Guidelines

Effective: November 1, 2026

H8330_2026_RXOPS474_C

ABILIFY MYCITE

Products Affected

- Abilify Mycite Maintenance Kit TBPK 10MG
- Abilify Mycite Starter Kit TBPK 15MG, 20MG, 2MG, 30MG, 5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet the following: 1) have a documented diagnosis of bipolar I disorder, major depressive disorder or schizophrenia 2) the member must have documentation of worsening symptoms with oral aripiprazole.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a psychiatrist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ABIRATERONE

Products Affected

- Abiraterone Acetate TABS 250MG, 500MG
- Abirtega
- Yonsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic castration-resistant prostate cancer (CRPC) or metastatic high-risk castration-sensitive prostate cancer and abiraterone is being used in combination with prednisone.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ADALIMUMAB

Products Affected

- Adalimumab-aaty 1-pen Kit
- Adalimumab-aaty 2-pen Kit
- Adalimumab-aaty 2-syringe
- Adalimumab-aaty Cd/uc/hs Starter
- Adalimumab-adbm
- Adalimumab-adbm Crohns/uc/hs Starter
- Adalimumab-adbm Psoriasis/uveitis Starter
- Adalimumab-adbm Starter Package For Crohns Disease/uc/hs
- Adalimumab-adbm Starter Package For Psoriasis/uveitis
- Humira INJ 10MG/0.1ML, 20MG/0.2ML, 40MG/0.4ML, 40MG/0.8ML
- Humira Pediatric Crohns Disease Starter Pack INJ 0, 80MG/0.8ML
- Humira Pen
- Humira Pen-cd/uc/hs Starter
- Humira Pen-pediatric Uc Starter Pack
- Humira Pen-ps/uv Starter INJ 0

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A

Required Medical Information	Ankylosing Spondylitis: The member must have a documented diagnosis of active ankylosing spondylitis. Crohn's Disease (CD): The member must have a documented diagnosis of CD. Ulcerative Colitis (UC): The member must have a documented diagnosis of moderately to severely active UC. Hidradenitis Suppurativa: The member must have a documented diagnosis of moderate-to-severe hidradenitis suppurativa. Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe chronic plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis. Rheumatoid Arthritis (RA): The member must have a documented diagnosis of moderately to severely active RA and has a trial and failure, contraindication, or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA): The member must have a documented diagnosis of PJIA and has a trial and failure, contraindication or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide or sulfasalazine. Uveitis: The member must have a documented diagnosis of non-infectious uveitis.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, gastroenterologist, ophthalmologist, or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AIMOVIG

Products Affected

- Aimovig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: The member must have a documented diagnosis of migraine. Subsequent: The member has had a clinically significant reduction in migraine days per month from baseline as determined by the prescriber.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial Approval: 6 months. Subsequent approval: 2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AKEEGA

Products Affected

- Akeega

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of prostate cancer and disease is either a) metastatic, castration-resistant, and deleterious or suspected deleterious BRCA-mutated (BRCAm) or b) metastatic, castration-sensitive, and deleterious or suspected deleterious BRCA2-mutated (BRCA2m) 2) Requested drug is being used in combination with prednisone and 3) 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	The prescriber must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALECENSA

Products Affected

- Alecensa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Anaplastic Lymphoma Kinase positive (ALK-positive), metastatic Non-small Cell Lung Cancer (NSCLC) or the requested drug is being used as adjuvant treatment following tumor resection of anaplastic lymphoma kinase (ALK)-positive non-small cell lung cancer (NSCLC).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALOSETRON

Products Affected

- Alosetron Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of severe diarrhea-predominant irritable bowel syndrome in female.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALUNBRIG

Products Affected

- Alunbrig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ARCALYST

Products Affected

- Arcalyst

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cryopyrin-associated periodic syndromes: The member must have a documented diagnosis of a Cryopyrin-Associated Periodic Syndrome, including Familial Cold Autoinflammatory Syndrome, or Muckle-Wells Syndrome. Deficiency of interleukin-1 receptor antagonist: The member must have a documented diagnosis of deficiency of interleukin-1 receptor antagonist and Arcalyst is being used for maintenance of remission in patients weighing 10kg or more. Recurrent Pericarditis (RP): The member must have a documented diagnosis of RP and Arcalyst is being used to reduce the risk of recurrence.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ARIKAYCE

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	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ARMODAFINIL AND MODAFINIL

Products Affected

- Armodafinil
- Modafinil TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage will not be approved for generalized fatigue, jet lag, or sleep-deprivation not associated with a covered diagnosis.
Required Medical Information	The member must have a documented diagnosis of narcolepsy, excessive sleepiness associated with obstructive sleep apnea, or shift-work sleep disorder.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AUGTYRO

Products Affected

- Augtyro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of 1) locally advanced or metastatic ROS1-positive non-small cell lung cancer (NSCLC) or 2) neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors that a) are locally advanced, metastatic, or where surgical resection is likely to result in severe morbidity and b) have progressed following treatment or have no satisfactory alternative therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescriber must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AUSTEDO

Products Affected

- Austedo
- Austedo Xr
- Austedo Xr Patient Titration Kit

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chorea Associated with Huntington's Disease: The member must have a documented diagnosis of chorea associated with Huntington's Disease. Tardive Dyskinesia: The member must have a documented diagnosis of Tardive Dyskinesia.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AVMAPKI FAKZYNJA

Products Affected

- Avmapki Fakzynja Co-pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of recurrent low-grade serous ovarian cancer (LGSOC) 2) Presence of a KRAS-mutation 3) Patient has received prior systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AYVAKIT

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): The member must have a documented diagnosis of PDGFRA Exon 18 mutation-positive, including PDGFRA D842V mutations, unresectable or metastatic GIST. Advanced Systemic Mastocytosis (AdvSM): The member must have a documented diagnosis of AdvSM, which includes aggressive systemic mastocytosis (ASM), systemic mastocytosis with an associated hematological neoplasm (SM-AHN), and mast cell leukemia (MCL). Indolent Systemic Mastocytosis (ISM): The member must have a document diagnosis of Indolent Systemic Mastocytosis
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an allergist, immunologist, or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BALVERSA

Products Affected

- Balversa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of locally advanced or metastatic urothelial carcinoma that has susceptible FGFR3 genetic alterations and 1) the member progressed during or following at least one line of prior systemic therapy and 2) the member had been treated with prior PD-1 inhibitor (e.g., nivolumab, pembrolizumab) or PD-L1 inhibitor therapy (e.g., avelumab) or the member is not a candidate for PD-1 or PD-L1 inhibitor therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BENLYSTA

Products Affected

- Benlysta INJ 200MG/ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Benlysta (belimumab) will not be approved as monotherapy, for members with severe active lupus nephritis or severe active central nervous system lupus, for members who are autoantibody negative, or in combination with other biologics or intravenous cyclophosphamide.
Required Medical Information	The member must have a documented diagnosis of active, autoantibody-positive (e.g., ANA, anti-ds-DNA, anti-Sm) systemic lupus erythematosus (SLE) or active lupus nephritis and is concurrently taking standard therapy (e.g., antimalarials, corticosteroids, or immunosuppressives).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a nephrologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BERINERT

Products Affected

- Berinert

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Hereditary Angioedema.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an allergist or immunologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BESREMI

Products Affected

- Besremi
- Besremi Pen

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of polycythemia vera or essential thrombocythemia.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BEXAROTENE GEL

Products Affected

- Bexarotene GEL

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of cutaneous T-cell lymphoma.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or dermatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BOSULIF

Products Affected

- Bosulif
- Bosutinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) chronic phase Philadelphia chromosome-positive (Ph+) chronic myelogenous leukemia (CML), newly-diagnosed or resistant or intolerant to at least one prior therapy (e.g. imatinib) or 2) accelerated, or blast phase Ph+ CML with resistance or intolerance to at least one prior therapy (e.g. imatinib).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BRAFTOVI

Products Affected

- Braftovi CAPS 75MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Braftovi is not indicated for treatment of patients with wild-type BRAF melanoma or wild-type BRAF CRC.
Required Medical Information	Metastatic Colorectal Cancer (CRC): The member must have a documented diagnosis of metastatic CRC with a BRAF V600E mutation 1) after prior therapy and will be taken in combination with cetuximab OR 2) and will be taken in combination with cetuximab and fluorouracil-based chemotherapy. Melanoma (unresectable or metastatic): The member must have a documented diagnosis of unresectable or metastatic melanoma with a BRAF V600E or V600K mutation and will be taken in combination with binimetinib. Non-Small Cell Lung Cancer (NSCLC): The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) with a BRAF V600E mutation and will be taken in combination with binimetinib.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BRUKINSA

Products Affected

- Brukinsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) Mantle Cell Lymphoma (MCL) and has received at least one prior therapy or 2) Relapsed or refractory Marginal Zone Lymphoma (MZL) and has received at least one anti-CD20-based regimen, 3) Waldenstrom's Macroglobulinemia, 4) chronic lymphocytic leukemia (CLL), 5) small lymphocytic lymphoma (SLL) or 6) Relapsed or refractory follicular lymphoma (FL), in combination with obinutuzumab, after two or more lines of systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CABOMETYX

Products Affected

- Cabometyx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Advanced Renal Cell Carcinoma (RCC): The member must have a documented diagnosis of advanced renal cell carcinoma (RCC). Differentiated Thyroid Cancer (DTC): The member must have a documented diagnosis of locally advanced or metastatic DTC that has progressed following prior VEGFR-targeted therapy and are radioactive iodine-refractory or ineligible. Hepatocellular Carcinoma (HCC): The member must have a documented diagnosis of HCC and has had a documented failure, contraindication, or intolerance with sorafenib. Pancreatic Neuroendocrine Tumors: The member must have a documented diagnosis of previously treated, unresectable, locally advanced or metastatic, well-differentiated pancreatic or extra-pancreatic neuroendocrine tumors (pNET or epNET).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CALQUENCE

Products Affected

- Calquence

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) or Small Lymphocytic Lymphoma (SLL): The member must have a documented diagnosis of CLL or SLL. Mantle Cell Lymphoma (MCL): The member must have a documented diagnosis of MCL and has received at least one prior therapy, or has previously untreated mantle cell lymphoma and is ineligible for autologous hematopoietic stem cell transplant.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CAPLYTA

Products Affected

- Caplyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) schizophrenia 2) bipolar depression or 3) major depressive disorder. For major depressive disorder, the requested drug is being used as adjunctive therapy with antidepressants.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a psychiatrist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CAPRELSA

Products Affected

- Caprelsa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of symptomatic or progressive medullary thyroid cancer with unresectable locally advanced or metastatic disease.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an endocrinologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CARGLUMIC

Products Affected

- Carglumic Acid

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have one of the following: 1) a documented diagnosis of chronic hyperammonemia due to N-acetylglutamate synthase (NAGS) deficiency. 2) a documented diagnosis of acute hyperammonemia due to N-acetylglutamate synthase (NAGS) deficiency and the requested drug is being used as adjunctive therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CAYSTON

Products Affected

- Cayston

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cystic fibrosis (CF): Diagnosis of CF AND Patient has evidence of Pseudomonas aeruginosa in the lungs.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

COBENFY

Products Affected

- Cobenfy
- Cobenfy Starter Pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of schizophrenia.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a psychiatrist.
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

COMETRIQ

Products Affected

- Cometriq

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of progressive, metastatic medullary thyroid cancer.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

COPIKTRA

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): The member must have a documented diagnosis of relapsed or refractory CLL or SLL and has received at least two prior therapies.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COSENTYX

Products Affected

- Cosentyx
- Cosentyx Sensoready Pen
- Cosentyx Unoready

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Ankylosing Spondylitis: The member must have a documented diagnosis of active ankylosing spondylitis. Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis. Non-radiographic Axial Spondyloarthritis: The member must have a documented diagnosis of active non-radiographic axial spondyloarthritis with objective signs of inflammation (e.g., 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.) and has had a minimum duration of one month trial and failure, contraindication, or intolerance to two NSAIDs (e.g., ibuprofen, meloxicam, naproxen) at maximally indicated doses. Enthesitis-related Arthritis (ERA): The member must have a documented diagnosis of active ERA and has had a minimum duration of one month trial and failure, contraindication, or intolerance to two NSAIDs (e.g., ibuprofen, meloxicam, naproxen). Hidradenitis Suppurativa: The member must have a documented diagnosis of moderate-to-severe hidradenitis suppurativa.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist or rheumatologist.
Coverage Duration	2 years

Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COTELLIC

Products Affected

- Cotellic

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Cotellic is not indicated for treatment of patients with wild-type BRAF melanoma.
Required Medical Information	The member must have a documented diagnosis of 1) unresectable or metastatic melanoma with a BRAF V600E or V600K mutation and is being taken in combination with Zelboraf (vemurafenib) or 2) histiocytic neoplasm and is being used as monotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CRESEMBA ORAL

Products Affected

- Cresemba CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion 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	1 year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DANZITEN

Products Affected

- Danziten

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Newly diagnosed Philadelphia chromosome positive chronic myeloid leukemia (Ph+ CML) in chronic phase or 2) Diagnosis of chronic phase and accelerated phase Ph+ CML resistant to or intolerant to prior therapy that included imatinib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DASATINIB

Products Affected

- Dasatinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic Myeloid Leukemia (CML): 1) Newly diagnosed Philadelphia chromosome-positive (Ph+) CML in chronic phase or 2) Diagnosis of chronic, accelerated, or myeloid or lymphoid blast phase Ph+ CML and documented resistance or intolerance to prior therapy, including imatinib mesylate or Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia (Ph+ALL): Diagnosis of Ph+ALL and documented resistance or intolerance to prior therapy. For Pediatric Members: 1) Diagnosis of Ph+CML in chronic phase or 2) newly diagnosed Ph+ALL and the requested drug is being used in combination with chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DAURISMO

Products Affected

- Daurismo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have newly diagnosed acute myelogenous leukemia (AML) and the requested drug is being used with low-dose cytarabine and the member meets one of the following: 1) is 75 years of age or older or 2) has comorbidities that make them ineligible for intensive induction chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DIACOMIT

Products Affected

- Diacomit

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of seizures associated with Dravet syndrome and is concurrently taking clobazam.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DICHLORPHENAMIDE

Products Affected

- Dichlorphenamide

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of primary hyperkalemic periodic paralysis, primary hypokalemic periodic paralysis, and related variants.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DOPTELET

Products Affected

- Doptelet

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of one of the following: 1) Thrombocytopenia associated with chronic liver disease (CLD) and is scheduled to undergo a procedure 2) Thrombocytopenia with chronic immune thrombocytopenia (ITP) who have had an insufficient response to a previous treatment.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DROXIDOPA

Products Affected

- Droxidopa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of neurogenic orthostatic hypotension (NOH).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DUPIXENT

Products Affected

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

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Dupixent will not be approved for the relief of acute bronchospasm or status asthmaticus.
Required Medical Information	Atopic Dermatitis: The member must have a documented diagnosis of moderate to severe atopic dermatitis with a trial and failure of a minimum 30-day supply (or 14-day supply for topical corticosteroids), contraindication, or intolerance to at least one of the following: Medium or higher potency topical corticosteroid, pimecrolimus cream, tacrolimus ointment, or crisaborole ointment. Asthma: The member must have a documented diagnosis of moderate-to- severe asthma with an eosinophilic phenotype or is dependent on oral corticosteroids. Rhinosinusitis (chronic) with nasal polyposis: The member must have a documented diagnosis of chronic rhinosinusitis with nasal polyposis (CRSwNP) and is inadequately controlled on current treatment alone. Prurigo Nodularis: The member must have a document diagnosis of Prurigo Nodularis. Eosinophilic Esophagitis: Diagnosis of Eosinophilic Esophagitis. COPD: Diagnosis of COPD with an eosinophilic phenotype and requested medication is being used as add-on maintenance treatment. Bullous Pemphigoid (BP): Diagnosis of BP. Allergic Fungal Rhinosinusitis (AFRS): Diagnosis of AFRS and a history of sino-nasal surgery.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an allergist, dermatologist, immunologist, otolaryngologist, pulmonologist, or gastroenterologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ELTROMBOPAG

Products Affected

- Eltrombopag Olamine

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Persistent or Chronic Immune Thrombocytopenic Purpura (ITP): The member must have a documented diagnosis of persistent or chronic ITP and has had an insufficient response or intolerance to corticosteroids, immunoglobulins, or splenectomy. Severe Aplastic Anemia: 1) The member must have a documented diagnosis of severe aplastic anemia and 2) will be taken in combination with, or in those who have had an insufficient response with, standard immunosuppressive therapy. Thrombocytopenia with Chronic Hepatitis C: The member must have a documented diagnosis of thrombocytopenia with chronic hepatitis C infection.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

EMGALITY

Products Affected

- Emgality

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial Approval: The member must have a documented diagnosis of one of the following: 1) Migraine 2) Episodic Cluster headache. Subsequent Approval: The member has had a clinically significant reduction in migraine days per month or the frequency of weekly cluster headache attacks from baseline as determined by the prescriber.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial Approval: 6 months. Subsequent Approval: Life of Plan.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EMPAVELI

Products Affected

- Empaveli

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Paroxysmal Nocturnal Hemoglobinuria (PNH) (initial): Diagnosis of PNH.
Age Restrictions	N/A
Prescriber Restrictions	PNH (initial): Prescribed by or in consultation with a hematologist or oncologist or nephrologist.
Coverage Duration	2 years
Other Criteria	PNH (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ENBREL

Products Affected

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

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Ankylosing Spondylitis: The member must have a documented diagnosis of active ankylosing spondylitis. Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Rheumatoid Arthritis (RA): The member must have a documented diagnosis of RA and has a trial and failure, contraindication, or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA): The member must have a documented diagnosis of PJIA and has a trial and failure, contraindication, or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ENSACOVE

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. 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	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EPIDIOLEX

Products Affected

- Epidiolex

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of seizures associated with Lennox-Gastaut syndrome (LGS), Dravet syndrome (DS) or tuberous sclerosis complex (TSC).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ERIVEDGE

Products Affected

- Erivedge

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic basal cell carcinoma or locally advanced basal cell carcinoma that has recurred following surgery or the member is not a candidate for surgery, and not a candidate for radiation.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a dermatologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ERLEADA

Products Affected

- Erleada

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of one of the following: 1) non-metastatic, castration-resistant prostate cancer 2) metastatic hormone-sensitive prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ESBRIET

Products Affected

- Pirfenidone CAPS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of idiopathic pulmonary fibrosis (IPF).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a pulmonologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EUCRISA

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	2 years
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.

EVEROLIMUS

Products Affected

- Everolimus TABS 10MG, 2.5MG, 5MG, 7.5MG
- Yulithira

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Advanced Hormone Receptor-Positive, HER2-Negative Breast Cancer (Advanced HR+ BC): The member must have a documented diagnosis of Advanced HR+ BC, the member is postmenopausal, concurrently taking exemestane and has a documented failure of letrozole or anastrozole. Advanced Renal Cell Carcinoma (ARCC): The member must have a documented diagnosis of ARCC and the member has a demonstrated disease progression or intolerance following a trial with sorafenib or sunitinib. Neuroendocrine Tumors (NET): The member must have a documented diagnosis of progressive neuroendocrine tumors of pancreatic origin (pNET) or progressive, well-differentiated, non-functional neuroendocrine tumors (NET) of gastrointestinal (GI) or lung origin, any of which are unresectable, locally advanced or metastatic. Renal Angiomyolipoma with Tuberous Sclerosis Complex (TSC): The member must have a documented presence of TSC and renal angiomyolipoma(s). Subependymal Giant Cell Astrocytoma (SEGA): The member must have a documented diagnosis of SEGA associated with TSC and the member is not a candidate for surgical resection.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A

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

EVEROLIMUS FOR ORAL SUSPENSION

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	Partial-onset Seizures Associated with Tuberous Sclerosis Complex (TSC): The member must have a documented diagnosis of partial-onset seizures associated with TSC. Subependymal Giant Cell Astrocytoma (SEGA): The member must have a documented diagnosis of SEGA associated with TSC and the member is not a candidate for surgical resection.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EXKIVITY

Products Affected

- Exkivity

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of locally advanced or metastatic non-small cell lung cancer (NSCLC) with epidermal growth factor receptor (EGFR) exon 20 insertion mutations and has progressed on or after platinum-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FASENRA

Products Affected

- Fasenra
- Fasenra Pen

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) severe asthma with an eosinophilic phenotype despite current treatment with both of the following medications: a) inhaled corticosteroids b) additional controller (Long-Acting Beta2-Agonist, Leukotriene Modifier, or Sustained Release Theophylline) or 2) eosinophilic granulomatosis with polyangiitis (EGPA) or 3) hypereosinophilic syndrome (HES) without an identifiable non-hematologic secondary cause (e.g., drug hypersensitivity, parasitic helminth infection, HIV infection, non-hematologic malignancy) and a) Pre-treatment blood eosinophil count greater than or equal to 1000 cells/microliter and b) Trial and failure, contraindication (CI), or intolerance to corticosteroid therapy (e.g., prednisone) or cytotoxic/immunosuppressive treatment (e.g., hydroxyurea, cyclosporine, imatinib).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an asthma specialist (e.g., allergist, immunologist, pulmonologist) or a rheumatologist or a hematologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

FINTEPLA

Products Affected

- Fintepla

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of seizures associated with Dravet syndrome or Lennox-Gastaut syndrome.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FOTIVDA

Products Affected

- Fotivda

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of relapsed or refractory advanced renal cell carcinoma following two or more prior systemic therapies.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

FRUZAQLA

Products Affected

- Fruzaqla

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of metastatic colorectal cancer. Patient has been previously treated with the following: A) Fluoropyrimidine-, oxaliplatin-, irinotecan-based chemotherapy, and B) Anti-VEGF biological therapy (e.g., bevacizumab, ramucirumab) and C) if RAS wild-type and medically appropriate, an anti-EGFR therapy
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GAVRETO

Products Affected

- Gavreto

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small Cell Lung Cancer (NSCLC): The member must have a documented diagnosis of metastatic rearranged during transfection (RET) fusion-positive NSCLC. Thyroid Cancer: The member must have advanced or metastatic RET fusion-positive thyroid cancer who require systemic therapy and who are radioactive iodine-refractory (if radioactive iodine is appropriate)
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GILOTRIF

Products Affected

- Gilotrif

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of one of the following: 1) Metastatic non-small cell lung cancer (NSCLC) and documented non-resistant epidermal growth factor receptor (EGFR) mutations 2) Metastatic, squamous cell NSCLC and documentation that the disease has progressed following platinum-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GLP1

Products Affected

- Mounjaro
- Ozempic INJ 2MG/3ML, 4MG/3ML, 8MG/3ML
- Ozempic TABS
- Rybelsus
- Trulicity

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Type 2 Diabetes (initial): Diagnosis of Type 2 Diabetes confirmed by a) documentation from the prescriber OR b) a record of a trial with a minimum 90-day supply with one product from any of the following drugs/classes: metformin-containing agent, DPP-4 inhibitors, DPP-4 inhibitor combinations, SGLT2 inhibitors, SGLT2 inhibitor combinations, alpha-glucosidase inhibitors, meglitinide analogues, sulfonylurea, or sulfonylurea combinations.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Type 2 diabetes mellitus (Reauthorization): Patient demonstrates positive clinical response to therapy
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GOMEKLI

Products Affected

- Gomekli

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of neurofibromatosis type 1 and 2) Has plexiform neurofibromas that are both of the following: a) inoperable and b) causing significant morbidity (e.g., disfigurement, motor dysfunction, pain, airway dysfunction, visual impairment).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GROWTH HORMONE REPLACEMENT THERAPY

Products Affected

- Genotropin
- Genotropin Miniquick

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Pediatric GHD, Initiation: Member has not attained epiphyseal closure as determined by X-ray, have failed to respond to at least TWO standard GH stimulation test, have documented gender-specific delayed bone age, have the height at initiation of therapy at greater than 2 standard deviations below normal mean for age and sex. Member must have one of the following: Chronic Renal Insufficiency prior to transplantation, Idiopathic Short Stature, Intrauterine Growth Retardation, Non-genetic GHD, Noonan Syndrome, Prader-Willi Syndrome, Short Stature Homeobox-containing gene (SHOX) deficiency, or Turner Syndrome. Pediatric GHD, Continuation: Documentation of the following is required: Medical history as it relates to growth, including any test results and growth chart, continuing care plan and an improvement in the annualized pre-treatment growth rate after the first six (6) months of therapy. Continuation of Therapy after Completion of Linear Growth: Member will be re-evaluated after GH treatments have been stopped for at least three (3) months to determine growth hormone status AND member must have failed to respond to at least one standard GH stimulation test. Acquired GHD: Member must have failed to respond to at least one standard GH stimulation test. AIDS Wasting Syndrome: Documented diagnosis of AIDS AND a weight loss of at least 10% from baseline weight OR a BMI of less than 20. Short Bowel Syndrome: Documented diagnosis of Short Bowel Syndrome from a gastroenterologist AND a documented dependence on IPN for nutritional support.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years

Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HAEGARDA

Products Affected

- Haegarda

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Hereditary Angioedema.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an allergist or immunologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HERNEXEOS

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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HYRNUO

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. Patient has received prior systemic therapy (e.g., chemotherapy).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IBRANCE

Products Affected

- Ibrance

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must 1) have a documented diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer and the requested drug is being used in combination with an aromatase inhibitor or 2) have a documented diagnosis of HR-positive, HER2- negative advanced or metastatic breast cancer with disease progression following endocrine therapy and documentation the requested drug will be used in combination with fulvestrant or 3) have a documented diagnosis of endocrine-resistant, PIK3CA-mutated, HR-positive, HER2- negative, locally advanced or metastatic breast cancer with disease progression following endocrine therapy and documentation the requested drug will be used in combination with inavolisib or 4) have a documented diagnosis of HR-positive, HER2-positive locally advanced or metastatic breast cancer after completion of induction therapy, in combination with endocrine therapy and trastuzumab (with or without pertuzumab).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IBTROZI

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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ICATIBANT

Products Affected

- Icatibant Acetate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Icatibant will not be approved for members with acquired angioedema or concurrently taking an angiotensin converting enzyme (ACE) inhibitor.
Required Medical Information	The member must have a documented diagnosis of Hereditary Angioedema (HAE) with a history of at least one severe attack in the past six months. For HAE Types 1 & 2, the diagnosis must be confirmed by laboratory testing (e.g., low C4 level, reduced C1 esterase inhibitor level or function).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an allergist, hematologist, or immunologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ICLUSIG

Products Affected

- Iclusig

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Iclusig will not be approved for members with newly diagnosed chronic phase CML.
Required Medical Information	Acute Lymphoblastic Leukemia (ALL): The member must be 1) T315I-positive or have a documented diagnosis of Philadelphia chromosome-positive ALL (Ph+ALL) for which no other tyrosine kinase inhibitor therapy is indicated or 2) newly diagnosed and in combination with chemotherapy. Chronic Myeloid Leukemia (CML): 1) The member must be T315I-positive or 2) have a documented diagnosis of accelerated phase, or blast phase CML for which no other kinase inhibitor therapy is indicated or 3) The member must have a diagnosis of Chronic phase (CP) chronic myeloid leukemia (CML) with resistance or intolerance to at least two prior kinase inhibitors.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IDHIFA

Products Affected

- Idhifa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of relapsed or refractory acute myeloid leukemia (AML) with an isocitrate dehydrogenase-2 (IDH2) mutation.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IMBRUVICA

Products Affected

- Imbruvica CAPS
- Imbruvica SUSP
- Imbruvica TABS 140MG, 280MG, 420MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL): Diagnosis of CLL/SLL. Waldenstrom's macroglobulinemia: Diagnosis of Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma. 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	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist, oncologist, or transplant specialist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMKELDI

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 Myeloid Leukemia (CML): Diagnosis of Philadelphia chromosome positive CML (Ph+ CML). Acute Lymphoblastic Leukemia (ALL): Diagnosis of Philadelphia chromosome positive ALL (Ph+ 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	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INCRELEX

Products Affected

- Increlex

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage of Increlex will not be authorized for conditions resulting in secondary forms of IGF-1 deficiency that include, but are not limited to: GH deficiency, malnutrition, hypothyroidism, or chronic steroid therapy.
Required Medical Information	The member must have a documented diagnosis of severe primary IGF-1 deficiency as defined by a height SD score less than or equal to -3.0, a basal IGF-1 SD score less than or equal to -3.0 , normal or elevated GH level OR GH gene deletion and has developed neutralizing antibodies to GH. Radiographs documenting open epiphyses are required for members who are Tanner stage III or greater.
Age Restrictions	The member must be 2 years or age or older.
Prescriber Restrictions	The prescribing physician must be an endocrinologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INLURIYO

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). Disease has progressed following at least one line of endocrine therapy (e.g., fulvestrant, anastrozole, letrozole, exemestane).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INLYTA

Products Affected

- Inlyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of advanced renal cell carcinoma and one of the following two requirements: 1) The member is using Inlyta as first line treatment in combination with avelumab or pembrolizumab 2) The member is using Inlyta as a single agent and has failed a trial of at least one systemic therapy (including but not limited to everolimus, Nexavar, sunitinib, Torisel, Votrient).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INQOVI

Products Affected

- Inqovi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) myelodysplastic syndromes (MDS), including previously treated and untreated, de novo and secondary MDS with the following French-American-British subtypes (refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, and chronic myelomonocytic leukemia [CMML]) and intermediate-1, intermediate-2, and high-risk International Prognostic Scoring System groups or 2) newly diagnosed acute myelogenous leukemia (AML) and a) at least 75 years of age or has comorbidities that preclude the use of intensive induction chemotherapy and b) being used in combination with venetoclax.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INREBIC

Products Affected

- Inrebic

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of intermediate-2 or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IRESSA

Products Affected

- Gefitinib

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) in tumors that have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutations.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ITOVEBI

Products Affected

- Itovebi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of breast cancer. 2) Disease is locally advanced or metastatic. 3) Disease is PIK3CA-mutated, Hormone receptor (HR)-positive, Human epidermal growth-factor receptor 2 (HER2)-negative. 4) Used following recurrence on or after completing adjuvant endocrine therapy (e.g. goserelin, anastrozole, tamoxifen). 5) Used in combination with palbociclib and fulvestrant.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IWILFIN

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) with at least a partial response to prior multiagent, multimodality therapy including anti-GD2 immunotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescriber must be an oncologist.
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JAKAFI

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	Chronic Graft-versus-host Disease (GVHD): The member must have a documented diagnosis of chronic GVHD after failure of one or two lines of systemic therapy. Myelofibrosis: The member must have a documented diagnosis of intermediate or high-risk myelofibrosis, including primary myelofibrosis, post-polycythemia vera myelofibrosis, and post-essential thrombocythemia myelofibrosis. Polycythemia Vera: The member must have a documented diagnosis of polycythemia vera with an inadequate response, contraindication, or inability to tolerate hydroxyurea. Steroid-Refractory Acute Graft-versus-host Disease (GVHD): The member must have a document diagnosis of steroid-refractory acute GVHD.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 Years
Other Criteria	For Myelofibrosis: Subsequent authorization requires documentation of spleen size reduction or symptomatic improvement.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JAYPIRCA

Products Affected

- Jaypirca

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) relapsed or refractory mantle cell lymphoma (MCL) and has received at least two prior lines of systemic therapies including a BTK inhibitor or 2) chronic lymphocytic leukemia or small lymphocytic lymphoma (CLL/SLL) and previously been treated with a covalent BTK inhibitor.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JIDEYTRO

Products Affected

- Jideytro

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: Diagnosis of non-small cell lung cancer. Disease is i) locally advanced or ii) metastatic. Presence of ROS1 rearrangement positive tumor(s). Patient has received prior treatment with a ROS1 kinase inhibitor (e.g., Augtyro, Xalkori, Rozlytrek).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JUBBONTI

Products Affected

- Jubbonti

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Coverage of Jubbonti (denosumab) for the treatment of osteoporosis in men and postmenopausal women will be authorized when the following criteria are met: 1) The member is at high risk of fracture or 2) the member has had an inadequate response to, or is unable to tolerate therapy with at least one of the traditional osteoporosis treatments [including but not limited to: alendronate, calcitonin, ibandronate, raloxifene, risedronate, zoledronic acid] or 3) the member is a female at high risk for fracture receiving adjuvant aromatase inhibitor therapy for breast cancer and is using the requested drug as a treatment to increase bone mass. Coverage of Jubbonti may also be authorized for 1) men at high risk of fracture who are receiving androgen deprivation therapy for non-metastatic prostate cancer or 2) treatment for glucocorticoid-induced osteoporosis in men and women at high risk for fracture.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

JYNARQUE

Products Affected

- Jynarque TABS
- Tolvaptan TABS

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

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

KALYDECO

Products Affected

- Kalydeco

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Kalydeco is not effective in patients with cystic fibrosis who are homozygous for the F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene.
Required Medical Information	The member must have a documented diagnosis of cystic fibrosis (CF) and have one mutation in the CFTR gene that is responsive to Kalydeco based on clinical and/or in vitro assay data. If the patient's genotype is unknown, an FDA-cleared cystic fibrosis mutation test should be used to detect the presence of a CFTR mutation followed by verification with bi-directional sequencing when recommended by the mutation test instructions for use.
Age Restrictions	Granules: The member must be 1 month of age or older.
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KERENDIA

Products Affected

- Kerendia

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of chronic kidney disease associated with type 2 diabetes or heart failure with left ventricular ejection fraction greater than or equal to 40%,
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KESIMPTA

Products Affected

- Kesimpta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of relapsing forms of multiple sclerosis (MS), to include clinically isolated syndrome, relapsing-remitting disease, and active secondary progressive disease.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KINERET

Products Affected

- Kineret

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, Orencia (abatacept), Rinvoq (upadacitinib), Xeljanz/Xeljanz XR (tofacitinib), or attestation demonstrating a trial may be inappropriate, OR b) For continuation of prior therapy. Neonatal-Onset Multisystem Inflammatory Disease (NOMID) (initial): Diagnosis of NOMID AND dx of NOMID has been confirmed by one of the following: 1) NLRP-3 (nucleotide-binding domain, leucine rich family (NLR), pyrin domain containing 3) gene (also known as Cold-Induced Auto-inflammatory Syndrome-1 [CIAS1]) mutation OR 2) Both of the following: a) two of the following clinical symptoms: urticaria-like rash, cold/stress triggered episodes, sensorineural hearing loss, musculoskeletal symptoms (e.g., arthralgia, arthritis, myalgia), chronic aseptic meningitis, or skeletal abnormalities (e.g., epiphyseal overgrowth, frontal bossing) AND b) elevated acute phase reactants (eg, erythrocyte sedimentation rate [ESR], C-reactive protein [CRP], serum amyloid A [SAA]). Deficiency of Interleukin-1 Receptor Antagonist (DIRA): Diagnosis of DIRA.
Age Restrictions	N/A
Prescriber Restrictions	RA (initial): Prescribed by or in consultation with a rheumatologist. NOMID (initial): Prescribed by or in consultation with allergist/immunologist or rheumatologist or pediatrician.
Coverage Duration	1 year

Other Criteria	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. NOMID (reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KISQALI

Products Affected

- Kisqali
- Kisqali Femara 200 Dose
- Kisqali Femara 400 Dose
- Kisqali Femara 600 Dose

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer in combination with a) an aromatase inhibitor as initial endocrine-based therapy or b) fulvestrant as initial endocrine-based therapy or following disease progression on endocrine therapy or 2) HR-positive, HER2-negative stage II and III early breast cancer at high risk of recurrence and the requested drug is being used with an aromatase inhibitor.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KOMZIFTI

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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KOSELUGO

Products Affected

- Koselugo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Neurofibromatosis type 1 and have symptomatic, inoperable plexiform neurofibromas.
Age Restrictions	The member must be 1 year of age or older.
Prescriber Restrictions	The prescribing physician must be a neurologist or an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KRAZATI

Products Affected

- Krazati

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-Small Cell Lung Cancer (NSCLC): Diagnosis of NSCLC. Disease is 1) locally advanced or metastatic 2) KRAS G12C-mutated and 3) patient has received at least one prior systemic therapy (e.g., chemotherapy, immunotherapy). Colorectal Cancer (CRC): 1) Diagnosis of KRAS G12C mutated locally advanced or metastatic CRC and 2) the patient has received prior treatment with fluoropyrimidine-, oxaliplatin-, and irinotecan based chemotherapy and 3) the requested drug is being used In combination with cetuximab.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LAPATINIB

