Mass General Brigham
Health Plan

2027 One Care Prior Authorization Medical Necessity Guidelines

Effective: January 1, 2027

For more recent information or other questions, please contact Mass General Brigham Health Plan Member Services.
Visit **MGBAdvantage.org/dsnp/rx-information** for the most up to date information on Medicare Part D drug coverage.

You can reach Member Services
by calling: **888-816-6000** (TTY: 711)

October 1 – March 31
8:00 AM to 8:00 PM EST, Monday through Sunday

April 1 – September 30
8:00 AM to 8:00 PM EST, Monday through Friday

Mass General Brigham One Care (HMO D-SNP)

Mass General Brigham Health Plan is a Medicare Advantage organization with a Medicare contract offering One Care D-SNP plan. Enrollment in Mass General Brigham Health Plan depends on contract renewal.

H1611_0452PHY_C

ABIRATERONE

Products Affected

- Abiraterone Acetate TABS 250MG, 500MG
- Abirtega

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Node-positive (N1), non-metastatic (M0) prostate cancer, very-high-risk prostate cancer, non-metastatic high-risk prostate cancer, non-metastatic prostate cancer with prostate-specific antigen (PSA) persistence/recurrence after radical prostatectomy, salivary gland tumors.
Exclusion Criteria	N/A
Required Medical Information	The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy. For salivary gland tumors: the requested drug is being used for the treatment of recurrent androgen receptor positive disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ACTIMMUNE

Products Affected

- Actimmune INJ 100MCG/0.5ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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-fkjp
- Hulio
- Yuflyma 1-pen Kit
- Yuflyma 2-syringe Kit
- Yuflyma Cd/uc/hs Starter

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

ADBRY

Products Affected

- Adbry

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ADEMPAS

Products Affected

- Adempas

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

AGAMREE

Products Affected

- Agamree

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

AIMOVIG

Products Affected

- Aimovig

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For preventive treatment of migraine, continuation: The member received at least 3 months of treatment with the requested drug and had a reduction in migraine days per month from baseline.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 3 months, Continuation: Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For deleterious or suspected deleterious BRCA2-mutated (BRCA2m) metastatic castration-sensitive prostate cancer (mCSPC): The requested drug will be used in combination with prednisone. For deleterious or suspected deleterious BRCA-mutated (BRCAm) metastatic castration-resistant prostate cancer (mCRPC):The requested drug will be used in combination with prednisone AND a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent anaplastic lymphoma kinase (ALK)-positive non-small cell lung cancer (NSCLC), brain metastases from ALK-positive NSCLC, ALK-positive anaplastic large cell lymphoma (ALCL), Erdheim-Chester Disease (ECD) with ALK-fusion, inflammatory myofibroblastic tumors (IMT) with ALK translocation (including advanced, recurrent/metastatic, or inoperable uterine sarcoma for IMT with ALK translocation), ALK-positive large B-cell lymphoma
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): 1) the member meets either of the following: a) the disease is recurrent, advanced, or metastatic OR b) the requested drug will be used as adjuvant treatment following tumor resection, AND 2) the disease is anaplastic lymphoma kinase (ALK)-positive.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALPHA1-PROTEINASE INHIBITOR

Products Affected

- Aralast Np INJ 1000MG, 500MG
- Prolastin-c INJ 1000MG/20ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Anit-IgA Antibodies
Required Medical Information	For alpha1-proteinase inhibitor deficiency: Member must have 1) clinically evident emphysema, AND 2) pretreatment serum alpha1-proteinase inhibitor level less than 11 micromol/L (80 milligrams per deciliter [mg/dL] by radial immunodiffusion or 50 mg/dL by nephelometry).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent anaplastic lymphoma kinase (ALK)-positive non-small cell lung cancer (NSCLC), brain metastases from ALK-positive NSCLC, ALK-positive anaplastic large cell lymphoma (ALCL), inflammatory myofibroblastic tumors (IMT) with ALK translocation (including advanced, recurrent/metastatic, or inoperable uterine sarcoma for IMT with ALK translocation), Erdheim-Chester disease (ECD) with ALK-fusion
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): 1) the disease is recurrent, advanced, or metastatic, AND 2) the disease is anaplastic lymphoma kinase (ALK)-positive.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ALVAIZ

Products Affected

- Alvaiz

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

ALYFTREK

Products Affected

- Alyftrek

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis: the requested drug will not be used in combination with other CFTR (cystic fibrosis transmembrane conductance regulator) potentiating agents (e.g., ivacaftor, deutivacaftor).
Age Restrictions	6 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AMBRISENTAN

Products Affected

- Ambrisentan

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

AMITRIPTYLINE

Products Affected

- Amitriptyline Hcl TABS 150MG, 25MG, 75MG
- Amitriptyline Hydrochloride TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neuropathic pain, chronic tension-type headache prophylaxis, chronic neck pain
Exclusion Criteria	N/A
Required Medical Information	For depression: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following alternative drugs: SSRIs (selective serotonin reuptake inhibitors), SNRIs (serotonin-norepinephrine reuptake inhibitors), including but not limited to bupropion, mirtazapine, or trazodone. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AMOXAPINE

Products Affected

- Amoxapine

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ANZUPGO

Products Affected

- Anzupgo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage will be denied when used in conjunction with other JAK inhibitors or potent immunosuppressants.
Required Medical Information	Member meets all of the following criteria: 1) Documented diagnosis of moderate to severe chronic hand eczema (CHE) AND 2) has experienced an inadequate treatment response to a topical corticosteroid OR topical corticosteroids are not advisable for the member.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by in or consultation with a dermatologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

AQNEURSA

Products Affected

- Aqneursa

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For Niemann-Pick disease type C (NPC), initial: 1) The diagnosis was confirmed by genetic testing demonstrating a variant of either the NPC1 or NPC2 gene, 2) The member has neurological manifestations of disease (e.g., loss of fine motor skills, swallowing, speech, ambulation), AND 3) The requested medication will not be used in combination with Miplyffa (arimoclomol). For Niemann-Pick disease type C, continuation: The member is experiencing benefit from therapy (e.g., stabilization or improvement in fine motor skills, swallowing, speech, ambulation).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AQVESME

Products Affected

- Aqvesme

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Diagnosis of liver cirrhosis
Required Medical Information	Member has a diagnosis of alpha- or beta-thalassemia associated anemia.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hematologist.
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Prevention of gout flares in members initiating or continuing urate-lowering therapy
Exclusion Criteria	N/A
Required Medical Information	For prevention of gout flares in members initiating or continuing urate-lowering therapy (e.g., allopurinol) (new starts): 1) two or more gout flares within the previous 12 months, AND 2) inadequate response, intolerance, or contraindication to maximum tolerated doses of a non-steroidal anti-inflammatory drug (NSAID) and colchicine, AND 3) concurrent use with urate-lowering therapy. For prevention of gout flares in members initiating or continuing urate-lowering therapy (e.g., allopurinol) (continuation): 1) member must have achieved or maintained a clinical benefit (i.e., a fewer number of gout attacks or fewer flare days) compared to baseline, AND 2) continued use of urate-lowering therapy concurrently with the requested drug. For recurrent pericarditis: member must have had an inadequate response, intolerance, or contraindication to maximum tolerated doses of a NSAID and colchicine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ARIKAYCE

Products Affected

- Arikayce

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ARMODAFINIL

Products Affected

- Armodafinil

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

ATTRUBY

Products Affected

- Attruby

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cardiomyopathy of variant or wild-type transthyretin-mediated amyloidosis (ATTR-CM), initial therapy: 1) member exhibits clinical manifestation of disease (e.g., dyspnea, fatigue, orthostatic hypotension, syncope, peripheral edema), AND 2) cardiac involvement was confirmed by one of the following: a) imaging via echocardiography or cardiac magnetic resonance imaging (e.g., end-diastolic interventricular septal wall thickness exceeding 12 millimeters), or b) myocardial technetium-labeled scintigraphy, AND 3) member meets one of the following: a) if the request is for variant ATTR-CM the member is positive for a mutation of the transthyretin (TTR) gene, b) if the request is for wild-type ATTR-CM the member has transthyretin precursor proteins confirmed by testing. For ATTR-CM, continuation: member demonstrates a beneficial response to therapy (e.g., slowing of clinical decline).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must have a diagnosis of ROS1-positive non-small cell lung cancer(NSCLC) OR solid tumors that 1) have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion AND 2) are locally advanced or metastatic or where surgical resection is likely to result in severe morbidity AND3) have progressed following treatment or have no satisfactory alternative therapy
Age Restrictions	Solid Tumors: 12 years of age or older ROS1+ NSCLC: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Tourette's syndrome
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AVONEX

Products Affected

- Avonex INJ 30MCG/0.5ML
- Avonex Pen

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

AYVAKIT

Products Affected

- Ayvakit

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Myeloid and lymphoid neoplasms with eosinophilia, gastrointestinal stromal tumor (GIST) for residual, unresectable, tumor rupture, or recurrent/metastatic disease without platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation.
Exclusion Criteria	N/A
Required Medical Information	For myeloid and lymphoid neoplasms with eosinophilia, the member meets all of the following criteria: 1) The disease is FIP1L1- PDGFRA rearrangement-positive, AND 2) The disease harbors a PDGFRA D842V mutation, AND 3) The disease is resistant to imatinib. For GIST, the member meets either of the following criteria: 1) The disease harbors PDGFRA exon 18 mutation, including a PDGFRA D842V mutation, OR 2) The requested drug will be used after failure on at least two FDA approved therapies in residual, unresectable, tumor rupture, or recurrent/metastatic disease without PDGFRA exon 18 mutation. For systemic mastocytosis: 1) The member has a diagnosis of indolent systemic mastocytosis or advanced systemic mastocytosis (including aggressive systemic mastocytosis [ASM], systemic mastocytosis with associated hematological neoplasm [SM-AHN], and mast cell leukemia [MCL]) AND 2) The member has a platelet count of greater than or equal to 50,000/microliter (mcL).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BAFIERTAM

Products Affected

- Bafiertam

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For urothelial carcinoma: 1) disease has susceptible fibroblast growth factor receptor 3 (FGFR3) genetic alterations, AND 2) the requested drug will be used as subsequent therapy for any of the following: a) locally advanced, recurrent, or metastatic urothelial carcinoma, OR b) stage II-IV, recurrent, or persistent urothelial carcinoma of the bladder.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BAXDELA

Products Affected

- Baxdela TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The infection is proven or strongly suspected to be caused by susceptible bacteria based on: 1) culture and susceptibility information, OR 2) local epidemiology and susceptibility patterns.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist
Coverage Duration	14 days
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BENLYSTA

Products Affected

- Benlysta INJ 200MG/ML

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

BENZTROPINE

Products Affected

- Benztropine Mesylate TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	EPS (extrapyramidal symptoms): The member has had an inadequate treatment response, contraindication, or intolerance to non-HRM (high-risk medication) alternative drug, amantadine. Parkinson's: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following non-HRM (high-risk medication) alternative drugs: including but not limited to amantadine, carbidopa/levodopa, pramipexole, or ropinirole. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member, AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BESREMI

Products Affected

- Besremi
- Besremi Pen

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BETASERON

Products Affected

- Betaseron

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BEXAROTENE ORAL

Products Affected

- Bexarotene CAPS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Mycosis fungoides (MF)/Sezary syndrome (SS), CD30-positive primary cutaneous anaplastic large cell lymphoma (ALCL), CD30-positive lymphomatoid papulosis (LyP), subcutaneous panniculitis-like T-cell lymphoma
Exclusion Criteria	N/A
Required Medical Information	For cutaneous T-cell lymphomas: members have tried at least one prior systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BEXAROTENE TOPICAL