Products Affected

- Lapatinib Ditosylate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For estrogen receptor (ER)-positive, human epidermal growth factor receptor 2 (HER2) overexpressing advanced or metastatic breast cancer, the member must meet ALL of the following criteria: 1) Documented diagnosis of HER2 overexpressing advanced or metastatic breast cancer. 2) The member has failed prior therapy with an anthracycline and a taxane chemotherapeutic agent. 3) The member has failed prior therapy with trastuzumab. 4) The member is concurrently treated with capecitabine. Hormone Receptor Positive Metastatic Breast Cancer in Post-menopausal Women: The member must have a documented diagnosis of hormone receptor positive metastatic breast cancer that overexpresses the HER2 receptor and is concurrently being treated with letrozole
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LAZCLUZE

Products Affected

- Lazcluze

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of non-small cell lung cancer (NSCLC). 2) Disease is locally advanced or metastatic. 3) Used as first line treatment of NSCLC. 4) Used in combination with amivantamab. 5) Presence of epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R substitution mutations.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LENALIDOMIDE

Products Affected

- Lenalidomide

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Follicular Lymphoma (FL): The member must have a documented diagnosis of previously treated FL and the requested drug is being used in combination with a rituximab product. Mantle Cell Lymphoma (MCL): The member must have a documented diagnosis of MCL and the member's disease has relapsed or progressed after two prior therapies, one of which included Velcade (bortezomib). Marginal Zone Lymphoma (MZL): The member must have a documented diagnosis of previously treated MZL and the requested drug is being used in combination with a rituximab product. Multiple Myeloma: The member must have a documented diagnosis of multiple myeloma and requested drug is being used in combination with dexamethasone or as maintenance therapy in a member following autologous hematopoietic stem cell transplantation. Myelodysplastic Syndrome (MDS): The member must have a documented diagnosis of transfusion-dependent anemia due to low- or intermediate-1-risk MDS associated with the 5q-deletion cytogenetic abnormality.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LENVIMA

Products Affected

- Lenvima 10 Mg Daily Dose
- Lenvima 12mg Daily Dose
- Lenvima 14 Mg Daily Dose
- Lenvima 18 Mg Daily Dose
- Lenvima 20 Mg Daily Dose
- Lenvima 24 Mg Daily Dose
- Lenvima 4 Mg Daily Dose
- Lenvima 8 Mg Daily Dose

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Advanced Endometrial Carcinoma: Diagnosis of advanced endometrial carcinoma, that is that is mismatch repair proficient (pMMR) or not microsatellite instability-high (MSI-H) and has disease progression following prior systemic therapy and are not candidates for curative surgery or radiation and the requested drug will be used in combination with pembrolizumab. Advanced Renal Cell Carcinoma (ARCC): Diagnosis of ARCC and 1) has had one prior antiangiogenic therapy and requested drug is being used in combination with everolimus or 2) being used as first-line treatment in combination with pembrolizumab. Hepatocellular carcinoma (HCC): Diagnosis of unresectable hepatocellular carcinoma. Thyroid Cancer: Diagnosis of locally recurrent or metastatic, progressive, radioactive iodine-refractory differentiated thyroid cancer.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LIDOCAINE TRANSDERMAL PATCHES

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	For Postherpetic Neuralgia or Diabetic Neuropathy, the member must have had a failure, adverse reaction, or contraindication to gabapentin. Lidocaine transdermal patches will also be approved for members who are not candidates for opioid or other oral pain management therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Coverage will be authorized for new members if their pain is currently well-controlled on lidocaine transdermal patches.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LIFYORLI

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	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LIVTENCITY

Products Affected

- Livtencity

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member, weighing 35kg or more, must have a documented diagnosis of post-transplant cytomegalovirus (CMV) infection/disease, that is refractory to treatment (with or without genotypic resistance) with ganciclovir, valganciclovir, cidofovir, or foscarnet.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LONSURF

Products Affected

- Lonsurf

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Metastatic Colorectal Cancer (mCRC): The member must have a documented diagnosis of mCRC and has been previously treated with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy, an anti-vascular endothelial growth factor (VEGF) biological therapy, and if rat sarcoma viral oncogene (RAS) wild-type, an anti-epidermal growth factor receptor (EGFR) therapy. Metastatic Gastric or Gastroesophageal Junction Adenocarcinoma: The member must have a documented diagnosis of metastatic gastric or gastroesophageal junction adenocarcinoma and has been previously treated with at least two prior lines of chemotherapy that included a fluoropyrimidine, a platinum, either a taxane or irinotecan, and if appropriate, HER2/neu-targeted therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LORBRENA

Products Affected

- Lorbrena

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) whose tumors are anaplastic lymphoma kinase (ALK)-positive.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LUMAKRAS

Products Affected

- Lumakras

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of 1) KRAS G12C-mutated locally advanced or metastatic non-small cell lung cancer (NSCLC) and has received at least one prior systemic therapy or 2) KRAS G12C-mutated Metastatic Colorectal Cancer (mCRC) and the requested drug is being used in combination with panitumumab and has received prior fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LYBALVI

Products Affected

- Lybalvi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of schizophrenia or Bipolar I disorder.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a psychiatrist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LYNPARZA

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	Breast Cancer: Diagnosis of 1) deleterious or suspected deleterious gBRCA-mutated, HER2-negative metastatic breast cancer and has been treated with chemotherapy in the neoadjuvant, adjuvant, or metastatic setting and, if hormone receptor-positive, the member should have prior endocrine therapy or contraindication to or inability to tolerate endocrine therapy or 2) high risk early breast cancer, has been treated with neoadjuvant or adjuvant chemotherapy. Ovarian Cancer: 1) Maintenance treatment of deleterious or suspected deleterious germline or somatic BRCA-mutated advanced epithelial ovarian, fallopian tube, or primary peritoneal cancer who are in complete or partial response to first-line platinum-based chemotherapy or 2) Maintenance treatment of advanced epithelial ovarian, fallopian tube, or primary peritoneal cancer in combination with bevacizumab and the member is in complete or partial response to first-line platinum-based chemotherapy and the members cancer is associated with homologous recombination deficiency (HRD)-positive status or 3) Maintenance treatment of recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer and the member is in complete or partial response to platinum-based chemotherapy. Pancreatic Cancer: Diagnosis of deleterious or suspected deleterious germline BRCA-mutated metastatic pancreatic adenocarcinoma and no disease progression after at least 16 weeks of first-line platinum-based chemotherapy. Prostate Cancer: Diagnosis of deleterious or suspected deleterious germline or somatic homologous recombination repair (HRR) gene-mutated metastatic castration-resistant prostate cancer (mCRPC) who have progressed following prior treatment with enzalutamide or abiraterone or a documented diagnosis of deleterious or suspected deleterious BRCA-mutated (BRCAm) metastatic castration-resistant prostate cancer (mCRPC) and is being used in combination with abiraterone and prednisone or prednisolone.
Age Restrictions	N/A

Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LYTGOBI

Products Affected

- Lytgobi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of previously treated, unresectable, locally advanced or metastatic intrahepatic cholangiocarcinoma harboring fibroblast growth factor receptor 2 (FGFR2) gene fusions or other rearrangements
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MAVYRET

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 guidance.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a gastroenterologist, hepatologist, or an infectious disease specialist.
Coverage Duration	Criteria will be applied consistent with current AASLD-IDSA guidance.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MEDICATIONS FOR THE TREATMENT OF PULMONARY HYPERTENSION

Products Affected

- Adempas
- Alyq
- Ambrisentan
- Bosentan TABS
- Macitentan
- Opsumit
- Orenitram
- Orenitram Titration Kit Month 1
- Orenitram Titration Kit Month 2
- Orenitram Titration Kit Month 3
- Riociguat
- Sildenafil Citrate TABS 20MG
- Tadalafil TABS 20MG
- Ventavis
- Winrevair

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (PAH) The member must have a documented diagnosis of pulmonary arterial hypertension as confirmed by right heart catheterization. Chronic thromboembolic pulmonary hypertension (CTEPH) Diagnosis of persistent/recurrent CTEPH (after surgical treatment or inoperable)
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a cardiologist or pulmonologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MEKINIST

Products Affected

- Mekinist

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Mekinist will not be approved as a single agent for members who have received prior BRAF-inhibitor therapy.
Required Medical Information	Single Agent: The member must have a documented diagnosis of unresectable or metastatic melanoma with a BRAF V600E or V600K mutations. In Combination with Tafinlar: The member must have a documented diagnosis of one of the following: 1) Unresectable or metastatic melanoma with a BRAF V600E or V600K mutation. 2) Melanoma with a BRAF V600E or V600K mutation and involvement of lymph node(s) following complete resection. 3) Metastatic non-small cell lung cancer (NSCLC) with a BRAF V600E mutation. 4) Locally advanced or metastatic anaplastic thyroid cancer (ATC) with a BRAF V600E mutation with no satisfactory locoregional treatment options. 5) Unresectable or metastatic solid tumors with BRAF V600E mutation and has progressed following prior treatment and have no satisfactory alternative treatment options. 6) low grade glioma with BRAF V600E mutation that requires systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MEKTOVI

Products Affected

- Mektovi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) unresectable or metastatic melanoma with a BRAF V600E or V600K mutation, and will be taken in combination with encorafenib or 2) metastatic non-small cell lung cancer (NSCLC) with a BRAF V600E mutation, and will be taken in combination with encorafenib.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MIFEPRISTONE

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	The member must have a documented diagnosis of hyperglycemia secondary to hypercortisolism with endogenous Cushing's syndrome and type 2 diabetes mellitus OR glucose intolerance AND has failed surgery OR is not a candidate for surgery.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MODEYSO

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. Disease has progressed following prior therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NAYZILAM

Products Affected

- Nayzilam

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of a seizure disorder requiring acute treatment.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NERLYNX

Products Affected

- Nerlynx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Extended Adjuvant Treatment of Early-stage Breast Cancer: The member must have a documented diagnosis of early stage human epidermal growth receptor type 2 (HER2)-positive breast cancer and has had previous adjuvant treatment with Herceptin-based therapy. Advanced or Metastatic Breast Cancer: The member must have a documented diagnosis of advanced or metastatic HER2-positive breast cancer, is using Nerlynx in combination with capecitabine, and has received two or more prior anti-HER2 based regimens.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NEXLETOL

Products Affected

- Nexletol

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The following criteria must be met: 1) Patient has an elevated LDL-C level while being treated with maximally tolerated statin therapy or has an elevated LDL-C level and a contraindication/intolerance to statin therapy and 2) Patient must have a documented diagnosis of one of the following: a) Heterozygous Familial Hypercholesterolemia (HeFH) b) atherosclerotic cardiovascular disease or c) established cardiovascular disease (CVD) or is at high risk for a CVD event but without established CVD.
Age Restrictions	N/A
Prescriber Restrictions	The medication must be prescribed by or in consultation with a cardiologist, endocrinologist, lipidologist, or neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NEXLIZET

Products Affected

- Nexlizet

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The following criteria must be met: 1) Patient has an elevated LDL-C level while being treated with maximally tolerated statin therapy or has an elevated LDL-C level and a contraindication/intolerance to statin therapy and 2) Patient must have a documented diagnosis of one of the following: a) Heterozygous Familial Hypercholesterolemia (HeFH) b) atherosclerotic cardiovascular disease or c) established cardiovascular disease (CVD) or is at high risk for a CVD event but without established CVD.
Age Restrictions	N/A
Prescriber Restrictions	The medication must be prescribed by or in consultation with a cardiologist, endocrinologist, lipidologist, or neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NILOTINIB CAPSULES

Products Affected

- Nilotinib D-tartrate
- Nilotinib Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Documented diagnosis of Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NINLARO

Products Affected

- Ninlaro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of multiple myeloma and the requested drug is being used in combination with lenalidomide and dexamethasone in patients who have received at least one prior therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NITISINONE

Products Affected

- Nitisinone

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of hereditary tyrosinemia type-1 (HT-1).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NUBEQA

Products Affected

- Nubeqa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of 1) non-metastatic castration-resistant prostate cancer or 2) metastatic castration-sensitive prostate cancer (mCSPC). For mCSPC, the requested drug is used as monotherapy or with docetaxel.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NUEDEXTA

Products Affected

- Dextromethorphan Hydrobromide/quinidine Sulfate
- Nuedexta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of pseudobulbar affect (PBA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NUPLAZID

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	The member must have a documented diagnosis of Parkinson's disease and have hallucinations and delusions associated with Parkinson's disease psychosis.
Age Restrictions	N/A
Prescriber Restrictions	The medication must be prescribed by or in consultation with a neurologist or psychiatrist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NURTEC

Products Affected

- Nurtec

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) acute migraine and has had an inadequate response, intolerance, or contraindication to at least one triptan medication OR 2) episodic migraine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ODOMZO

Products Affected

- Odomzo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of locally advanced basal cell carcinoma and one of the following: 1) Documentation of disease recurrence following surgery or radiation therapy or 2) Documentation that the member is not a candidate for surgery or radiation therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OFEV

Products Affected

- Nintedanib Esylate
- Ofev

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of one of the following: 1) idiopathic pulmonary fibrosis (IPF) 2) systemic sclerosis-associated interstitial lung disease (SSc-ILD) or 3) chronic fibrosing interstitial lung diseases (ILDs) with a progressive phenotype.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a pulmonologist or a rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OGSIVEO

Products Affected

- Ogsiveo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of progressing desmoid tumors and 2) Patient requires systemic treatment.
Age Restrictions	N/A
Prescriber Restrictions	The prescriber must be an oncologist or sarcoma specialist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OJEMDA

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. Disease has a BRAF fusion or rearrangement, or BRAF V600 mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OJJAARA

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 intermediate or high-risk myelofibrosis (MF), including primary MF or secondary MF [post-polycythemia vera (PV) and post-essential thrombocythemia (ET)], in adults with anemia.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ONUREG

Products Affected

- Onureg

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of acute myeloid leukemia and has achieved first complete remission (CR) or complete remission with incomplete blood count recovery (CRi) following intensive induction chemotherapy and are not able to complete intensive curative therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OPIPZA

Products Affected

- Opipza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of schizophrenia or 2) Diagnosis of major depressive disorder (MDD) and being used as adjunctive treatment or 3) Drug is being used for irritability associated with autistic disorder in pediatric patients or 4) Diagnosis of Tourette's disorder in pediatric patients
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a psychiatrist or a neurologist.
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORENCIA

Products Affected

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

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Rheumatoid Arthritis (RA): Diagnosis of moderately to severely active RA and a trial and failure, contraindication, or intolerance to one of the following conventional therapies: methotrexate, leflunomide, sulfasalazine. Polyarticular Juvenile Idiopathic Arthritis (PJIA): Diagnosis of moderately to severely active PJIA and has a trial and failure, contraindication, or intolerance to one of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine. Psoriatic Arthritis (PsA): Diagnosis of active psoriatic arthritis.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, rheumatologist, oncologist, or transplant specialist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ORGOVYX

Products Affected

- Orgovyx

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of advanced prostate cancer.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORKAMBI

Products Affected

- Orkambi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of cystic fibrosis (CF) and have documentation that the member is homozygous for the F508del mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORSERDU

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 advanced or metastatic breast cancer that is estrogen receptor (ER)-positive and human epidermal growth factor receptor 2 (HER2)-negative in a postmenopausal woman or adult man. Documentation that the member has estrogen receptor (ESR1) mutated disease. Disease has progressed following at least one line of endocrine therapy.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OTEZLA

Products Affected

- Otezla

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Psoriatic Arthritis (PsA): The member must have a documented diagnosis of psoriatic arthritis. Plaque psoriasis (PsO): The member must have a documented diagnosis of plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Oral ulcers associated with Behcet's Disease: The member must have a documented diagnosis of Behcet's Disease. The member has active oral ulcers.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PEMAZYRE

Products Affected

- Pemazyre

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) unresectable, locally advanced or metastatic Cholangiocarcinoma with a fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement and has been previously treated or 2) relapsed or refractory myeloid/lymphoid neoplasms (MLNs) with fibroblast growth factor receptor 1 (FGFR1) rearrangement.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PIQRAY

Products Affected

- Piqray 200mg Daily Dose
- Piqray 250mg Daily Dose
- Piqray 300mg Daily Dose

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet the following criteria: 1) The member must have a documented diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, PIK3CA-mutated, advanced or metastatic breast cancer. 2) The member has progressed on or after an endocrine-based regimen. 3) Piqray is being used in combination with fulvestrant.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PIRFENIDONE

Products Affected

- Pirfenidone TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of idiopathic pulmonary fibrosis (IPF).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a pulmonologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

POMALYST

Products Affected

- Pomalidomide
- Pomalyst

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Kaposi Sarcoma: The member must have a documented diagnosis Kaposi sarcoma (KS) or AIDS-related Kaposi sarcoma (KS) after failure of highly active antiretroviral therapy (HAART). Multiple Myeloma: The member must have a documented diagnosis of multiple myeloma and has received at least two prior therapies including lenalidomide and a proteasome inhibitor (including but not limited to: Kyprolis, Ninlaro, or Velcade) and has demonstrated disease progression on or within 60 days of completion of the last therapy AND the requested drug is being used in combination with dexamethasone.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PRALUENT

Products Affected

- Praluent

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet the following criteria: 1) Member has an elevated LDL-C level while being treated with a high-intensity statin (i.e. atorvastatin or rosuvastatin) or has an elevated LDL-C level and a contraindication/intolerance to statin therapy. 2) The member must have a documented diagnosis of one of the following: a) Cardiovascular disease b) Primary hyperlipidemia including Heterozygous Familial Hypercholesterolemia (HeFH) as confirmed by genetic testing or clinical criteria. c) Homozygous Familial Hypercholesterolemia (HoFH) as confirmed by genetic testing or clinical criteria.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PREVYMIS

Products Affected

- Prevymis PACK
- Prevymis TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have documentation of 1) having had, or is scheduled to receive, an allogeneic hematopoietic stem cell transplant (HSCT) and the member is at risk for cytomegalovirus (CMV) infection or 2) having had, or is scheduled to receive, a kidney transplant and the member is at risk for cytomegalovirus (CMV) infection
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PROLASTIN

Products Affected

- Prolastin-c INJ 1000MG/20ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of hereditary deficiency of alpha-1 antitrypsin with clinical evidence of emphysema.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

QINLOCK

Products Affected

- Qinlock

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of advanced gastrointestinal stromal tumor (GIST) who have received prior treatment with 3 or more kinase inhibitors, including imatinib.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

QUININE SULFATE

Products Affected

- Quinine Sulfate CAPS 324MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage will not be approved for the treatment or prevention of nocturnal leg cramps.
Required Medical Information	The member is using the medication for treatment of uncomplicated Plasmodium falciparum malaria.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

QULIPTA

Products Affected

- Qulipta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of migraines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RADICAVA ORAL SUSPENSION

Products Affected

- Radicava Ors
- Radicava Ors Starter Kit

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of amyotrophic lateral sclerosis (ALS).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REPATHA

Products Affected

- Repatha
- Repatha Pushtronex System
- Repatha Sureclick

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet the following criteria: 1) Member has an elevated LDL-C level while being treated with a high-intensity statin (i.e. atorvastatin or rosuvastatin) or has an elevated LDL-C level and a contraindication/intolerance to statin therapy. 2) The member a) has a documented diagnosis of Primary hyperlipidemia including Heterozygous Familial Hypercholesterolemia (HeFH) as confirmed by genetic testing or clinical criteria. b) has a documented diagnosis of Homozygous Familial Hypercholesterolemia (HoFH) as confirmed by genetic testing or clinical criteria or c) is at increased risk for major adverse cardiovascular (CV) events.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RETEVMO

Products Affected

- Retevmo

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: The member must have a documented diagnosis of metastatic RET fusion-positive non-small cell lung cancer (NSCLC). RET-mutant Medullary Thyroid Cancer: The member must have a documented diagnosis of advanced or metastatic RET-mutant medullary thyroid cancer (MTC) who require systemic therapy. RET Fusion-Positive Thyroid Cancer: The member must have a documented diagnosis of advanced or metastatic RET fusion-positive thyroid cancer who require systemic therapy and who are radioactive iodine-refractory (if radioactive iodine is appropriate). Solid Tumors: The member must have a documented diagnosis of locally advanced or metastatic solid tumors with a RET gene fusion that have progressed on or following prior systemic treatment or who have no satisfactory alternative treatment options.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RETINOIDS FOR THE TOPICAL TREATMENT OF ACNE VULGARIS AND PSORIASIS

Products Affected

- Adapalene GEL
- Tazarotene CREA
- Tazarotene GEL
- Tretinoin CREA

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage of topical acne products will not be authorized for cosmetic purposes.
Required Medical Information	For all retinoids, the member must have a documented diagnosis of acne vulgaris, comedones (white heads), or actinic keratosis. Tazarotene may also be covered if the member has a physician-documented diagnosis of plaque psoriasis or documented diagnosis of skin cancer provided effective treatment with tazarotene is recognized for treatment of such indication in one of the standard reference compendia, or in the medical literature.
Age Restrictions	This criterion only applies to members age 26 or older. Authorization is not required for members 25 years of age or younger.
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REVCOVI

Products Affected

- Revcovi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion 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	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REVUFORJ

Products Affected

- Revuforj

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of 1) relapsed or refractory acute leukemia with a lysine methyltransferase 2A gene (KMT2A) translocation or 2) relapsed or refractory acute myeloid leukemia with a susceptible nucleophosmin 1 (NPM1) mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZDIFFRA

Products Affected

- Rezdiffra

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: Diagnosis of metabolic dysfunction-associated steatohepatitis (MASH), formerly known as nonalcoholic steatohepatitis (NASH). Patient does not have cirrhosis (e.g., decompensated cirrhosis). Disease is fibrosis stage F2 or F3 as confirmed by one of the following: 1) Both of the following: A) Serum biomarker [e.g., enhanced liver fibrosis (ELF) test, fibrosis-4 index (FIB-4)], and B) Imaging biomarker [e.g., FibroScan, magnetic resonance imaging-proton density fat fraction (MRI-PDFF)], or 2) One of the following: A) FibroScan aspartate aminotransferase (FAST), B) MRI-aspartate aminotransferase (MAST), C) Magnetic Resonance Elastography combined with fibrosis-4 index (MEFIB), or D) Liver biopsy within the past 12 months. Presence of greater than or equal to 1 metabolic risk factor (e.g., Type 2 diabetes, hypertension, obesity).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist or hepatologist.
Coverage Duration	2 years
Other Criteria	Reauth: Patient demonstrates positive response to therapy. Patient has not progressed to cirrhosis.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZLIDHIA

Products Affected

- Rezlidhia

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of relapsed or refractory acute myeloid leukemia and documentation cancer has susceptible IDH1 mutation
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or a hematologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZUROCK

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): Diagnosis of cGVHD. Trial and failure of two or more lines of systemic therapy [e.g., corticosteroids (e.g., prednisone, methylprednisolone), mycophenolate].
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with one of the following: hematologist, oncologist, or physician experienced in the management of transplant patients.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RINVOQ

Products Affected

- Rinvoq
- Rinvoq Lq

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Ankylosing Spondylitis(AS): Diagnosis of active AS. Atopic Dermatitis: Diagnosis of refractory, moderate to severe atopic dermatitis with 1) a trial and failure of a minimum 30-day supply (or 14-day supply for topical corticosteroids), contraindication, or intolerance to at least one of the following: Medium or higher potency topical corticosteroid, pimecrolimus cream, tacrolimus ointment, crisaborole ointment and 2) a trial and failure of a minimum 12-week supply of at least one systemic drug product for the treatment of atopic dermatitis (examples include, but are not limited to, tralokinumab-ldrm, dupilumab, etc.). Psoriatic Arthritis: Diagnosis of psoriatic arthritis and has an inadequate response or intolerance to one or more TNF inhibitors. Rheumatoid Arthritis (RA): 1) Diagnosis of moderately to severely active RA and 2) has a trial and failure, contraindication, or intolerance (T/F/C/I) to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine and 3) has an inadequate response or intolerance to one or more TNF inhibitors. Ulcerative Colitis (UC): 1) Diagnosis moderately to severely active UC and 2) has an inadequate response or intolerance to one or more TNF inhibitors OR, if TNF blockers are clinically inadvisable, patients should have received at least one approved systemic therapy. Non-radiographic Axial Spondyloarthritis: Diagnosis of Non-radiographic Axial Spondyloarthritis and has an inadequate response or intolerance to TNF blocker therapy. Crohn's disease: 1) Diagnosis of moderate to severe CD and 2) inadequate response or intolerance to one of more TNF blockers OR, if TNF blockers are clinically inadvisable, patients should have received at least one approved systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a allergist/immunologist, dermatologist, gastroenterologist, or rheumatologist.

Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ROMVIMZA

Products Affected

- Romvimza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of symptomatic tenosynovial giant cell tumor (TGCT) and 2) Surgical resection will potentially cause worsening functional limitation or severe morbidity.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ROZLYTREK

Products Affected

- Rozlytrek

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Non-small Cell Lung Cancer (NSCLC): The member must have a documented diagnosis of metastatic NSCLC with ROS1-positive tumors. Solid Tumors: The member must have a documented diagnosis of solid tumors that 1) have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion without a known acquired resistance mutation, 2) are metastatic or where surgical resection is likely to result in severe morbidity, and 3) have progressed following treatment or have no satisfactory alternative therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RUBRACA

Products Affected

- Rubraca

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Rubraca will not be approved for concurrent use with other chemotherapy agents.
Required Medical Information	Recurrent Ovarian Cancer (maintenance): The member must have a documented diagnosis of deleterious BRCA mutation (germline and/or somatic)- associated recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer and is in a complete or partial response to platinum-based chemotherapy. Prostate Cancer: The member must have a documented diagnosis of deleterious BRCA mutation (germline and/or somatic)-associated metastatic castration-resistant prostate cancer (mCRPC) who have been treated with androgen receptor directed therapy and a taxane-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RYDAPT

Products Affected

- Rydapt

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Rydapt will not be approved as single-agent induction therapy for the treatment of patients with AML.
Required Medical Information	Acute Myeloid Leukemia (AML): The member must have a documented diagnosis of AML that is FLT3 mutation-positive and Rydapt is being used as first-line therapy in combination with standard cytarabine and daunorubicin induction and cytarabine consolidation. Mast Cell Leukemia (MCL): The member must have a documented diagnosis of MCL. Systemic Mastocytosis: The member must have a documented diagnosis of aggressive systemic mastocytosis (ASM) or systemic mastocytosis with associated hematological neoplasm (SM-AHN).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist or an allergist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SAPROPTERIN

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	The member must have a documented diagnosis of hyperphenylalaninemia (HPA) due to tetrahydrobiopterin (BH4)-responsive phenylketonuria (PKU).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a specialist in metabolic diseases or a geneticist.
Coverage Duration	Initial authorization is for 8 weeks. Subsequent authorization is for 2 years.
Other Criteria	Coverage will be authorized for continuing therapy if the member has experienced improvement.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SCEMBLIX

Products Affected

- Scemblix

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP) and is either 1) newly diagnosed or 2) previously treated or 3) has a documented T315I mutation.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SIGNIFOR

Products Affected

- Signifor

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Cushing's disease and pituitary surgery is not an option or has not been curative.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an endocrinologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SIRTURO

Products Affected

- Sirturo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of pulmonary multi-drug resistant tuberculosis (MDR-TB).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SKYRIZI

Products Affected

- Skyrizi INJ 150MG/ML, 180MG/1.2ML, 360MG/2.4ML, 55MG/0.37ML, 600MG/10ML
- Skyrizi Pen

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis. Crohn's disease: 1) diagnosis of moderate to severe CD. Ulcerative Colitis: Diagnosis of moderately to severely active UC .
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, rheumatologist, or gastroenterologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SODIUM OXYBATE

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	Diagnosis of 1) narcolepsy with cataplexy or excessive daytime sleepiness (EDS) or 2) idiopathic hypersomnia
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SOMAVERT

Products Affected

- Somavert

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of acromegaly and has had a failure of, or is unable to tolerate, a treatment regimen that includes octreotide, and the member is not a candidate for or has had an inadequate response to surgery and/or radiation.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an endocrinologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SORAFENIB

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	Advanced Renal Cell Carcinoma (ARCC): The member must have a documented diagnosis of ARCC. Hepatocellular Carcinoma (HCC): The member must have a documented diagnosis of unresectable HCC. Thyroid Carcinoma (TC): The member must have a documented diagnosis of locally recurrent or metastatic, progressive, differentiated TC refractory to radioactive iodine treatment.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a nephrologist, oncologist, or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

STELARA

Products Affected

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

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): The member must have a documented diagnosis of moderate to severe CD. Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis. Ulcerative Colitis (UC): The member must have a documented diagnosis of moderate to severe UC. For all indications: The member must also have a trial, intolerance, or contraindication to Yesintek or Steqeyma.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, gastroenterologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

STIVARGA

Products Affected

- Stivarga

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Gastrointestinal Stromal Tumors (GIST): Diagnosis of locally advanced, unresectable, or metastatic GIST and documented failure, contraindication, or intolerance to both imatinib mesylate and sunitinib malate. Hepatocellular Carcinoma: Diagnosis of hepatocellular carcinoma and had a documented failure, contraindication, or intolerance to sorafenib. Metastatic Colorectal Cancer (MCC): Diagnosis of MCC and has been previously treated with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy, an anti-vascular endothelial growth factor (VEGF) therapy, and, if RAS wild type, an anti-epidermal growth factor receptor (EGFR) therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SUNITINIB

Products Affected

- Sunitinib Malate

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Advanced Renal Cell Carcinoma (ARCC): Diagnosis of ARCC. Gastrointestinal Stromal Tumor (GIST): Diagnosis of GIST and has a demonstrated disease progression or intolerance with imatinib mesylate. Progressive Neuroendocrine Tumors (pNET): Diagnosis of unresectable, locally advanced, or metastatic pNET located in the pancreas. Renal Cell Carcinoma (RCC): Diagnosis of RCC with high risk of recurrence following nephrectomy
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TABRECTA

Products Affected

- Tabrecta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) with a mutation that leads to mesenchymal-epithelial transition (MET) exon 14 skipping.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TADALAFIL

Products Affected

- Tadalafil TABS 2.5MG, 5MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Tadalafil is excluded from coverage for the treatment of Erectile Dysfunction.
Required Medical Information	The member must have a documented diagnosis or signs and symptoms of Benign Prostatic Hyperplasia (BPH) and has had a documented failure, adverse reaction, or contraindication to a 30-day trial of at least two of the following medications: alfuzosin, doxazosin, dutasteride, dutasteride-tamsulosin, finasteride, tamsulosin, or terazosin.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TAFINLAR

Products Affected

- Tafinlar

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Tafinlar is not indicated for the treatment of patients with wild-type BRAF mutations.
Required Medical Information	Single Agent: The member must have a documented diagnosis of unresectable or metastatic melanoma with a BRAF V600E mutation. In Combination with trametinib: The member must have a documented diagnosis of one of the following: 1) Unresectable or metastatic melanoma with a BRAF V600E or V600K mutation. 2) Melanoma with a BRAF V600E or V600K mutation and involvement of lymph node(s) following complete resection. 3) Metastatic non-small cell lung cancer (NSCLC) with a BRAF V600E mutation. 4) Locally advanced or metastatic anaplastic thyroid cancer (ATC) with a BRAF V600E mutation with no satisfactory locoregional treatment options. 5) Unresectable or metastatic solid tumors with BRAF V600E mutation and has progressed following prior treatment and have no satisfactory alternative treatment options 6) low grade glioma with BRAF V600E mutation that requires systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAGRISSO

Products Affected

- Tagrisso

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) metastatic non-small cell lung cancer (NSCLC) whose tumors have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations and the requested drug is being used first line OR 2) metastatic EGFR T790M mutation-positive NSCLC whose disease has progressed on or after EGFR tyrosine kinase inhibitor (TKI) therapy or 3) NSCLC whose tumors have EGFR exon 19 deletions or exon 21 L858R mutations and Tagrisso is being used as a) adjuvant therapy after tumor resection b) first-line treatment in combination with pemetrexed and platinum-based chemotherapy or 4) locally advanced, unresectable (stage III) NSCLC and disease has not progressed during or following concurrent or sequential platinum-based chemoradiation therapy and whose tumors have EGFR exon 19 deletions or exon 21 L858R mutations
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TALZENNA

Products Affected

- Talzenna

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) deleterious or suspected deleterious germline BRCA-mutated, HER2-negative locally advanced or metastatic breast cancer or 2) HRR gene-mutated metastatic castration-resistant prostrate cancer (mCRPC) and is being used in combination with enzalutamide.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TASIMELTEON

Products Affected

- Tasimelteon

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage will not be authorized for the diagnosis of insomnia.
Required Medical Information	The member must have a documented diagnosis of Smith-Magenis Syndrome (SMS) and be experiencing nighttime sleep disturbances or the member must be completely blind and have a documented diagnosis of non-24-hour sleep-wake disorder (non-24).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist or sleep specialist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAVNEOS

Products Affected

- Tavneos

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) 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) and 2) Patient is receiving concurrent immunosuppressant therapy with one of the following: a) cyclophosphamide OR b) rituximab and 3) 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	Prescribed by or in consultation with a nephrologist, pulmonologist, or rheumatologist
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TAZVERIK

Products Affected

- Tazverik

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have one of the following requirements: 1) The member must have a documented diagnosis of metastatic or locally advanced epithelioid sarcoma not eligible for complete resection. 2) The member must have a documented diagnosis of relapsed or refractory follicular lymphoma whose tumors are positive for an EZH2 mutation and has received at least two prior systemic therapies. 3) The member must have a documented diagnosis of relapsed or refractory follicular lymphoma who have no satisfactory alternative treatment options.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TEPMETKO

Products Affected

- Tepmetko

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) harboring mesenchymal-epithelial transition (MET) exon 14 skipping alterations.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TERIPARATIDE

Products Affected

- Bonsity
- Teriparatide

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Documentation of 1) a postmenopausal woman with osteoporosis 2) a man with primary or hypogonadal osteoporosis or 3) man or woman with osteoporosis associated with sustained systemic glucocorticoid therapy. For either condition previously listed, the patient must be at high risk of fracture and must have had an inadequate response to, or is unable to tolerate therapy with at least one of the traditional osteoporosis treatments (including but not limited to: alendronate, calcitonin, denosumab, ibandronate, raloxifene, risedronate or zoledronic acid.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TETRABENAZINE

Products Affected

- Tetrabenazine

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of chorea associated with Huntington's Disease.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TIBSOVO

Products Affected

- Tibsovo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Relapsed or Refractory Acute Myeloid Leukemia (AML): The member must have a documented diagnosis of relapsed or refractory AML with a susceptible isocitrate dehydrogenase-1 (IDH1) mutation. Acute Myeloid Leukemia (AML): 1) The member must have a new diagnosis of AML with a susceptible isocitrate dehydrogenase-1 (IDH1) mutation and 2) Requested drug is being used as monotherapy or in combination with azacitidine and the member meets one of the following: a) is 75 years of age or older or b) has comorbidities that make them ineligible for intensive induction chemotherapy. Cholangiocarcinoma: The member must have a documented diagnosis of locally advanced or metastatic cholangiocarcinoma who have been previously treated. Relapsed or refractory Myelodysplastic Syndromes (MDS): The member must have a documented diagnosis of MDS
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TRANSMUCOSAL IMMEDIATE-RELEASE FENTANYL (TIRF)

Products Affected

- Fentanyl Citrate Oral Transmucosal

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	The Transmucosal Immediate-Release Fentanyl (TIRF) products will not be covered for any non-cancer pain indication.
Required Medical Information	The Transmucosal Immediate-Release Fentanyl (TIRF) products will be covered for the management of breakthrough pain in adult cancer patients who are already receiving and who are tolerant to around-the-clock opioid therapy for their underlying persistent cancer pain. (applies to Fentanyl lozenges only) Approvable for pediatric patients 16 years of age and older.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or a pain management specialist.
Coverage Duration	2 years
Other Criteria	Patients considered opioid tolerant are those who are taking around-the-clock medicine consisting of, but not limited to, morphine oral 60 mg daily or more, fentanyl transdermal 25 mcg/hour or more, oxycodone oral 30 mg daily or more, hydromorphone oral 8 mg daily or more, or an equianalgesic dose of another opioid daily for a week or longer. Patients must remain on around-the-clock opioids when taking fentanyl transmucosal.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRIKAFTA

Products Affected

- Trikafta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Cystic Fibrosis (CF) (initial): Diagnosis of CF. Patient has at least one of the following mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene: F508del mutation OR a mutation in the CFTR gene that is responsive based on clinical and/or in vitro data.
Age Restrictions	CF (initial): For granule packets: patient is at least 2 to less than 6 years of age. For tablets: patient is 6 years of age or older.
Prescriber Restrictions	CF (initial): Prescribed by or in consultation with a pulmonologist or specialist affiliated with a CF care center.
Coverage Duration	2 years
Other Criteria	CF (reauth): Patient demonstrates positive clinical response to therapy (e.g., improvement in lung function [percent predicted forced expiratory volume in one second {PPFEV1}] or decreased number of pulmonary exacerbations).
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRUQAP

Products Affected

- Truqap

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis of breast cancer. Disease is one of the following: locally advanced or metastatic. Will be taken in combination with fulvestrant. Disease is hormone receptor (HR)-positive. Disease is human epidermal growth factor receptor 2 (HER2)-negative. Patient has one or more PIK3CA/AKT1/PTEN-alterations. One of the following: A) Following progression on at least one endocrine-based regimen in the metastatic setting (e.g., anastrozole, letrozole, exemestane, tamoxifen, etc.) OR B) Recurrence on or within 12 months of completing adjuvant therapy (e.g., chemotherapy). OR 2) Diagnosis of prostate cancer. Disease is metastatic, PTEN-deficient, androgen pathway modulation-naive or sensitive. Used in combination with abiraterone and prednisone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TUKYSA

Products Affected

- Tukysa

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) advanced unresectable or metastatic HER2-positive breast cancer, including patients with brain metastases, who have received one or more prior anti-HER2-based regimens in the metastatic setting and be taking in combination with trastuzumab and capecitabine or 2) RAS wild-type HER-2 positive unresectable or metastatic colorectal cancer that has progressed following treatment with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy and be taking in combination trastuzumab.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TURALIO

Products Affected

- Turalio CAPS 125MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of symptomatic tenosynovial giant cell tumor (TGCT) associated with severe morbidity or functional limitations and the condition is not amenable to improvement with surgery.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TYENNE SC

Products Affected

- Tyenne INJ 162MG/0.9ML

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion 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, Orencia (abatacept), Rinvoq (upadacitinib), Xeljanz/Xeljanz XR (tofacitinib) or attestation demonstrating a trial may be inappropriate, OR b) For continuation of prior therapy. Giant Cell Arteritis (GCA) (Initial): Diagnosis of GCA. TF/C/I to a glucocorticoid (eg, prednisone). 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), minimum duration of a 3-month trial of methotrexate, 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, Orencia (abatacept), Rinvoq/LQ, or Xeljanz (tofacitinib), OR b) for continuation of prior therapy. Systemic sclerosis-associated interstitial lung disease (SSc-ILD) (Initial): Diagnosis of SSc-ILD as documented by the following: a) Exclusion of other known causes of ILD AND b) One of the following: i) In patients not subjected to surgical lung biopsy, the presence of idiopathic interstitial pneumonia (eg, fibrotic nonspecific interstitial pneumonia [NSIP], usual interstitial pneumonia [UIP] and centrilobular fibrosis) pattern on high-resolution computed tomography (HRCT) revealing SSc-ILD or probable SSc-ILD, OR ii) In patients subjected to a lung biopsy, both HRCT and surgical lung biopsy pattern revealing SSc-ILD or probable SSc-ILD.
Age Restrictions	N/A

Prescriber Restrictions	RA, GC, SJIA, PJIA (initial): Prescribed by or in consultation with a rheumatologist. SSc-ILD (initial): Prescribed by or in consultation with a pulmonologist or rheumatologist.
Coverage Duration	1 year
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. GC, SSc-ILD (Reauth): Patient demonstrates positive clinical response to therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

UBRELVY

Products Affected

- Ubrelvy

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of migraines and has had an inadequate response, intolerance, or contraindication to at least one triptan medication.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

USTEKINUMAB

Products Affected

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

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): The member must have a documented diagnosis of moderate to severe CD. Plaque Psoriasis: The member must have a documented diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: The member must have a documented diagnosis of psoriatic arthritis. Ulcerative Colitis (UC): The member must have a documented diagnosis of moderate to severe UC. For all indications: The member must also have a trial, intolerance, or contraindication to Yesintek or Steqeyma.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, gastroenterologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VALTOCO

Products Affected

- Valtoco 10 Mg Dose
- Valtoco 15 Mg Dose
- Valtoco 20 Mg Dose
- Valtoco 5 Mg Dose

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of a seizure disorder requiring acute treatment.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VANFLYTA

Products Affected

- Vanflyta

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of newly diagnosed Acute Myeloid Leukemia (AML) that is FLT3 internal tandem duplication (ITD) positive.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or a hematologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VENCLEXTA

Products Affected

- Venclexta
- Venclexta Starting Pack

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Acute Myeloid Leukemia (AML): The member must have a documented diagnosis of AML and the requested drug is being used as first-line therapy in combination with azacitidine, decitabine, or low-dose cytarabine and the member meets one of the following: 1) is 75 years of age or older or 2) has comorbidities that make them ineligible for intensive induction chemotherapy. Chronic Lymphocytic Leukemia (CLL) or Small Lymphocytic Lymphoma (SLL): The member must have a documented diagnosis of CLL or SLL.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VEOZAH

Products Affected

- Veozah

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of moderate to severe vasomotor symptoms due to menopause.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VERZENIO

Products Affected

- Verzenio

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For Monotherapy: Diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer with disease progression following endocrine therapy and prior chemotherapy. For Combination Therapy with Faslodex (fulvestrant): Diagnosis of HR-positive, HER2-negative advanced or metastatic breast cancer with disease progression following endocrine therapy. For Combination Therapy with an Aromatase Inhibitor as initial endocrine-based therapy: Diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer. Combination with Endocrine Therapy (tamoxifen or an aromatase inhibitor): Diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, node-positive, early breast cancer at high risk of recurrence.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VIGAFYDE

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.
Age Restrictions	Patient is 1 month to 2 years of age.
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VITRAKVI

Products Affected

- Vitrakvi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic or surgically unresectable neurotrophic receptor tyrosine kinase (NTRK) gene fusion-positive solid tumors with no known acquired resistance mutation and with no satisfactory alternative treatments or the member has progressed following treatment.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VIZIMPRO

Products Affected

- Vizimpro

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of metastatic non-small cell lung cancer (NSCLC) in tumors that have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutations.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VONJO

Products Affected

- Vonjo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of intermediate or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis with a platelet count less than 50 × 10^9/L.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOQUEZNA

Products Affected

- Voquezna
- Voquezna Dual Pak
- Voquezna Triple Pak

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Helicobacter pylori (H. pylori) (Voquezna Dual Pak, Voquezna Triple Pak): Diagnosis of H. pylori infection. Trial and failure, contraindication, or intolerance to bismuth quadruple therapy (e.g., bismuth and metronidazole and tetracycline and proton pump inhibitor [PPI]). H. pylori (Voquezna): Diagnosis of H. pylori infection. One of the following: a) Used in combination with amoxicillin and clarithromycin for the treatment of H. pylori infection, or b) Used in combination with amoxicillin for the treatment of H. pylori infection. Trial and failure, contraindication, or intolerance to bismuth quadruple therapy (e.g., bismuth and metronidazole and tetracycline and proton pump inhibitor [PPI]). Healing and Relief of Heartburn associated with Erosive Esophagitis (HRH) (Voquezna): Diagnosis of erosive esophagitis. Used for healing of all grades of erosive esophagitis and relief of heartburn associated with erosive esophagitis. Trial and inadequate response, contraindication, or intolerance to TWO of the following generic PPIs: a) omeprazole, b) esomeprazole, c) pantoprazole, d) lansoprazole, e) rabeprazole, or f) dexlansoprazole. Maintenance of Healing and Relief of Heartburn associated with Erosive Esophagitis (MHRH) (Voquezna): Used to maintain healing and relief of heartburn associated with erosive esophagitis. Trial and inadequate response, contraindication, or intolerance to TWO of the following generic PPIs: a) omeprazole, b) esomeprazole, c) pantoprazole, d) lansoprazole, e) rabeprazole, or f) dexlansoprazole.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	H. pylori, NERD: 1 mo. HRH: 2 months. MHRH: 6 mos.

Other Criteria	Relief of Heartburn associated with Non-Erosive Gastroesophageal Reflux Disease (NERD): Diagnosis of non-erosive Gastroesophageal Reflux Disease. Both of the following: a) Patient has history of heartburn for at least 6 months and b) Heartburn symptoms are present for at least 4 days during any consecutive 7-day period. Trial and inadequate response, contraindication, or intolerance to TWO of the following generic PPIs: a) omeprazole, b) esomeprazole, c) pantoprazole, d) lansoprazole, e) rabeprazole, or f) dexlansoprazole.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VORANIGO

Products Affected

- Voranigo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of grade 2 astrocytoma or oligodendroglioma. Presence of a susceptible isocitrate dehydrogenase-1 (IDH1) or isocitrate dehydrogenase-2 (IDH2) mutation. History of one of the following: a) Biopsy, b) Sub-total resection, or c) Gross total resection.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of therapy
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VORICONAZOLE FOR IV INJECTION

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	The member must have a documented diagnosis of 1) infection caused by Aspergillus or Candida, including candidemia or other serious invasive candidiasis infection, invasive aspergillosis, CNS infection (i.e. meningitis), cardiovascular system infection (i.e. endocarditis, myocarditis, pericarditis, infected pacemaker, implantable cardiac defibrillator, or ventricular assist devices), esophageal candidiasis, invasive pulmonary aspergillosis and other Aspergillus respiratory infection (i.e. pneumonia, tracheobronchitis, sinusitis, aspergilloma), intrabdominal infections, bone and joint infection, fungal skin and skin structure infection or 2) serious fungal infection caused by Scedosporium apiospermum or Fusarium and intolerant of, or refractory to, other therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOSEVI

Products Affected

- Vosevi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Criteria will be applied consistent with current AASLD-IDSA guidance.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a gastroenterologist, hepatologist, or an infectious disease specialist.
Coverage Duration	Criteria will be applied consistent with current AASLD-IDSA guidance.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOTRIENT

Products Affected

- Pazopanib Hydrochloride

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Advanced Renal Cell Carcinoma (RCC): The member must have a documented diagnosis of advanced RCC. Advanced Soft Tissue Sarcoma (STS): The member must have a documented diagnosis of advanced STS and has received prior chemotherapy, including anthracycline treatment, or was unsuited for such therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VOWST

Products Affected

- Vowst

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of recurrent clostridioides difficile infection (CDI) and the patient has completed one of the following antibiotic therapies 2-4 days prior to initiating Vowst: oral vancomycin or Dificid (fidaxomicin)
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist or infectious disease specialist.
Coverage Duration	14 days
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VYNDAMAX

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), 2) Cardiac or noncardiac tissue biopsy demonstrating histologic confirmation of TTR amyloid deposits, OR 3) Both of the following: i) cardiac magnetic resonance imaging or scintigraphy scan suggestive of amyloidosis, and ii) absence of light-chain amyloidosis. Documentation that patient has New York Heart Association (NYHA) Functional Class I, II, or III heart failure.
Age Restrictions	N/A
Prescriber Restrictions	ATTR-CM (initial, reauth): Prescribed by a cardiologist
Coverage Duration	ATTR-CM (initial, reauth): Plan year.
Other Criteria	ATTR-CM (reauth): Patient continues to have New York Heart Association (NYHA) Functional Class I, II, or III heart failure.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

WELIREG

Products Affected

- Welireg

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) von Hippel-Lindau disease and require therapy for associated renal cell carcinoma, CNS hemangioblastomas, or pancreatic neuroendocrine tumors, not requiring immediate surgery or 2) Advanced Renal Cell Carcinoma (RCC) following a programmed death receptor-1 (PD-1) or programmed death-ligand 1 (PD-L1) inhibitor and a vascular endothelial growth factor tyrosine kinase inhibitor (VEGF-TKI) or 3) locally advanced, unresectable, or metastatic pheochromocytoma or paraganglioma (PPGL).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

WYOST

Products Affected

- Wyost

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Coverage for Wyost (denosumab) will be authorized if one of the following is met: 1) for prevention of skeletal-related events in patients with multiple myeloma or with bone metastases from solid tumors 2) the member is being treated for unresectable giant cell tumor of bone (GCTB) or surgical resection of GCTB is likely to result in severe morbidity 3) for the treatment of hypercalcemia of malignancy refractory to bisphosphonate therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XALKORI

Products Affected

- Xalkori

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK)-positive or the member has documented ROS1-positive tumors, 2) relapsed or refractory, systemic anaplastic large cell lymphoma (ALCL) that is ALK-positive, or 3) unresectable, recurrent, or refractory inflammatory myofibroblastic tumor (IMT) that is ALK-positive.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XDEMVY

Products Affected

- Xdemvy

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of Demodex blepharitis.
Age Restrictions	N/A
Prescriber Restrictions	The medication must be prescribed by or in consultation with an ophthalmologist or optometrist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XELJANZ