Products Affected

- Bexarotene GEL

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Mycosis fungoides (MF)/Sezary syndrome (SS), chronic or smoldering adult T-cell leukemia/lymphoma (ATLL), primary cutaneous marginal zone lymphoma, primary cutaneous follicle center lymphoma
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BOSENTAN

Products Affected

- Bosentan

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

BOSULIF

Products Affected

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

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

BRAFTOVI

Products Affected

- Braftovi CAPS 75MG

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

BRINSUPRI

Products Affected

- Brinsupri

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Must have a diagnosis of non-cystic fibrosis bronchiectasis
Age Restrictions	12 years or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BRIVARACETAM

Products Affected

- Brivaracetam SOLN
- Brivaracetam TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of partial-onset seizures (i.e., focal-onset seizures): 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND 2) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to any of the following: Aptiom (if 4 years of age or older), Xcopri (if 18 years of age or older), Spritam (if 4 years of age or older).
Age Restrictions	1 month of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BRUKINSA

Products Affected

- Brukinsa

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For mantle cell lymphoma and chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL): the member has experienced an intolerable adverse event or has a contraindication to Calquence (acalabrutinib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

BYLVAY

Products Affected

- Bylvay
- Bylvay (pellets)

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of pruritis in progressive familial intrahepatic cholestasis (PFIC) (initial requests): 1) diagnosis of PFIC has been confirmed by genetic testing, 2) the member does not have PFIC type 2 with ABCB11 variants resulting in non-functional or complete absence of bile salt export pump protein (BSEP-3), AND 3) the member does not have any other concomitant liver disease. For treatment of pruritis in PFIC (continuation requests): the member has experienced benefit from therapy (for example, improvement in pruritis). For treatment of cholestatic pruritus with Alagille Syndrome (ALGS) (continuation): the member has experienced benefit from therapy (for example, improvement in pruritis).
Age Restrictions	For PFIC: 3 months of age or older, For ALGS: 12 months of age or older
Prescriber Restrictions	Prescribed by or in consultation with a hepatologist or gastroenterologist
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

BYSANTI

Products Affected

- Bysanti
- Bysanti Titration Pack A
- Bysanti Titration Pack B
- Bysanti Titration Pack C

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet all of the following requirements: 1) have a documented diagnosis of schizophrenia OR 2) the requested drug is being used for acute treatment of manic or mixed episodes associated with bipolar I disorder AND 3) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to at least two or more of the following Second Generation Antipsychotic (SGA) medications: aripiprazole, lurasidone, olanzapine, quetiapine, quetiapine ER, risperidone or ziprasidone.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist or psychiatrist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CABLIVI

Products Affected

- Cablivi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CABOMETYX

Products Affected

- Cabometyx

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-small cell lung cancer, Ewing sarcoma, osteosarcoma, gastrointestinal stromal tumor, endometrial carcinoma, soft tissue sarcoma (alveolar soft part sarcoma and extraskeletal myxoid chondrosarcoma subtypes)
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma: The disease is advanced, relapsed, or stage IV (including brain metastases). For non-small cell lung cancer: 1) the disease is rearranged during transfection (RET) positive AND 2) the disease is recurrent, advanced, or metastatic. For hepatocellular carcinoma: the requested drug will be used as subsequent therapy. For gastrointestinal stromal tumor (GIST): 1) the disease is residual, unresectable, recurrent, or metastatic/tumor rupture, AND 2) the disease has progressed after at least two FDA-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib). For Ewing sarcoma and osteosarcoma: the requested drug will be used as subsequent therapy. For differentiated thyroid cancer (DTC) (follicular, papillary, oncocytic): 1) the disease is locally advanced or metastatic, AND 2) the disease has progressed after a vascular endothelial growth factor receptor (VEGFR)- targeted therapy, AND 3) the member is refractory to radioactive iodine therapy (RAI) or ineligible for RAI. For endometrial carcinoma: 1) the disease is recurrent, AND 2) the requested drug will be used as subsequent therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A

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

CALQUENCE

Products Affected

- Calquence TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Waldenstrom macroglobulinemia (lymphoplasmacytic lymphoma), marginal zone lymphoma (including extranodal marginal zone lymphoma of the stomach, extranodal marginal zone lymphoma of nongastric sites, nodal marginal zone lymphoma, splenic marginal zone lymphoma)
Exclusion Criteria	N/A
Required Medical Information	For marginal zone lymphoma (including extranodal marginal zone lymphoma of the stomach, extranodal marginal zone lymphoma of nongastric sites, nodal marginal zone lymphoma, and splenic marginal zone lymphoma): the requested drug is being used for the treatment of relapsed, refractory, or progressive disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CAMZYOS

Products Affected

- Camzyos

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For obstructive hypertrophic cardiomyopathy: 1) before initiating therapy, member has left ventricular ejection fraction (LVEF) of 55 percent or greater, AND 2) member has New York Heart Association (NYHA) class II-III symptoms.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CAPLYTA

Products Affected

- Caplyta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Thyroid carcinomas (follicular, oncocytic, papillary).
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CARGLUMIC ACID

Products Affected

- Carglumic Acid

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

CAYSTON

Products Affected

- Cayston

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	FEV1 less than 25% or greater than 75% predicted OR Colonization with Burkholderia cepacia
Required Medical Information	For treatment of respiratory symptoms in cystic fibrosis members: 1) Pseudomonas aeruginosa is present in the member's airway cultures, OR 2) The member has a history of pseudomonas aeruginosa infection or colonization in the airways.
Age Restrictions	7 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CEQUR

Products Affected

- Cequr Simplicity 1u Extnd Wear
- Cequr Simplicity 2u
- Cequr Simplicity 2u Extnd Wear
- Cequr Simplicity Extended Wear Inserter
- Cequr Simplicity Inserter

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: 1) The member has diabetes requiring insulin management with multiple daily injections AND 2) The member is self-testing glucose levels 4 or more times per day OR the member is using a continuous glucose monitor AND 3) The member has experienced any of the following with the current diabetes regimen: inadequate glycemic control, recurrent hypoglycemia, wide fluctuations in blood glucose, dawn phenomenon with persistent severe early morning hyperglycemia, severe glycemic excursions.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CERDELGA

Products Affected

- Cerdelga

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For type 1 Gaucher disease (GD1): 1) Diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing, and 2) Member's CYP2D6 metabolizer status has been established using an FDA-cleared test, and 3) Member is a CYP2D6 extensive metabolizer, an intermediate metabolizer, or a poor metabolizer.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CEREZYME

Products Affected

- Cerezyme

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Type 2 Gaucher disease, Type 3 Gaucher disease.
Exclusion Criteria	N/A
Required Medical Information	For Gaucher disease: Diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CHOLBAM

Products Affected

- Cholbam

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For bile acid synthesis disorders due to single enzyme defects (SEDs) and adjunctive treatment of peroxisomal disorders (PDs): Diagnosis was confirmed by mass spectrometry or other biochemical or genetic testing. For bile acid synthesis disorders due to SEDs and adjunctive treatment of PDs, continuation of therapy: Member has achieved and maintained improvement in liver function.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CIBINQO

Products Affected

- Cibinqo

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

CINRYZE

Products Affected

- Cinryze

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the prophylaxis of angioedema attacks due to hereditary angioedema (HAE): 1) the member has HAE with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing, OR 2) the member has HAE with normal C1 inhibitor confirmed by laboratory testing and one of the following: a) the member tested positive for an F12, angiopoietin-1, plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosamine 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation, b) the member has a family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy for at least one month.
Age Restrictions	6 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an immunologist, allergist, or rheumatologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

CLOBAZAM

Products Affected

- Clobazam SUSP 2.5MG/ML
- Clobazam TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Seizures associated with Dravet syndrome
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Seizures associated with Lennox-Gastaut syndrome (LGS): 2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CLORAZEPATE

Products Affected

- Clorazepate Dipotassium TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the management of anxiety disorders: 1) The requested drug is being used concurrently with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the SSRI/SNRI becomes effective for the symptoms of anxiety OR 2) The member has experienced an inadequate treatment response, contraindication, or intolerance to to at least two agents from the following classes: a) selective serotonin reuptake inhibitors (SSRIs) b) serotonin-norepinephrine reuptake inhibitors (SNRIs). For all indications: 1) The prescriber must acknowledge the benefit of therapy with this prescribed medication outweighs the potential risks for the member AND 2) if the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COBENFY

Products Affected

- Cobenfy
- Cobenfy Starter Pack

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Urinary retention, moderate or severe hepatic impairment, gastric retention, untreated narrow-angle glaucoma
Required Medical Information	For treatment of schizophrenia: 1) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following generic products: aripiprazole, asenapine, lurasidone, olanzapine, quetiapine, risperidone, ziprasidone, AND 2) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following brand products: Caplyta, Lybalvi, Rexulti, Secuado, Vraylar.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COMETRIQ

Products Affected

- Cometriq

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-small cell lung cancer (NSCLC), thyroid carcinomas (follicular, oncocytic, papillary).
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): Disease is positive for rearranged during transfection (RET) rearrangements.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Hepatosplenic T-Cell lymphoma, breast implant-associated anaplastic large cell lymphoma (ALCL), peripheral T-Cell lymphoma
Exclusion Criteria	N/A
Required Medical Information	For chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL), breast implant-associated anaplastic large cell lymphoma (ALCL), and peripheral T-Cell lymphoma: the member has relapsed or refractory disease. For hepatosplenic T-Cell lymphoma: the member has refractory disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CORTROPHIN

Products Affected

- Cortrophin

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the following diagnoses, member has experienced an inadequate treatment response to a parenteral or an oral glucocorticoid (for ophthalmic diseases only, inadequate response to a trial of a topical ophthalmic glucocorticoid is also acceptable): 1) For rheumatic disorders (e.g., psoriatic arthritis, rheumatoid arthritis, ankylosing spondylitis, acute gouty arthritis): The requested drug must be used as adjunctive treatment, 2) For nephrotic syndrome: the requested drug must be requested for induction of diuresis or for remission of proteinuria, 3) For multiple sclerosis (MS): member has an acute exacerbation of MS, 4) Collagen diseases (e.g., systemic lupus erythematosus, dermatomyositis, or polymyositis), 5) Dermatologic diseases (e.g., severe erythema multiforme, Stevens-Johnson syndrome, severe psoriasis), 6) Ophthalmic diseases, acute or chronic (e.g., iritis, keratitis, optic neuritis), 7) Symptomatic sarcoidosis, 8) Allergic states (e.g., serum sickness, atopic dermatitis).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	MS exacerbation: 3 weeks, Allergic states: 1 month, All other diagnoses: 3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

COSENTYX

Products Affected

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

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

COTELLIC

Products Affected

- Cotellic

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Central nervous system (CNS) cancer (i.e., glioma, glioblastoma), adjuvant systemic therapy for cutaneous melanoma.
Exclusion Criteria	N/A
Required Medical Information	For central nervous system (CNS) cancer (i.e., glioma, glioblastoma): 1) The tumor is positive for BRAF V600E activating mutation, AND 2) The requested drug will be used in combination with vemurafenib. For melanoma: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), AND 2) The requested drug will be used in combination with vemurafenib, AND 3) The requested drug will be used for either of the following: a) unresectable, limited resectable, or metastatic disease, b) adjuvant or neoadjuvant systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CRENESSITY

Products Affected

- Crenessity

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For congenital adrenal hyperplasia (CAH), initial: 1) Member has a confirmed diagnosis of CAH by any of the following: a) genetic testing to confirm the presence of pathogenic variants in CYP21A1, b) confirmed 21-hydroxylase deficiency (e.g., cosyntropin [ACTH] stimulation test, baseline morning serum 17-hydroxyprogesterone [17-OHP] measurement by liquid chromatography-tandem mass spectrometry), AND 2) Member is currently on supraphysiological glucocorticoid therapy.
Age Restrictions	4 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CRINONE

Products Affected

- Crinone

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Prophylaxis for premature birth in women with a short cervix
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CTEXLI

Products Affected

- Ctexli

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must have a documented diagnosis of cerebrotendinous xanthomatosis (CTX).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CUVRIOR

Products Affected

- Cuvrior

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CYCLOBENZAPRINE

Products Affected

- Cyclobenzaprine Hydrochloride TABS 10MG, 5MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is using one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, hydroxyzine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Prior authorization applies to greater than cumulative 90 days of therapy per year.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CYPROHEPTADINE

Products Affected

- Cyproheptadine Hcl SYRP
- Cyproheptadine Hydrochloride TABS

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

CYSTADROPS

Products Affected

- Cystadrops

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystinosis: 1) Diagnosis was confirmed by ANY of the following: a) the presence of increased cystine concentration in leukocytes, OR b) genetic testing, OR c) demonstration of corneal cystine crystals by slit lamp examination, AND 2) the member has corneal cystine crystal accumulation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CYSTAGON

Products Affected

- Cystagon

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For nephropathic cystinosis: Diagnosis was confirmed by ANY of the following: 1) the presence of increased cystine concentration in leukocytes, OR 2) genetic testing, OR 3) demonstration of corneal cystine crystals by slit lamp examination.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

CYSTARAN

Products Affected

- Cystaran

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystinosis: 1) Diagnosis was confirmed by ANY of the following: a) the presence of increased cystine concentration in leukocytes, OR b) genetic testing, OR c) demonstration of corneal cystine crystals by slit lamp examination, AND 2) the member has corneal cystine crystal accumulation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DALFAMPRIDINE

Products Affected

- Dalfampridine Er

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For multiple sclerosis, member must meet the following (for new starts): prior to initiating therapy, member demonstrates sustained walking impairment. For multiple sclerosis (continuation): member must have experienced an improvement in walking speed OR other objective measure of walking ability since starting the requested drug.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan 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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), pigmented villonodular synovitis/tenosynovial giant cell tumor
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML), including members newly diagnosed with CML and members who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, AND 2) If member experienced resistance to an alternative tyrosine kinase inhibitor for CML, member is negative for T315I, Y253H, E255K/V, and F359V/C/I mutations. For acute lymphoblastic leukemia (ALL), including members who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, AND 2) If the member has experienced resistance to an alternative tyrosine kinase inhibitor for ALL, member is negative for T315I, Y253H, E255K/V, F359V/C/I and G250E mutations.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DASATINIB

Products Affected

- Dasatinib

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Gastrointestinal stromal tumor (GIST), metastatic and/or widespread chondrosarcoma, recurrent chordoma, T-cell acute lymphoblastic leukemia (ALL), and Philadelphia (Ph)-like B-ALL, myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in the chronic phase or blast phase, cutaneous melanoma
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML), including members who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia (Ph) chromosome or BCR-ABL gene AND 2) If member experienced resistance to an alternative tyrosine kinase inhibitor, member is negative for all of the following mutations: T315I/A, F317L/V/I/C, and V299L. For acute lymphoblastic leukemia (ALL), the member has a diagnosis of one of the following: 1) Philadelphia chromosome positive ALL, including members who have received a hematopoietic stem cell transplant: Diagnosis that has been confirmed by detection of the Ph chromosome or BCR-ABL gene AND if member experienced resistance to an alternative tyrosine kinase inhibitor, member is negative for all of the following mutations: T315I/A, F317L/V/I/C, and V299L OR 2) Ph-like B-ALL with ABL-class kinase fusion OR 3) Relapsed or refractory T-cell ALL with ABL-class translocation. For gastrointestinal stromal tumor (GIST): 1) Member meets all of the following: A) Disease is residual, unresectable, recurrent/progressive, or metastatic/tumor rupture, B) Member has received prior therapy with avapritinib AND C) Member is positive for platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutations. For cutaneous melanoma: 1) Disease is metastatic or unresectable, 2) Disease is positive for c-KIT activating mutations AND 3) Requested drug will be used as subsequent therapy AND 4) Member has had disease progression, intolerance, or risk of progression with BRAF-targeted therapy.
Age Restrictions	N/A

Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Post-induction therapy/consolidation following response to previous therapy with the same regimen for acute myeloid leukemia (AML)
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML): 1) the requested drug must be used in combination with cytarabine, 2) the member is 75 years of age or older OR has comorbidities that preclude intensive chemotherapy, AND 3) the requested drug will be used as treatment for induction therapy, post-induction/consolidation therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DAYBUE

Products Affected

- Daybue
- Daybue Stix

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DEFERASIROX

Products Affected

- Deferasirox

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For chronic iron overload due to blood transfusions: pretreatment serum ferritin level is greater than 1000 mcg/L. For chronic iron overload with non-transfusion-dependent thalassemia (NTDT) syndromes: 1) liver iron (Fe) concentration (LIC) of at least 5 mg Fe per gram of dry weight (Fe/g dw) AND 2) a serum ferritin greater than 300 mcg/L
Age Restrictions	Chronic iron overload due to blood transfusions: 2 years of age and older. Chronic iron overload with non-transfusion-dependent thalassemia (NTDT) syndromes: 10 years of age and older.
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DEFERIPRONE

Products Affected

- Deferiprone

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DEFLAZACORT

Products Affected

- Deflazacort
- Jaythari
- Kymbee TABS
- Pyquvi

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

DENOSUMAB

Products Affected

- Jubbonti
- Stoboclo

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

DESIPRAMINE

Products Affected

- Desipramine Hydrochloride

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neuropathic pain
Exclusion Criteria	N/A
Required Medical Information	For depression: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following alternative drugs: SSRIs (selective serotonin reuptake inhibitors), SNRIs (serotonin-norepinephrine reuptake inhibitors), including but not limited to, bupropion, mirtazapine, or trazodone. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DESMODA

Products Affected

- Desmoda

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Diagnosis of nephrogenic diabetes insipidus
Required Medical Information	Member has 1) Diagnosis of central diabetes insipidus (also known as arginine vasopressin deficiency) AND 2) meets one of the following: a) member has experienced an inadequate treatment response, intolerance, or has a contraindication to generic desmopressin tablet or spray OR b) member is unable to swallow a solid dosage form (e.g., age, physical impairment, dysphagia, patient is using a feeding tube or nasal gastric tube) OR c) dosage requirements cannot be met with a solid dosage form.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DIACOMIT

Products Affected

- Diacomit

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	To be taken in conjunction with clobazam.
Age Restrictions	6 months of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DIAZEPAM

Products Affected

- Diazepam CONC
- Diazepam SOLN 5MG/5ML
- Diazepam TABS
- Diazepam Intensol

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the management of anxiety disorders: 1) The requested drug is being used concurrently with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the SSRI/SNRI becomes effective for the symptoms of anxiety OR 2) The member has experienced an inadequate treatment response, contraindication, or intolerance to to at least two agents from the following classes: a) selective serotonin reuptake inhibitors (SSRIs), b) serotonin-norepinephrine reuptake inhibitors (SNRIs). For all indications: 1) The prescriber must acknowledge the benefit of therapy with this prescribed medication outweighs the potential risks for the member AND 2) if the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Applies to greater than cumulative 5 days of therapy per year.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

DICHLORPHENAMIDE

Products Affected

- Dichlorphenamide
- Ormalvi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For primary HYPOkalemic periodic paralysis: 1) The diagnosis was supported by genetic test results, OR 2) Member has a family history of primary hypokalemic periodic paralysis, OR 3) Member's attacks are associated with hypokalemia AND and thyrotoxic periodic paralysis has been ruled out. For primary HYPERkalemic periodic paralysis: 1) The diagnosis was supported by genetic test results, OR 2) Member has a family history of primary hyperkalemic periodic paralysis, OR 3) Member's attacks are associated with hyperkalemia. For continuation of therapy for primary HYPOkalemic and primary HYPERkalemic periodic paralysis: Member is demonstrating a response to therapy with the requested drug as demonstrated by a decrease in the number or severity of attacks.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 2 months. Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DICLOFENAC EPOLAMINE PATCH

Products Affected

- Diclofenac Epolamine

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Must have a documented diagnosis of acute pain due to one of the following: minor strain, sprain, or contusion.
Age Restrictions	6 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DICYCLOMINE

Products Affected

- Dicyclomine Hcl SOLN
- Dicyclomine Hydrochloride CAPS
- Dicyclomine Hydrochloride TABS 20MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DIMETHYL FUMARATE

Products Affected

- Dimethyl Fumarate CPDR
- Dimethyl Fumarate Starterpack CDPK 0

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DIPYRIDAMOLE

Products Affected

- Dipyridamole TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DOPTELET

Products Affected

- Doptelet
- Doptelet Sprinkle

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

DOXEPIN

Products Affected

- Doxepin Hcl CAPS 75MG
- Doxepin Hcl CONC
- Doxepin Hydrochloride CAPS 100MG, 10MG, 150MG, 25MG, 50MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For depression: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following alternative drugs: SSRIs (selective serotonin reuptake inhibitors), SNRIs (serotonin-norepinephrine reuptake inhibitors), including but not limited to, bupropion, mirtazapine, or trazodone. For anxiety: 1) The member has experienced an inadequate treatment response, contraindication, or intolerance to to at least two of the following agents: including but not limited to buspirone, duloxetine, escitalopram, sertraline, or venlafaxine extended-release. For all indications: 1) The prescriber must acknowledge the benefit of therapy with this prescribed medication outweighs the potential risks for the member AND 2) if the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.