Products Affected

- Tofacitinib Citrate SOLN
- Tofacitinib Citrate TABS
- Tofacitinib Citrate Er
- Xeljanz
- Xeljanz Xr

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Ankylosing spondylitis: Diagnosis of active ankylosing spondylitis and has an inadequate response or intolerance to one or more TNF inhibitors (e.g., Enbrel, Humira). Rheumatoid Arthritis (RA): 1) Diagnosis of moderately to severe active RA and 2) has a trial and failure, contraindication, or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine and 3) has an inadequate response or intolerance to one or more TNF inhibitors (e.g., Enbrel, Humira). Polyarticular Juvenile Idiopathic Arthritis (PJIA): 1) Diagnosis of active PJIA and 2) has a trial and failure, contraindication, or intolerance to ONE of the following conventional therapies: methotrexate, leflunomide, or sulfasalazine and 3) has an inadequate response or intolerance to one or more TNF inhibitors (e.g., Enbrel, Humira). Psoriatic Arthritis: Diagnosis of active psoriatic arthritis and has an inadequate response or intolerance to one or more TNF inhibitors (e.g., Enbrel, Humira). Ulcerative Colitis (UC): 1) Diagnosis of moderately to severely active UC and 2) has an inadequate response or intolerance to one or more TNF inhibitors (e.g., Humira).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A

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

XERMELO

Products Affected

- Xermelo

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of carcinoid syndrome diarrhea that is inadequately controlled by somatostatin analog (SSA) therapy alone and Xermelo is being used in combination with an SSA (e.g. Sandostatin LAR).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a gastroenterologist, hematologist, or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XIFAXAN 550 MG

Products Affected

- Xifaxan TABS 550MG

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Hepatic Encephalopathy: The member must have a documented diagnosis of hepatic encephalopathy and has had an inadequate response or a contraindication to lactulose. Irritable Bowel Syndrome with Diarrhea (IBS-D): The member must have a documented diagnosis of IBS-D.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XOLAIR

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: The member must 1) have a documented diagnosis of moderate-to-severe persistent asthma 2) has had a failure of a treatment regimen that included two or more of the following medications: inhaled corticosteroids, oral corticosteroids, leukotriene modifiers and inhaled long-acting bronchodilators, or is unable to tolerate these medications. 3) shows a definitive sensitivity on allergy testing to one or more perennial allergens and 4) The member has a pre-treatment serum IgE level equal to or greater than 30 IU/mL and less than or equal to 1,300 IU/mL. Chronic Spontaneous Urticaria (CSU): 1) The member has a documented diagnosis of CSU and 2) the physician has documented that the member remains symptomatic despite H1 antihistamine treatment. Nasal polyps: The member must have a documented diagnosis of nasal polyps with inadequate response to nasal corticosteroids. IgE-mediated food allergy: For the reduction of allergic reactions (Type I), including anaphylaxis, that may occur with accidental exposure to one or more foods.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an allergist, dermatologist, immunologist, otolaryngologist or pulmonologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XOSPATA

Products Affected

- Xospata

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of relapsed or refractory acute myeloid leukemia (AML) with a FLT3 mutation.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XPOVIO

Products Affected

- Xpovio
- Xpovio 60 Mg Twice Weekly
- Xpovio 80 Mg Twice Weekly

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	In combination with dexamethasone: The member must meet ALL of the following criteria: 1) Documented diagnosis of relapsed or refractory multiple myeloma. 2) Has received at least four prior therapies. 3) The member's disease is refractory to at least two proteasome inhibitors, at least two immunomodulatory agents, and an anti-CD38 monoclonal antibody. In combination with (Velcade) bortezomib and dexamethasone: The member must have a documented diagnosis of multiple myeloma and has received at least one prior therapy. Relapsed or Refractory Diffuse Large B-cell Lymphoma (DLBCL): The member must have a documented diagnosis of DLBCL, not otherwise specified, including DLBCL arising from follicular lymphoma, and has received at least 2 lines of systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XTANDI

Products Affected

- Xtandi

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of 1) non-metastatic castration-resistant prostate cancer or 2) metastatic hormone-sensitive prostate cancer and the requested drug is being used in combination with docetaxel.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or urologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

YESINTEK AND STEQEYMA

Products Affected

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

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. Plaque Psoriasis: Diagnosis of moderate-to-severe plaque psoriasis and has failed to respond to or has been unable to tolerate treatment with one of the following: Corticosteroids (e.g., betamethasone, clobetasol, desonide), Vitamin D analogs (e.g., calcitriol, calcipotriene), Tazarotene, Calcineurin inhibitors (e.g., tacrolimus, pimecrolimus), Anthralin, or Coal tar. Psoriatic Arthritis: Diagnosis of active psoriatic arthritis. Ulcerative Colitis (UC): Diagnosis of moderately to severely active UC.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, gastroenterologist or rheumatologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZEJULA

Products Affected

- Zejula TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet one of the two following requirements: 1) The member must have a documented diagnosis of advanced or recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer and is experiencing complete or partial response to platinum-based chemotherapy 2) The member must have a documented diagnosis of recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer with a deleterious or suspected deleterious germline BRCA mutation and who are in a complete or partial response to platinum-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZELBORAF

Products Affected

- Zelboraf

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	Zelboraf is not indicated for treatment of patients with wild-type BRAF melanoma.
Required Medical Information	Erdheim-Chester Disease (ECD): The member must have a documented diagnosis of ECD with a BRAF V600 mutation. Unresectable or Metastatic Melanoma: The member must have a documented diagnosis of unresectable or metastatic melanoma that is BRAF V600E mutation-positive.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist or a hematologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZENBEXUS

Products Affected

- Zenbexus

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of multiple myeloma. Patient has received at least one prior line of therapy including both of the following: a) One proteasome inhibitor (e.g., bortezomib), and b) One immunomodulatory agent (e.g., pomalidomide, lenalidomide). Used in combination with both of the following: a) Darzalex (daratumumab) or Darzalex Faspro (daratumumab and hyaluronidase-fihj), and b) Dexamethasone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	2 years
Other Criteria	Approve for continuation of prior therapy.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZOLINZA

Products Affected

- Zolinza

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of cutaneous T-cell lymphoma with progressive, persistent or recurrent disease and documented current or prior treatment or treatment failure with at least two systemic chemotherapeutic agents for cutaneous T-cell lymphoma.
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZTALMY

Products Affected

- Ztalmy

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a diagnosis of seizures associated with cyclin-dependent kinase-like 5 (CDKL5) deficiency disorder (CDD).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZURZUVAE

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 postpartum Depression
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.

ZYDELIG

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): The member must have a documented diagnosis of relapsed CLL and Zydelig is being used in combination with Rituxan (rituximab).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be a hematologist or oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZYKADIA

Products Affected

- Zykadia TABS

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a documented diagnosis of anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC).
Age Restrictions	N/A
Prescriber Restrictions	The prescribing physician must be an oncologist.
Coverage Duration	2 years
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PART B VERSUS PART D

Products Affected

- Abelcet
- Acetylcysteine SOLN
- Acyclovir Sodium INJ 50MG/ML
- Albuterol Sulfate NEBU 0.083%, 0.63MG/3ML, 1.25MG/3ML, 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, 993MG/100ML; 1018MG/100ML; 700MG/100ML; 738MG/100ML; 500MG/100ML; 300MG/100ML; 660MG/100ML; 1000MG/100ML; 1050MG/100ML; 172MG/100ML; 270MG/100ML; 298MG/100ML; 722MG/100ML; 530MG/100ML; 400MG/100ML; 200MG/100ML; 500MG/100ML
- Aminosyn-pf INJ 46MEQ/L; 698MG/100ML; 1227MG/100ML; 527MG/100ML; 820MG/100ML; 385MG/100ML; 312MG/100ML; 760MG/100ML; 1200MG/100ML; 677MG/100ML; 180MG/100ML; 427MG/100ML; 812MG/100ML; 495MG/100ML; 70MG/100ML; 512MG/100ML; 180MG/100ML; 44MG/100ML; 673MG/100ML
- Aminosyn-pf 7% INJ 32.5MEQ/L; 490MG/100ML; 861MG/100ML; 370MG/100ML; 576MG/100ML; 270MG/100ML; 220MG/100ML; 534MG/100ML; 831MG/100ML; 475MG/100ML; 125MG/100ML; 300MG/100ML; 570MG/100ML; 347MG/100ML; 50MG/100ML; 360MG/100ML; 125MG/100ML; 44MG/100ML; 452MG/100ML
- Amphotericin B INJ
- Amphotericin B Liposome
- Aprepitant
- Arformoterol Tartrate
- Azathioprine TABS
- Bivigam INJ 10%, 5GM/50ML
- Budesonide SUSP
- Clinimix 6/5
- Clinimix 8/10
- Clinimix E 8/10
- Cromolyn Sodium NEBU
- Cuvitru
- Cyclophosphamide CAPS
- Cyclophosphamide TABS
- Cyclosporine CAPS
- Cyclosporine Modified
- Dronabinol CAPS
- Engerix-b
- Envarsus Xr
- Everolimus TABS 0.25MG, 0.5MG, 0.75MG, 1MG
- Flebogamma Dif INJ 10GM/100ML, 10GM/200ML, 2.5GM/50ML, 20GM/200ML, 20GM/400ML, 5GM/100ML, 5GM/50ML
- Formoterol Fumarate NEBU
- Gammagard Liquid INJ 10GM/100ML, 2.5GM/25ML, 20GM/200ML, 30GM/300ML, 5GM/50ML

- Gammaplex INJ 10GM/100ML, 10GM/200ML, 20GM/200ML, 20GM/400ML, 5GM/100ML, 5GM/50ML
- Gengraf CAPS 100MG, 25MG
- Gengraf SOLN
- Granisetron Hydrochloride TABS
- Heplisav-b
- Hizentra
- Imovax Rabies (h.d.c.v.)
- Intralipid INJ 20GM/100ML, 30GM/100ML
- Ipratropium Bromide INHALATION SOLN 0.02%
- Ipratropium Bromide/albuterol Sulfate
- Levalbuterol NEBU
- Levalbuterol Hcl NEBU
- Levalbuterol Hydrochloride NEBU 0.63MG/3ML
- Mycophenolate Mofetil CAPS
- Mycophenolate Mofetil SUSR
- Mycophenolate Mofetil TABS
- Mycophenolic Acid Dr
- Nutrilipid
- Octagam INJ 10GM/100ML, 10GM/200ML, 1GM/20ML, 2.5GM/50ML, 20GM/200ML, 2GM/20ML, 30GM/300ML, 5GM/100ML, 5GM/50ML
- Ondansetron Hcl SOLN 4MG/5ML
- Ondansetron Hcl TABS 24MG
- 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
- Prehevbrio
- Premasol INJ 52MEQ/L; 1760MG/100ML; 880MG/100ML; 34MEQ/L; 1760MG/100ML; 372MG/100ML; 406MG/100ML; 526MG/100ML; 492MG/100ML; 492MG/100ML; 526MG/100ML; 356MG/100ML; 356MG/100ML; 390MG/100ML; 34MG/100ML; 152MG/100ML
- Privigen
- Prograf PACK
- Prosol
- Pulmozyme SOLN 2.5MG/2.5ML
- Rabavert
- Recombivax Hb
- Sirolimus SOLN
- Sirolimus TABS
- Tacrolimus CAPS
- Travasol INJ 52MEQ/L; 1760MG/100ML; 880MG/100ML; 34MEQ/L; 1760MG/100ML; 372MG/100ML; 406MG/100ML; 526MG/100ML; 492MG/100ML; 492MG/100ML; 526MG/100ML; 356MG/100ML; 500MG/100ML; 356MG/100ML; 390MG/100ML; 34MG/100ML; 152MG/100ML
- Trophamine INJ 0.54GM/100ML; 1.2GM/100ML; 0.32GM/100ML; 0; 0; 0.5GM/100ML; 0.36GM/100ML; 0.48GM/100ML; 0.82GM/100ML; 1.4GM/100ML; 1.2GM/100ML; 0.34GM/100ML; 0.48GM/100ML; 0.68GM/100ML; 0.38GM/100ML; 5MEQ/L; 0.025GM/100ML; 0.42GM/100ML; 0.2GM/100ML; 0.24GM/100ML; 0.78GM/100ML

Details

This drug may be covered under Medicare Part B or D depending upon the circumstances. Information may need to be submitted describing the use and setting of the drug to make the determination.

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