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

DROXIDOPA

Products Affected

- Droxidopa

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For neurogenic orthostatic hypotension (nOH): For initial therapy, member has a persistent, consistent decrease in systolic blood pressure of at least 20 mmHg OR decrease in diastolic blood pressure of at least 10 mmHg within 3 minutes of standing or head-up tilt test. For continuation of therapy, member has experienced a sustained reduction in symptoms of nOH (i.e., decrease in dizziness, lightheadedness, or feeling faint). For both initial and continuation of therapy, the requested drug will be used for members with neurogenic orthostatic hypotension associated with one of the following diagnoses: 1) primary autonomic failure due to Parkinson's disease, multiple system atrophy, or pure autonomic failure, OR 2) dopamine beta-hydroxylase deficiency, OR 3) non-diabetic autonomic neuropathy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

DUPIXENT

Products Affected

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

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

EBGLYSS

Products Affected

- Ebglyss

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For atopic dermatitis, initial therapy: 1) Member has moderate-to-severe disease, AND 2) Member has experienced an inadequate treatment response to either a topical corticosteroid or a topical calcineurin inhibitor OR topical corticosteroids and topical calcineurin inhibitors are not advisable for the member. For atopic dermatitis, continuation of therapy: The member achieved or maintained positive clinical response.
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Initial: 4 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

EDARAVONE

Products Affected

- Edaravone
- Radicava Ors
- Radicava Ors Starter Kit

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

EGRIFTA

Products Affected

- Egrifta Sv
- Egrifta Wr

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Use for weight loss
Required Medical Information	For human immunodeficiency virus (HIV)-infected members with lipodystrophy: Member is receiving anti-retroviral therapy. For members who have received at least 6 months of the requested drug: Member has demonstrated clear clinical improvement from baseline that is supported by a waist circumference measurement or computed tomography (CT) scan.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist or endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ELIGARD

Products Affected

- Eligard

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Salivary gland tumors
Exclusion Criteria	N/A
Required Medical Information	For salivary gland tumors: 1) the disease is androgen receptor positive AND 2) the disease is recurrent, metastatic, or unresectable.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EMGALITY

Products Affected

- Emgality

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

ENBREL

Products Affected

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

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

ENSACOVE

Products Affected

- Ensacove

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ENSPRYNG

Products Affected

- Enspryng

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For neuromyelitis optica spectrum disorder (continuation): 1) there is no evidence of unacceptable toxicity or disease progression while on the current regimen, AND 2) the member has demonstrated a positive response to therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ENTYVIO

Products Affected

- Entyvio Pen

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

EOHILIA

Products Affected

- Eohilia

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For eosinophilic esophagitis (EoE): 1) Diagnosis has been confirmed by esophageal biopsy characterized by greater than or equal to 15 intraepithelial esophageal eosinophils per high power field, AND 2) The member is exhibiting clinical manifestations of the disease (for example, dysphagia).
Age Restrictions	11 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, allergist, or immunologist
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

EPIDIOLEX

Products Affected

- Epidiolex

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ERGOTAMINE

Products Affected

- Ergotamine Tartrate/caffeine

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Coverage will be denied when used in conjunction with potent CYP3A4 inhibitors (e.g., ritonavir, nelfinavir, indinavir, erythromycin, clarithromycin).
Required Medical Information	The member has experienced an inadequate treatment response, intolerance, or has a contraindication to at least ONE triptan 5-HT1 agonist.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ERIVEDGE

Products Affected

- Erivedge

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Adult medulloblastoma
Exclusion Criteria	N/A
Required Medical Information	For adult medulloblastoma: member has received prior systemic therapy AND has tumor(s) with mutations in the sonic hedgehog pathway.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ERLEADA

Products Affected

- Erleada

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ERLOTINIB

Products Affected

- Erlotinib Hydrochloride TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent non-small cell lung cancer (NSCLC), recurrent chordoma, relapsed or stage IV renal cell carcinoma (RCC), brain metastases from non-small cell lung cancer (NSCLC), recurrent pancreatic cancer
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC) (including brain metastases from NSCLC): 1) the disease is recurrent, advanced, or metastatic, AND 2) the member has sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease. For pancreatic cancer: the disease is locally advanced, unresectable, recurrent, or metastatic.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ESZOPICLONE

Products Affected

- Eszopiclone

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For insomnia: 1) The member has had an inadequate treatment response, contraindication, or intolerance to non-HRM (high-risk medication) alternative drug doxepin (3 mg or 6 mg) and ramelteon AND 2) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 3) If the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

EUCRISA

Products Affected

- Eucrisa

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

EVENITY

Products Affected

- Evenity

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

EVEROLIMUS

Products Affected

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

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Classic Hodgkin lymphoma, thymomas and thymic carcinomas, previously treated Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma, soft tissue sarcoma (perivascular epithelioid cell tumors (PEComa) and lymphangioleiomyomatosis subtypes), gastrointestinal stromal tumors, neuroendocrine tumors of the thymus, well differentiated grade 3 neuroendocrine tumors, thyroid carcinoma (papillary, oncocytic, and follicular), endometrial carcinoma, uterine sarcoma, breast cancer (in combination with fulvestrant or tamoxifen), histiocytic neoplasms (Rosai-Dorfman Disease, Erdheim-Chester Disease, Langerhans Cell Histiocytosis), meningiomas.
Exclusion Criteria	N/A
Required Medical Information	For breast cancer: 1) The disease is recurrent unresectable, advanced, or metastatic hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, AND 2) The requested drug is prescribed in combination with exemestane, fulvestrant, or tamoxifen, AND 3) The requested drug is used for subsequent treatment. For renal cell carcinoma: The disease is relapsed, advanced, or stage IV. For subependymal giant cell astrocytoma (SEGA): The requested drug is given as adjuvant treatment. For gastrointestinal stromal tumor: 1) The disease is residual, recurrent, unresectable, or metastatic/tumor rupture, AND 2) The disease has progressed after use of at least two FDA-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib). For Erdheim-Chester Disease (ECD), Rosai-Dorfman Disease, and Langerhans Cell Histiocytosis (LCH): the member must have a phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit alpha (PIK3CA) mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A

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

EVRYSDI

Products Affected

- Evrysdi

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

FABHALTA

Products Affected

- Fabhalta

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

FABRAZYME

Products Affected

- Fabrazyme

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For Fabry disease, the member meets ANY of the following: 1) diagnosis of Fabry disease was confirmed by an enzyme assay demonstrating a deficiency of alpha-galactosidase enzyme activity or by genetic testing, OR 2) the member is a symptomatic obligate carrier.
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

FENTANYL PATCH

Products Affected

- Fentanyl PT72 100MCG/HR, 12MCG/HR, 25MCG/HR, 50MCG/HR, 75MCG/HR

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the member meets all of the following: 1) The requested drug is being prescribed for pain severe and persistent enough to require an extended treatment period with a daily opioid analgesic in a member who has been taking an opioid AND 2) The member can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The member has been evaluated and the member will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a member who has been receiving an extended-release opioid agent for at least 30 days OR the member has taken an immediate-release opioid for at least one week.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

FILSPARI

Products Affected

- Filspari

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For members with primary immunoglobulin A nephropathy (IgAN) at risk of disease progression: 1) The member had an inadequate response to therapy with a maximally tolerated dose of a renin-angiotensin system (RAS) inhibitor (e.g., angiotensin-converting enzyme [ACE] inhibitor or angiotensin-receptor blocker [ARB]), OR 2) The member experienced an intolerance or has a contraindication to RAS inhibitors. For members with focal segmental glomerulosclerosis (FSGS) without nephrotic syndrome to reduce proteinuria: diagnosis confirmed by biopsy OR a genetic mutation known to cause FSGS.
Age Restrictions	FSGS: 8 years of age or older IgAN: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

FINGOLIMOD

Products Affected

- Fingolimod Hydrochloride

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	10 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FINTEPLA

Products Affected

- Fintepla

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FIRMAGON

Products Affected

- Firmagon INJ 120MG/VIAL, 80MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma: 1) The disease is advanced, relapsed, refractory or Stage IV, AND 2) The member has received two or more prior systemic therapies.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

FULPHILA

Products Affected

- Fulphila

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Stem cell transplantation-related indications
Exclusion Criteria	N/A
Required Medical Information	If receiving chemotherapy, the requested drug will be administered at least 24 hours after chemotherapy. For prophylaxis of myelosuppressive chemotherapy-induced febrile neutropenia: the member must meet both of the following: 1) Member has a solid tumor or non-myeloid cancer, AND 2) Member is currently receiving or will be receiving treatment with myelosuppressive anti-cancer therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GABAPENTIN

Products Affected

- Gabapentin Once-daily TABS 450MG, 750MG, 900MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For postherpetic neuralgia: The member has experienced an inadequate treatment response or intolerance to gabapentin immediate-release.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GAMASTAN

Products Affected

- Gamastan

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GATTEX

Products Affected

- Gattex

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For short bowel syndrome (SBS) initial therapy: 1) for an adult member, the member has been dependent on parenteral support OR 2) for a pediatric member, the member is dependent on parenteral support. For SBS continuation: requirement for parenteral support has decreased from baseline while on therapy with the requested drug.
Age Restrictions	1 year of age or older
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, gastrointestinal surgeon, or nutritional support specialist.
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent rearranged during transfection (RET) rearrangement-positive non-small cell lung cancer, RET gene fusion positive gallbladder cancer
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer, member must meet all of the following: 1) The disease is recurrent, advanced, or metastatic, AND 2) The tumor is rearranged during transfection (RET) fusion-positive or RET rearrangement-positive.
Age Restrictions	Non-small cell lung cancer: 18 years of age or older, Thyroid cancer: 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GEFITINIB

Products Affected

- Gefitinib

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Sensitizing epidermal growth factor receptor (EGFR) mutation-positive recurrent non-small cell lung cancer (NSCLC)
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): 1) the disease is recurrent, advanced, or metastatic, AND 2) the member must have a sensitizing epidermal growth factor receptor (EGFR) mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC), member meets either of the following: 1) has sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease AND a) has experienced an intolerable adverse event or contraindication to erlotinib, gefitinib or osimertinib, OR 2) has metastatic squamous NSCLC that progressed after platinum-based chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

GLATIRAMER

Products Affected

- Glatiramer Acetate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GLYCEROL PHENYLBUTYRATE

Products Affected

- Glycerol Phenylbutyrate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For urea cycle disorders (UCD) that cannot be managed by dietary protein restriction and/or amino acid supplementation alone: Diagnosis of UCD was confirmed by enzymatic, biochemical, or genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GOMEKLI

Products Affected

- Gomekli

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

GROWTH HORMONE

Products Affected

- Genotropin
- Genotropin Miniquick
- Humatrope INJ 12MG, 24MG, 6MG
- Norditropin Flexpro INJ 10MG/1.5ML, 15MG/1.5ML, 5MG/1.5ML
- Nutropin Aq Nuspin 10
- Nutropin Aq Nuspin 20
- Nutropin Aq Nuspin 5
- Omnitrope
- Zomacton

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

GUANFACINE

Products Affected

- Guanfacine Hydrochloride
- Guanfacine Hydrochloride Er

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HARLIKU

Products Affected

- Harliku

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HERCEPTIN

Products Affected

- Herceptin INJ 150MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-positive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	All indications: the member had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information. For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with paclitaxel and continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

HERCEPTIN HYLECTA

Products Affected

- Herceptin Hylecta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-positive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	All indications: the member had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information. For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with paclitaxel and continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

HERNEXEOS

Products Affected

- Hernexeos

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HERZUMA

Products Affected

- Herzuma

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-positive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	All indications: the member had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information. For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with paclitaxel and continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

HETLIOZ LQ

Products Affected

- Hetlioz Lq

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For nighttime sleep disturbances in Smith-Magenis Syndrome (SMS): 1) For initial therapy and continuation therapy, the member has a confirmed diagnosis of SMS, AND 2) If currently on therapy with the requested drug, the member experienced improvement in the quality of sleep since starting therapy.
Age Restrictions	3 to 15 years of age
Prescriber Restrictions	Prescribed by or in consultation with a sleep disorder specialist, neurologist, or psychiatrist
Coverage Duration	Initiation: 6 months, Renewal: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

HYDROXYZINE

Products Affected

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

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

HYRNUO

Products Affected

- Hyrnuo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IBRANCE

Products Affected

- Ibrance

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

IBTROZI

Products Affected

- Ibtrozi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ICATIBANT

Products Affected

- Icatibant Acetate
- Sajazir

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of acute angioedema attacks due to hereditary angioedema (HAE): 1) the member has HAE with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR 2) the member has HAE with normal C1 inhibitor confirmed by laboratory testing and one of the following: a) the member tested positive for an F12, angiopoietin-1, plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosamine 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation, b) the member has a family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy for at least one month.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an immunologist, allergist, or rheumatologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ICLUSIG

Products Affected

- Iclusig

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Myeloid and/or lymphoid neoplasms with eosinophilia and FGFR1 or ABL1 rearrangement in the chronic phase or blast phase, Gastrointestinal Stromal Tumors
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML), including members who have received a hematopoietic stem cell transplant: 1) Member has accelerated or blast phase CML and no other kinase inhibitor is indicated, OR 2) Member has chronic phase CML and has experienced resistance or intolerance to at least 2 prior kinase inhibitors AND at least one of those was imatinib, dasatinib, or nilotinib, OR 3) Member is positive for the T315I mutation. For acute lymphoblastic leukemia (ALL), including members who have received a hematopoietic stem cell transplant: Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For gastrointestinal stromal tumors (GIST): 1) Disease meets any of the following: A) residual, B) unresectable, C) recurrent, D) metastatic/tumor rupture, AND 2) Disease has progressed after use of at least two FDA approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Newly-diagnosed acute myeloid leukemia
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML) with an isocitrate dehydrogenase-2 (IDH2) mutation: 1) member has newly-diagnosed AML and is not a candidate for intensive induction therapy, OR 2) the requested drug will be used as post-induction therapy following response to induction therapy with the requested drug, OR 3) member has relapsed or refractory AML, OR 4) the requested drug will be used as consolidation therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMATINIB

Products Affected

- Imatinib Mesylate TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Desmoid tumors, pigmented villonodular synovitis/tenosynovial giant cell tumor (PVNS/TGCT), recurrent chordoma, cutaneous melanoma, Kaposi sarcoma, chronic graft versus host disease (cGVHD), T-cell acute lymphoblastic leukemia with ABL-class translocation, aggressive systemic mastocytosis for well-differentiated systemic mastocytosis (WDSM) or when eosinophilia is present with FIP1L1-PDGFRA fusion gene, myeloid and/or lymphoid neoplasms with eosinophilia and ABL1, FIP1L1-PDGFRA, or PDGFRB rearrangement in the chronic phase or blast phase.
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), including members who have received a hematopoietic stem cell transplant: Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML: Member did not fail (excluding failure due to intolerance) prior therapy with a tyrosine kinase inhibitor. For cutaneous melanoma: 1) Disease is metastatic or unresectable AND 2) Disease is positive for c-KIT activating mutations AND 3) Requested medication will be used as subsequent therapy AND 4) Member has had disease progression, intolerance, or risk of progression with BRAF-targeted therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A

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

IMBRUVICA

Products Affected

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

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Hairy cell leukemia, lymphoplasmacytic lymphoma, primary central nervous system (CNS) lymphoma, human immunodeficiency virus (HIV)-related B-cell lymphoma, diffuse large B-cell lymphoma, post-transplant lymphoproliferative disorders, high-grade B-cell lymphoma, mantle cell lymphoma, marginal zone lymphoma (including extranodal marginal zone lymphoma of the stomach, extranodal marginal zone lymphoma of nongastric sites, nodal marginal zone lymphoma, splenic marginal zone lymphoma), brain metastases in lymphoma
Exclusion Criteria	N/A
Required Medical Information	For mantle cell lymphoma: 1) the requested drug will be used as subsequent therapy OR 2) the requested drug will be used in combination with rituximab as pretreatment to induction therapy with RHyperCVAD (rituximab, cyclophosphamide, vincristine, doxorubicin, and dexamethasone) regimen, OR 3) the requested drug will be used as aggressive induction therapy. For marginal zone lymphoma (including extranodal marginal zone lymphoma of the stomach, extranodal marginal zone lymphoma of nongastric sites, nodal marginal zone lymphoma, and splenic marginal zone lymphoma): the requested drug will be used as second-line or subsequent therapy. For hairy cell leukemia: the requested drug will be used as a single agent for disease progression. For primary CNS lymphoma: 1) the disease is relapsed or refractory OR 2) the requested drug is used for induction therapy as a single agent. For diffuse large B-cell lymphoma, high-grade B-cell lymphoma, human immunodeficiency virus (HIV)-related B-cell lymphoma: The requested drug will be used as a single agent and as second-line or subsequent therapy for relapsed or refractory disease. For post-transplant lymphoproliferative disorders: the requested drug will be used in members who have received prior chemoimmunotherapy.
Age Restrictions	N/A

Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMIPRAMINE

Products Affected

- Imipramine Hcl TABS 25MG, 50MG
- Imipramine Hydrochloride TABS 10MG
- Imipramine Pamoate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neuropathic pain
Exclusion Criteria	N/A
Required Medical Information	For depression: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following alternative drugs: SSRIs (selective serotonin reuptake inhibitors), SNRIs (serotonin-norepinephrine reuptake inhibitors), including but not limited to bupropion, mirtazapine, or trazodone. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMKELDI

Products Affected

- Imkeldi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent chordoma, cutaneous melanoma, Kaposi sarcoma
Exclusion Criteria	N/A
Required Medical Information	For all indications: The member is unable to use imatinib tablets. For chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), including members who have received a hematopoietic stem cell transplant: Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML: Member did not fail (excluding failure due to intolerance) prior therapy with a tyrosine kinase inhibitor. For cutaneous melanoma: 1) Disease is metastatic or unresectable AND 2) Disease is positive for c-KIT activating mutations AND 3) Requested medication will be used as subsequent therapy AND 4) Member has had disease progression, intolerance, or risk of progression with BRAF-targeted therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

IMVEXXY

Products Affected

- Imvexxy Maintenance Pack
- Imvexxy Starter Pack

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INBRIJA

Products Affected

- Inbrija

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INCRELEX

Products Affected

- Increlex

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Pediatric members with closed epiphyses
Required Medical Information	For growth failure due to severe primary insulin-like growth factor-1 (IGF-1) deficiency or growth hormone (GH) gene deletion in member who have developed neutralizing antibodies to GH, member meets all of the following prior to beginning therapy with the requested drug (new starts only): 1) height 3 or more standard deviations (SD) below the mean for children of the same age and gender AND 2) basal IGF-1 level 3 or more SD below the mean for children of the same age and gender AND 3) provocative growth hormone test showing a normal or elevated growth hormone level. For growth failure due to severe primary IGF-1 deficiency or GH gene deletion in members who have developed neutralizing antibodies to GH, continuation of therapy: member is experiencing improvement.
Age Restrictions	2 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INFLIXIMAB

Products Affected

- Infliximab
- Remicade

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Behcet's syndrome, hidradenitis suppurativa, sarcoidosis, Takayasu's arteritis, uveitis.
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): 1) Member meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) or leflunomide OR b) intolerance or contraindication to MTX AND leflunomide, AND 2) Member meets ANY of the following: a) inadequate treatment response, intolerance or contraindication to MTX OR b) inadequate treatment response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis (new starts only): an inadequate treatment response or intolerance to a non-steroidal anti-inflammatory drug (NSAID) OR contraindication that would prohibit a trial of NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at time of diagnosis, AND 2) member meets ANY of the following: a) member has experienced inadequate treatment response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with MTX, cyclosporine, or acitretin, OR b) pharmacologic treatment with MTX, cyclosporine, or acitretin is contraindicated, OR c) member has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of BSA or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	For hidradenitis suppurativa (new starts only): Member has severe, refractory disease. For uveitis (new starts only): Inadequate treatment response or intolerance or has a contraindication to a trial of immunosuppressive therapy for uveitis. For all indications: The member experienced an intolerable adverse event to Renflexis and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

INLURIYO

Products Affected

- Inluriyo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INLYTA

Products Affected

- Inlyta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Thyroid carcinoma (papillary, oncocytic, or follicular), alveolar soft part sarcoma
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma: the disease is advanced, relapsed, or Stage IV.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INQOVI

Products Affected

- Inqovi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and janus kinase 2 (JAK2) rearrangement, accelerated or blast phase myeloproliferative neoplasms
Exclusion Criteria	N/A
Required Medical Information	For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement: the disease is in chronic or blast phase.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

INSULIN SUPPLIES

Products Affected

- Alcohol Prep Pads PADS 70%
- Curity Gauze Pads 2"x2" 12 Ply
- Insulin Syringes, Needles

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested product is being used with insulin.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ISOTRETINOIN

Products Affected

- Accutane
- Amnesteem
- Claravis
- Isotretinoin CAPS
- Zenatane

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Severe acne vulgaris, severe refractory rosacea, neuroblastoma, cutaneous T-cell lymphoma (CTCL) (e.g., mycosis fungoides, Sezary syndrome), high risk for developing skin cancer (squamous cell cancers), transient acantholytic dermatosis (Grover's Disease), keratosis follicularis (Darier Disease), lamellar ichthyosis, pityriasis rubra pilaris
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ISTURISA

Products Affected

- Isturisa TABS 1MG, 5MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IVERMECTIN TAB

Products Affected

- Ivermectin TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Ascariasis, Cutaneous larva migrans, Mansonelliasis, Scabies, Gnathostomiasis, Pediculosis
Exclusion Criteria	N/A
Required Medical Information	The requested drug is not being prescribed for the prevention or treatment of coronavirus disease 2019 (COVID-19).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 month
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

IWILFIN

Products Affected

- Iwilfin

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

JAKAFI

Products Affected

- Jakafi
- Jakafi Xr

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Lower-risk myelofibrosis, accelerated or blast phase myeloproliferative neoplasms, acute lymphoblastic leukemia (ALL), chronic myelomonocytic leukemia (CMML)-2, myelodysplastic syndrome/myeloproliferative neoplasm (MDS/MPN) with neutrophilia, essential thrombocythemia, myeloid, lymphoid or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement, T-cell prolymphocytic leukemia, T-cell large granular lymphocytic leukemia
Exclusion Criteria	N/A
Required Medical Information	For polycythemia vera: 1) member had an inadequate response or intolerance to hydroxyurea and Besremi (ropeginterferon alfa-2b-njft), OR 2) member has high risk disease. For acute lymphoblastic leukemia: member has a cytokine receptor-like factor 2 (CRLF2) mutation or a mutation associated with activation of the Janus kinase/signal transducers and activators of transcription (JAK/STAT) pathway. For CMML-2: the requested drug is used in combination with a hypomethylating agent. For myelodysplastic syndrome/myeloproliferative neoplasm (MDS/MPN) with neutrophilia: the requested drug is used as a single agent or in combination with a hypomethylating agent. For essential thrombocythemia: member had an inadequate response or loss of response to hydroxyurea, interferon therapy, or anagrelide. For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement: the disease is in chronic or blast phase.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A

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

JASCAYD

Products Affected

- Jascayd

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

JAYPIRCA

Products Affected

- Jaypirca

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

JIDEYTRO

Products Affected

- Jideytro

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

JOENJA

Products Affected

- Joenja TABS 70MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For activated phosphoinositide 3-kinase delta syndrome (APDS): the diagnosis was confirmed by genetic testing demonstrating variant in either PIK3CD or PIK3R1.
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

JOURNAVX

Products Affected

- Journavx

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must use for the treatment of new onset moderate to severe acute pain, including postoperative pain.
Age Restrictions	18 years of age or older
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.

JUXTAPID

Products Affected

- Juxtapid CAPS 10MG, 20MG, 2MG, 30MG, 5MG

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

JYLAMVO

Products Affected

- Jylamvo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KALYDECO

Products Affected

- Kalydeco

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis (CF): The requested medication will not be used in combination with other medications containing ivacaftor.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KANJINTI

Products Affected

- Kanjinti

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-positive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	All indications: the member had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information. For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with paclitaxel and continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KERENDIA

Products Affected

- Kerendia

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KEVZARA

Products Affected

- Kevzara

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): Member has experienced an inadequate treatment response, intolerance, or has a contraindication to two of the following products: Yuflyma (adalimumab-aaty), Enbrel (etanercept), Hulio (adalimumab-fkjp), Rinvoq (upadacitinib), Xeljanz (tofacitinib)/Xeljanz XR (tofacitinib extended-release). For polymyalgia rheumatica (PMR) (new starts only): 1) Member has experienced an inadequate treatment response to corticosteroids OR 2) Member has experienced a disease flare while attempting to taper corticosteroids OR 3) Member has a contraindication that would prohibit a trial of corticosteroids. For active polyarticular juvenile idiopathic arthritis (new starts only): Member has experienced an inadequate treatment response, intolerance, or has a contraindication to two of the following products: Yuflyma (adalimumab-aaty), Enbrel (etanercept), Hulio (adalimumab-fkjp), Rinvoq (upadacitinib)/Rinvoq LQ (upadacitinib), Xeljanz (tofacitinib)/Xeljanz XR (tofacitinib extended-release).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

KINERET

Products Affected

- Kineret

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

KISQALI

Products Affected

- Kisqali

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent hormone receptor-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer, in combination with an aromatase inhibitor, or fulvestrant. Endometrial cancer, in combination with letrozole, for estrogen receptor positive tumors.
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KLISYRI

Products Affected

- Klisyri

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member has experienced an inadequate treatment response, intolerance, or has a contraindication to ONE of the following: A) imiquimod 5 percent cream, B) fluorouracil cream or solution.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For relapsed or refractory acute myeloid leukemia (AML): 1) Member must have a susceptible nucleophosmin 1 (NPM1) mutation AND 2) Members must have no satisfactory alternative treatment options.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

KOSELUGO

Products Affected

- Koselugo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	BRAF fusion or BRAF V600E activating mutation-positive recurrent or progressive circumscribed glioma, Langerhans cell histiocytosis.
Exclusion Criteria	N/A
Required Medical Information	For neurofibromatosis type 1 (NF1) members have symptomatic, inoperable plexiform neurofibromas (PN).
Age Restrictions	For neurofibromatosis type 1: 1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent KRAS G12C-positive non-small cell lung cancer (NSCLC), Central nervous system (CNS) brain metastases from KRAS G12C-positive NSCLC, KRAS G12C-positive pancreatic adenocarcinoma, KRAS G12C-positive ampullary adenocarcinoma, KRAS G12C-positive appendiceal adenocarcinoma, KRAS G12C-positive Biliary Tract Cancer (intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma, gall bladder cancer)
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LANREOTIDE

Products Affected

- Lanreotide Acetate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Tumor control of neuroendocrine tumors (NETs) (including tumors of the lung, thymus, well-differentiated grade 3 NETs not of gastroenteropancreatic origin with favorable biology, and pheochromocytoma/paraganglioma)
Exclusion Criteria	N/A
Required Medical Information	For acromegaly, initial: 1) Member has a high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range, AND 2) Member had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the member has not had surgery or radiotherapy. For acromegaly, continuation of therapy: Member's IGF-1 level has decreased or normalized since initiation of therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LAPATINIB

Products Affected

- Lapatinib Ditosylate

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

LAZCLUZE

Products Affected

- Lazcluze

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LEDIPASVIR/SOFOSBUVIR

Products Affected

- Ledipasvir/sofosbuvir

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases and Infectious Diseases Society of America (AASLD-IDSA) treatment guidelines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Criteria applied consistent w/ current AASLD-IDSA guidance. Reminder for 8wk option if appropriate.
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LENALIDOMIDE

Products Affected

- Lenalidomide

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Systemic light chain amyloidosis, classical Hodgkin lymphoma, myelodysplastic syndrome without the 5q deletion cytogenetic abnormality, POEMS (polyneuropathy, organomegaly, endocrinopathy, monoclonal protein, skin changes) syndrome, myeloproliferative neoplasms, Kaposi Sarcoma, Langerhans cell histiocytosis, Rosai-Dorfman disease, peripheral T-Cell lymphomas not otherwise specified, angioimmunoblastic T-cell lymphoma (AITL), enteropathy-associated T-cell lymphoma, monomorphic epitheliotropic intestinal T-cell lymphoma, nodal peripheral T-cell lymphoma, adult T-cell leukemia/lymphoma, hepatosplenic T-cell lymphoma, primary central nervous system (CNS) lymphoma, chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL), human immunodeficiency virus (HIV)-related B-cell lymphomas, monomorphic post-transplant lymphoproliferative disorder, diffuse large B-cell lymphoma, multicentric Castlemans disease, high-grade B-cell lymphomas, histologic transformation of indolent lymphoma to diffuse large B-cell lymphoma
Exclusion Criteria	N/A
Required Medical Information	For myelodysplastic syndrome (MDS): member has lower risk MDS with symptomatic anemia per the Revised International Prognostic Scoring System (IPSS-R), International Prognostic Scoring System (IPSS), or World Health organization (WHO) classification-based Prognostic Scoring System (WPSS).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A

Prerequisite Therapy Required	Criteria DOES NOT 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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Medullary thyroid carcinoma, recurrent endometrial carcinoma, thymic carcinoma, unresectable or metastatic cutaneous melanoma.
Exclusion Criteria	N/A
Required Medical Information	For differentiated thyroid cancer (follicular, papillary, or oncocytic): disease is not amenable to radioactive iodine therapy and unresectable, locally recurrent, persistent, or metastatic. For hepatocellular carcinoma (HCC): disease is unresectable, extrahepatic/metastatic, or liver-confined. For renal cell carcinoma (RCC): the disease is advanced, relapsed, or stage IV. For endometrial carcinoma (EC), the member meets ALL of the following: 1) The disease is advanced, recurrent, or metastatic, 2) The requested drug will be used in combination with pembrolizumab, 3) The member experienced disease progression following prior systemic therapy. For anaplastic thyroid carcinoma, the member meets ALL of the following: 1) The disease is metastatic, 2) The requested drug will be used in combination with pembrolizumab.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LEQEMBI

Products Affected

- Leqembi Iqlik

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

LEUKERAN

Products Affected

- Leukeran

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

LEUPROLIDE

Products Affected

- Leuprolide Acetate INJ 1MG/0.2ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Use in combination with growth hormone for children with growth failure and advancing puberty, recurrent androgen receptor positive salivary gland tumors, central precocious puberty
Exclusion Criteria	N/A
Required Medical Information	For central precocious puberty (CPP): Members not currently receiving therapy must meet all of the following criteria: 1) Diagnosis of CPP was confirmed by a pubertal response to a gonadotropin releasing hormone (GnRH) agonist test OR a pubertal level of a third generation luteinizing hormone (LH) assay, 2) Assessment of bone age versus chronological age supports the diagnosis of CPP, 3) The onset of secondary sexual characteristics occurred prior to 8 years of age for female members OR prior to 9 years of age for male members.
Age Restrictions	CPP: Member must be less than 12 years old if female and less than 13 years old if male
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIDOCAINE

Products Affected

- Lidocaine OINT 5%
- Lidocaine PTCH 5%
- Lidocaine Hydrochloride SOLN
- Lidocan

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Pain associated with diabetic neuropathy, pain associated with cancer-related neuropathy (including treatment-related neuropathy [e.g., neuropathy associated with radiation treatment or chemotherapy]).
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIDOCAINE/PRILOCAINE

Products Affected

- Lidocaine/prilocaine CREA

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Pain associated with diabetic neuropathy, pain associated with cancer-related neuropathy (including treatment-related neuropathy [e.g., neuropathy associated with radiation treatment or chemotherapy]).
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIFYORLI

Products Affected

- Lifyorli

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must meet all of the following requirements: 1) Have a documented diagnosis of platinum-resistant epithelial ovarian, fallopian tube, or primary peritoneal cancer, AND 2) Has received at least one prior systemic therapy that includes bevacizumab, AND 3) the requested drug is used combination with nab-paclitaxel (Abraxane).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LIRAGLUTIDE

Products Affected

- Liraglutide INJ 6MG/ML

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

LIVDELZI

Products Affected

- Livdelzi

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

LIVMARLI

Products Affected

- Livmarli

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of cholestatic pruritis in a member with Alagille syndrome (ALGS) (continuation): the member has experienced benefit from therapy (for example, improvement in pruritis). For treatment of cholestatic pruritis in a member with Progressive Familial Intrahepatic Cholestasis (PFIC), (initial): 1) diagnosis of PFIC has been confirmed by genetic testing, 2) the member does not have PFIC type 2 with ABCB11 variants resulting in non-functional or complete absence of bile salt export pump (BSEP) protein, AND 3) the member does not have any other concomitant liver disease. For treatment of cholestatic pruritis in a member with PFIC (continuation): the member has experienced benefit from therapy (for example, improvement in pruritis).
Age Restrictions	For ALGS: 3 months of age or older, For PFIC: 12 months of age or older
Prescriber Restrictions	Prescribed by or in consultation with a hepatologist or gastroenterologist.
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LIVTENCITY

Products Affected

- Livtencity

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Weight less than 35kg
Required Medical Information	N/A
Age Restrictions	12 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist, transplant specialist, hematologist, or oncologist.
Coverage Duration	3 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LONSURF

Products Affected

- Lonsurf

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Unresectable locally advanced, recurrent, or metastatic esophageal cancer. Unresectable locally advanced or recurrent gastric cancer and gastroesophageal junction cancers. Advanced or metastatic appendiceal adenocarcinoma.
Exclusion Criteria	N/A
Required Medical Information	For colorectal cancer (including appendiceal adenocarcinoma): The disease is advanced or metastatic. For gastric, esophageal, or gastroesophageal junction adenocarcinoma, ALL of the following criteria must be met: 1) The disease is unresectable locally advanced, recurrent, or metastatic, and 2) The member has been previously treated with at least two prior lines of chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Anaplastic lymphoma kinase (ALK)-positive recurrent non-small cell lung cancer (NSCLC), proto-oncogene tyrosine-protein kinase ROS1 (ROS1) rearrangement-positive recurrent, advanced, or metastatic NSCLC, symptomatic or relapsed/refractory ALK-positive Erdheim-Chester Disease, inflammatory myofibroblastic tumor (IMT) with ALK translocation (including advanced, recurrent/metastatic, or inoperable uterine sarcoma for IMT with ALK translocation), central nervous system (CNS) brain metastases from ALK rearrangement-positive NSCLC, relapsed or refractory ALK-positive Diffuse Large B-Cell Lymphoma, relapsed or refractory ALK-positive Peripheral T-Cell Lymphoma
Exclusion Criteria	N/A
Required Medical Information	For recurrent, advanced, or metastatic non-small cell lung cancer: 1) Disease is ALK-positive, OR 2) Disease is positive for ROS1 rearrangement and the requested drug is being used following disease progression on one of the following: crizotinib, entrectinib, ceritinib, or repotrectinib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LUMAKRAS

Products Affected

- Lumakras

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent KRAS G12C-positive non-small cell lung cancer (NSCLC), recurrent, locally advanced, or metastatic KRAS G12C-positive pancreatic adenocarcinoma, advanced or unresectable KRAS G12C-positive colorectal cancer (including appendiceal adenocarcinoma), progressive KRAS G12C-positive ampullary adenocarcinoma
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): 1) the disease is recurrent, advanced, or metastatic, AND 2) the member has KRAS G12C mutation-positive disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LUMIZYME

Products Affected

- Lumizyme

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For Pompe disease: Diagnosis was confirmed by an enzyme assay demonstrating a deficiency of acid alpha-glucosidase (GAA) enzyme activity or by genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LUPKYNIS

Products Affected

- Lupkynis

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Use in combination with cyclophosphamide
Required Medical Information	For lupus nephritis: 1) member is currently receiving background immunosuppressive therapy regimen for lupus nephritis (for example, mycophenolate mofetil, corticosteroids) OR 2) member has an intolerance or has a contraindication to background immunosuppressive therapy regimen for lupus nephritis. For lupus nephritis continuation: member is receiving benefit from therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

LUPRON

Products Affected

- Lupron Depot (1-month)
- Lupron Depot (3-month)
- Lupron Depot (4-month)
- Lupron Depot (6-month)

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LYBALVI

Products Affected

- Lybalvi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

LYNKUET

Products Affected

- Lynkuet

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent HER2-negative, BRCA 1/2-germline mutated breast cancer, recurrent or metastatic HER2-positive, BRCA 1/2-germline mutated breast cancer, uterine leiomyosarcoma.
Exclusion Criteria	N/A
Required Medical Information	For recurrent or metastatic breast cancer: the disease is BRCA 1/2-germline mutated. For prostate cancer: 1) The member has a BRCA mutation and the requested drug will be used in combination with abiraterone and an oral corticosteroid OR 2) The member has progressed on prior treatment with an androgen receptor-directed therapy. For ovarian, fallopian tube, or primary peritoneal cancer: The requested drug is used for maintenance therapy for stage II-IV or recurrent disease who are in complete or partial response to chemotherapy. For uterine leiomyosarcoma: 1) the member has had at least one prior therapy AND 2) the member has BRCA-altered disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh [CTP] class B or C).
Required Medical Information	For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [CTP class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases and Infectious Diseases Society of America (AASLD-IDSA) treatment guidelines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
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.

MECLIZINE

Products Affected

- Meclizine Hcl TABS
- Meclizine Hydrochloride TABS 25MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member, AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Prior Authorization applies to greater than cumulative 30 days of therapy per year.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MEKINIST

Products Affected

- Mekinist

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Langerhans cell histiocytosis, Erdheim-Chester disease, Rosai-Dorfman disease, hairy cell leukemia.
Exclusion Criteria	N/A
Required Medical Information	For melanoma: 1) The tumor is positive for a BRAF mutation, AND 2) The requested drug will be used as a single agent or in combination with dabrafenib, AND 3) The requested drug will be used for either of the following: a) unresectable, limited resectable, or metastatic disease, b) adjuvant or neoadjuvant systemic therapy. For uveal melanoma: The requested drug will be used as a single agent. For ovarian cancer, fallopian tube cancer, and primary peritoneal cancer: The requested drug will be used to treat persistent or recurrent disease. For papillary, follicular, and oncocytic thyroid carcinoma: 1) The disease is positive for BRAF V600E mutation, AND 2) The disease is not amenable to radioactive iodine (RAI) therapy, AND 3) The requested drug will be used in combination with dabrafenib. For hairy cell leukemia: 1) the requested drug will be used in combination with dabrafenib, AND 2) the member has not had previous treatment with BRAF inhibitor therapy. For solid tumors: 1) The tumor is positive for a BRAF V600E mutation, AND 2) The requested drug will be used in combination with dabrafenib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Adjuvant systemic therapy for cutaneous melanoma, Langerhans Cell Histiocytosis, recurrent non-small cell lung cancer (NSCLC)
Exclusion Criteria	N/A
Required Medical Information	For melanoma: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), AND 2) The requested drug will be used in combination with encorafenib, AND 3) The requested drug will be used for either of the following: a) unresectable, limited resectable, or metastatic disease, b) adjuvant systemic therapy. For non-small cell lung cancer: 1) The tumor is positive for BRAF V600E mutation, AND 2) The requested drug will be used in combination with encorafenib, AND 3) The disease is advanced, recurrent, or metastatic.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

METHADONE - IR BEFORE ER

Products Affected

- Methadone Hcl SOLN 5MG/5ML
- Methadone Hcl TABS
- Methadone Hydrochloride SOLN 10MG/5ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the member meets all of the following: 1) The requested drug is being prescribed for pain severe and persistent enough to require an extended treatment period with a daily opioid analgesic in a member who has been taking an opioid AND 2) The member can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The member has been evaluated and will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a member who has been receiving an extended-release opioid agent for at least 30 days OR the member has taken an immediate-release opioid for at least one week.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

METHOCARBAMOL

Products Affected

- Methocarbamol TABS 500MG, 750MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is using one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, hydroxyzine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Prior authorization applies to greater than cumulative 90 days of therapy per year.
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MIGLUSTAT

Products Affected

- Miglustat
- Yargesa

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For type 1 Gaucher disease (GD1): The diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MIPLYFFA

Products Affected

- Miplyffa

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

MODAFINIL

Products Affected

- Modafinil TABS

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

MODEYSO

Products Affected

- Modeyso

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MORPHINE - IR BEFORE ER

Products Affected

- Morphine Sulfate Er TBCR

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the member meets all of the following: 1) The requested drug is being prescribed for pain severe and persistent enough to require an extended treatment period with a daily opioid analgesic in a member who has been taking an opioid AND 2) The member can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The member has been evaluated and will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a member who has been receiving an extended-release opioid agent for at least 30 days OR the member has taken an immediate-release opioid for at least one week.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

MOUNJARO

Products Affected

- Mounjaro

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

MOVANTIK

Products Affected

- Movantik

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Must have documented diagnosis of opioid-induced constipation caused by opioids taken for chronic non-cancer pain, including members with chronic pain related to prior cancer or its treatment who do not require frequent (e.g., weekly) opioid dosage escalation
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MYCAPSSA

Products Affected

- Mycapssa

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

MYFEMBREE

Products Affected

- Myfembree

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For heavy menstrual bleeding associated with uterine leiomyomas (fibroids) and moderate to severe pain associated with endometriosis in a premenopausal member: the member has not already received greater than or equal to 24 months of treatment with the requested drug.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months, max 24 months total
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

MYQORZO

Products Affected

- Myqorzo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Concomitant use with rifampin
Required Medical Information	To improve functional capacity and symptoms in obstructive hypertrophic cardiomyopathy: 1) before initiating therapy, member has left ventricular ejection fraction (LVEF) of 55 percent or greater 2) member has New York Heart Association (NYHA) class II-III symptoms and 3) member has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following at a maximally tolerated dose: a) beta blocker (e.g., bisoprolol, propranolol) or b) calcium channel blocker (e.g., verapamil, diltiazem) AND 4) Requested drug will not be used in combination with another cardiac myosin inhibitor (e.g., Camzyos).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

NEMLUVIO

Products Affected

- Nemluvio

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

NERLYNX

Products Affected

- Nerlynx

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent human epidermal growth factor receptor 2 (HER2)-positive breast cancer, brain metastases from HER2-positive breast cancer.
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NEXLETOL

Products Affected

- Nexletol

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NEXLIZET

Products Affected

- Nexlizet

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NGENLA

Products Affected

- Ngenla

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

NILOTINIB

Products Affected

- Nilotinib D-tartrate
- Nilotinib Hydrochloride

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), gastrointestinal stromal tumor (GIST), myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in the chronic phase or blast phase, pigmented villonodular synovitis/tenosynovial giant cell tumor, cutaneous melanoma.
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML), including members newly diagnosed with CML and members who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, AND 2) If member experienced resistance to an alternative tyrosine kinase inhibitor for CML, member is negative for T315I, Y253H, E255K/V, and F359V/C/I mutations. For acute lymphoblastic leukemia (ALL), including members who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, AND 2) If the member has experienced resistance to an alternative tyrosine kinase inhibitor for ALL, member is negative for T315I, Y253H, E255K/V, F359V/C/I and G250E mutations. For gastrointestinal stromal tumor (GIST): 1) Disease is residual, unresectable, recurrent/progressive, or metastatic/tumor rupture, AND 2) Disease has progressed on at least 2 FDA approved therapies (e.g. imatinib, sunitinib, regorafenib, ripretinib). For cutaneous melanoma: 1) Disease is metastatic or unresectable, AND 2) Disease is positive for c-KIT activating mutations, AND 3) Requested drug will be used as subsequent therapy, AND 4) Member has had disease progression, intolerance, or risk of progression with BRAF-targeted therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A

Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Relapsed/refractory systemic light chain amyloidosis, Waldenstrom macroglobulinemia, lymphoplasmacytic lymphoma
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NINTEDANIB

Products Affected

- Nintedanib Esylate
- Ofev

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For idiopathic pulmonary fibrosis (new starts only): 1) a high-resolution computed tomography (HRCT) study of the chest or a lung biopsy reveals the usual interstitial pneumonia (UIP) pattern, OR 2) HRCT study of the chest reveals a result other than the UIP pattern (e.g., probable UIP, indeterminate for UIP) and the diagnosis is supported either by a lung biopsy or by a multidisciplinary discussion between at least a radiologist and pulmonologist who are experienced in idiopathic pulmonary fibrosis if a lung biopsy has not been conducted.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NITISINONE

Products Affected

- Nitisinone
- Orfadin SUSP

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hereditary tyrosinemia type 1 (HT-1): Diagnosis of HT-1 is confirmed by one of the following: 1) biochemical testing (e.g., detection of succinylacetone in urine) OR 2) DNA testing (mutation analysis).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

NUCALA

Products Affected

- Nucala

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

NUEDEXTA

Products Affected

- Dextromethorphan Hydrobromide/quinidine Sulfate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For pseudobulbar affect (PBA) (continuation): The member has experienced a decrease in pseudobulbar affect (PBA) episodes since starting therapy with the requested drug.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 4 months, Continuation: Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hallucinations and delusions associated with Parkinson's disease psychosis, the diagnosis of Parkinson's disease must be made prior to the onset of psychotic symptoms.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

OCTREOTIDE

Products Affected

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

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

ODOMZO

Products Affected

- Odomzo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OGIVRI

Products Affected

- Ogivri

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-positive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	All indications: the member had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information. For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the disease is HER2-positive AND 2) the requested drug is used in combination with paclitaxel and continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OGSIVEO

Products Affected

- Ogsiveo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For relapsed or refractory pediatric low-grade glioma (LGG): the member's tumor is positive for either a) BRAF fusion or rearrangement OR b) BRAF V600 mutation.
Age Restrictions	6 months of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Accelerated or blast phase myeloproliferative neoplasms
Exclusion Criteria	N/A
Required Medical Information	For myelofibrosis, member meets ALL of the following: 1) the member has a diagnosis of intermediate or high-risk primary myelofibrosis or secondary myelofibrosis (i.e., post-polycythemia vera or post-essential thrombocythemia), AND 2) the member has anemia defined as hemoglobin less than 10 grams per deciliter (g/dL) or having transfusion-dependent anemia, AND 3) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to Jakafi (ruxolitinib) OR has hemoglobin less than 8 g/dL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OMNIPOD

Products Affected

- Omnipod 5 Dexcom G7g6 Intro Kit (gen 5)
- Omnipod 5 Dexcom G7g6 Pods (gen 5)
- Omnipod 5 Libre Intro Kit
- Omnipod 5 Libre Pods
- Omnipod Dash Intro Kit (gen 4)
- Omnipod Dash Pdm Kit (gen 4)
- Omnipod Dash Pods (gen 4)

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: 1) The member has diabetes requiring insulin management with multiple daily injections AND 2) The member is self-testing glucose levels 4 or more times per day OR the member is using a continuous glucose monitor AND 3) The member has experienced any of the following with the current diabetes regimen: inadequate glycemic control, recurrent hypoglycemia, wide fluctuations in blood glucose, dawn phenomenon with persistent severe early morning hyperglycemia, severe glycemic excursions.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Peripheral T-cell lymphoma
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OPIPZA

Products Affected

- Opipza

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of schizophrenia, 1) the member meets both of the following: a) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following generic products: aripiprazole, asenapine, lurasidone, olanzapine, quetiapine, risperidone, ziprasidone, AND b) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following brand products: Caplyta, Lybalvi, Rexulti, Secuado, Vraylar, OR 2) The member is unable to swallow oral formulations. For adjunctive treatment of major depressive disorder (MDD), 1) the member meets both of the following: a) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following generic products: aripiprazole, olanzapine, quetiapine, AND b) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following brand products: Rexulti, Vraylar, OR 2) The member is unable to swallow oral formulations. For treatment of irritability associated with autistic disorder: 1) The member experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following generic products: aripiprazole, risperidone, OR 2) The member is unable to swallow oral formulations. For the treatment of Tourette's disorder: 1) The member experienced an inadequate treatment response or intolerance to generic aripiprazole, OR 2) The member is unable to swallow oral formulations.
Age Restrictions	Schizophrenia: 13 years of age or older, Adjunctive treatment of major depressive disorder (MDD): 18 years of age or older, Irritability associated with autistic disorder AND Tourette's disorder: 6 years of age or older
Prescriber Restrictions	N/A

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

ORENITRAM

Products Affected

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

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

ORGOVYX

Products Affected

- Orgovyx

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORIAHNN

Products Affected

- Oriahnn

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For heavy menstrual bleeding associated with uterine leiomyomas (fibroids) in a premenopausal member: the member has not already received greater than or equal to 24 months of treatment with any elagolix-containing drug.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months, max 24 months total
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis (CF): The requested medication will not be used in combination with other medications containing ivacaftor.
Age Restrictions	1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ORLYNVAH

Products Affected

- Orlynvah

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The infection is proven or strongly suspected to be caused by susceptible bacteria based on: 1) culture and susceptibility OR 2) local epidemiology and susceptibility patterns.
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.

ORSERDU

Products Affected

- Orserdu

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OSPHENA

Products Affected

- Osphena

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

OTEZLA

Products Affected

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

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

OXYBATE

Products Affected

- Sodium Oxybate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of excessive daytime sleepiness in a member with narcolepsy, initial request: 1) The diagnosis has been confirmed by sleep lab evaluation, AND 2) The member meets one of the following criteria: a) If the member is 18 years of age or older, the member has experienced an inadequate treatment response or intolerance to at least one central nervous system (CNS) wakefulness promoting drug (e.g., armodafinil, modafinil), OR has a contraindication that would prohibit a trial of central nervous system (CNS) wakefulness promoting drugs (e.g., armodafinil, modafinil). For the treatment of cataplexy in a member with narcolepsy, initial request: The diagnosis has been confirmed by sleep lab evaluation. If the request is for a continuation of therapy, then the member experienced a decrease in daytime sleepiness with narcolepsy or a decrease in cataplexy episodes with narcolepsy.
Age Restrictions	7 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with a sleep disorder specialist or neurologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

OZEMPIC

Products Affected

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

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

PALSONIFY

Products Affected

- Palsonify

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PALYNZIQ

Products Affected

- Palynziq

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PANRETIN

Products Affected

- Panretin

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Topical treatment of cutaneous lesions in members with non-AIDS-related Kaposi sarcoma
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PAROXETINE

Products Affected

- Paroxetine Hcl TABS 30MG, 40MG
- Paroxetine Hcl Er
- Paroxetine Hydrochloride SUSP
- Paroxetine Hydrochloride TABS 10MG, 20MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PAZOPANIB

Products Affected

- Pazopanib Hydrochloride

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Thyroid carcinoma (follicular, papillary, oncocytic, or medullary), uterine sarcoma, chondrosarcoma, gastrointestinal stromal tumor, dedifferentiated liposarcoma
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma: 1) the disease is advanced, relapsed, or stage IV, OR 2) the requested drug will be used for von Hippel-Lindau (VHL)-associated renal cell carcinoma. For gastrointestinal stromal tumor (GIST): 1) the disease is residual, unresectable, recurrent, or metastatic/tumor rupture AND 2) the member meets one of the following: a) the disease has progressed after at least two FDA-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib), b) the disease is succinate dehydrogenase (SDH)-deficient GIST. For soft tissue sarcoma (STS): the member does not have an adipocytic soft tissue sarcoma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PEGASYS

Products Affected

- Pegasys

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

PEMAZYRE

Products Affected

- Pemazyre

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PERAMPANEL

Products Affected

- Perampanel

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of partial-onset seizures (i.e., focal-onset seizures): 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND 2) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to any of the following: Aptiom, Xcopri, Spritam. For adjunctive treatment of primary generalized tonic-clonic seizures: 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND 2) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to Spritam.
Age Restrictions	Partial-onset seizures (i.e., focal-onset seizures): 4 years of age or older. Primary generalized tonic-clonic seizures: 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PERPHENAZINE/AMITRIPTYLINE

Products Affected

- Perphenazine/amitriptyline

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For depression: The member has had an inadequate treatment response, contraindication, or intolerance to two of the following alternative drugs: SSRIs (selective serotonin reuptake inhibitors), SNRIs (serotonin-norepinephrine reuptake inhibitors), including but not limited to bupropion, mirtazapine, or trazodone. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PHENOBARBITAL

Products Affected

- Phenobarbital ELIX 20MG/5ML
- Phenobarbital TABS 100MG, 15MG, 16.2MG, 30MG, 32.4MG, 60MG, 64.8MG, 97.2MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For all indications: Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PHENYLBUTYRATE

Products Affected

- Olpruva
- Sodium Phenylbutyrate POWD 3GM/TSP
- Sodium Phenylbutyrate TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For urea cycle disorders (UCD) that cannot be managed by dietary protein restriction and/or amino acid supplementation alone: Diagnosis of UCD was confirmed by enzymatic, biochemical, or genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT 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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, phosphatidylinositol-3-kinase catalytic alpha subunit (PIK3CA)-mutated breast cancer in combination with fulvestrant
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PIRFENIDONE

Products Affected

- Pirfenidone

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For idiopathic pulmonary fibrosis (new starts only): 1) a high-resolution computed tomography (HRCT) study of the chest or a lung biopsy reveals the usual interstitial pneumonia (UIP) pattern, OR 2) HRCT study of the chest reveals a result other than the UIP pattern (e.g., probable UIP, indeterminate for UIP) and the diagnosis is supported either by a lung biopsy or by a multidisciplinary discussion between at least a radiologist and pulmonologist who are experienced in idiopathic pulmonary fibrosis if a lung biopsy has not been conducted.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

POMALIDOMIDE

Products Affected

- Pomalidomide
- Pomalyst

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Relapsed/refractory systemic light chain amyloidosis, primary central nervous system (CNS) lymphoma
Exclusion Criteria	N/A
Required Medical Information	For multiple myeloma, member has previously received at least two prior therapies, including an immunomodulatory agent AND a proteasome inhibitor.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

POSACONAZOLE

Products Affected

- Posaconazole SUSP

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug will be used orally. For treatment of oropharyngeal candidiasis: member has experienced an inadequate treatment response, intolerance, or has a contraindication to fluconazole.
Age Restrictions	13 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Oropharyngeal candidiasis: 1 month. All other indications: 6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PREGABALIN

Products Affected

- Pregabalin CAPS
- Pregabalin SOLN
- Pregabalin Er

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

PREVYMIS

Products Affected

- Prevymis PACK
- Prevymis TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For prophylaxis of cytomegalovirus (CMV) infection or disease in hematopoietic stem cell transplant (HSCT): 1) the member is CMV-seropositive, AND 2) the member is a recipient of an allogeneic HSCT. For prophylaxis of CMV disease in kidney transplant: 1) the member is CMV-seronegative, AND 2) the member is a high risk recipient of kidney transplant.
Age Restrictions	For prophylaxis of cytomegalovirus (CMV) infection or disease in hematopoietic stem cell transplant (HSCT): 6 months of age or older, For prophylaxis of CMV disease in kidney transplant: 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	7 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

PROCRIT

Products Affected

- Procrit

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

PROMETHAZINE

Products Affected

- Promethazine Hydrochloride SOLN 6.25MG/5ML
- Promethazine Hydrochloride TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For rhinitis: 1) The member has had an inadequate treatment response, contraindication, or intolerance to two of the following non-HRM (high-risk medications) alternative drugs: including but not limited to levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 2) If the member is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Prior Authorization applies to greater than cumulative 30 days of therapy per year.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PYRUKYND

Products Affected

- Pyrukynd
- Pyrukynd Taper Pack

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hemolytic anemia in a member with pyruvate kinase (PK) deficiency: Diagnosis was confirmed by an enzyme assay demonstrating deficiency of PK enzyme activity or by genetic testing. For hemolytic anemia in a member with PK deficiency (continuation of therapy): Member achieved or maintained a positive clinical response (e.g., improvement in hemoglobin levels, reduction in blood transfusions).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 7 months, Continuation: Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Gastrointestinal stromal tumor (GIST) for residual, unresectable, tumor rupture, recurrent, or progressive disease. Metastatic or unresectable cutaneous melanoma.
Exclusion Criteria	N/A
Required Medical Information	For residual, unresectable, tumor rupture, advanced, recurrent/metastatic, or progressive gastrointestinal stromal tumor (GIST): 1) Member has received prior treatment with 3 or more kinase inhibitors, including imatinib OR 2) Member has experienced disease progression following treatment with avapritinib and dasatinib OR 3) Member has received prior treatment with imatinib and is intolerant of second-line sunitinib. For cutaneous melanoma: 1) Disease is metastatic or unresectable AND 2) Disease is positive for KIT activating mutations AND 3) Requested drug will be used as subsequent therapy AND 4) Member has had disease progression, intolerance, or risk of progression with BRAF-targeted therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Babesiosis, uncomplicated Plasmodium vivax malaria.
Exclusion Criteria	N/A
Required Medical Information	For babesiosis: the requested drug is used in combination with clindamycin.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 month
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

RECORLEV

Products Affected

- Recorlev

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REDEMPLO

Products Affected

- Redemplo

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

RELISTOR INJ

Products Affected

- Relistor INJ

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of opioid-induced constipation in a member with chronic non-cancer pain, including chronic pain related to prior cancer or its treatment who does not require frequent (e.g., weekly) opioid dosage escalation: 1) the member is unable to tolerate oral medications, OR 2) the member meets one of the following criteria: A) experienced an inadequate treatment response or intolerance to an oral drug indicated for opioid-induced constipation in a member with chronic non-cancer pain (e.g., Movantik), OR B) the member has a contraindication that would prohibit a trial of an oral drug indicated for opioid-induced constipation in a member with chronic non-cancer pain (e.g., Movantik).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RELISTOR TAB

Products Affected

- Relistor TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RENFLEXIS

Products Affected

- Renflexis

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Behcet's disease, hidradenitis suppurativa, sarcoidosis, Takayasu's arteritis, uveitis
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): 1) Member meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) or leflunomide OR b) intolerance or contraindication to MTX AND leflunomide, AND 2) member meets ANY of the following: a) inadequate treatment response, intolerance or contraindication to MTX OR b) inadequate treatment response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis (new starts only): an inadequate treatment response or intolerance to a non-steroidal anti-inflammatory drug (NSAID) OR contraindication that would prohibit a trial of NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at time of diagnosis, AND 2) member meets ANY of the following: a) member has experienced inadequate treatment response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with MTX, cyclosporine, or acitretin, OR b) pharmacologic treatment with MTX, cyclosporine, or acitretin is contraindicated, OR c) member has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of BSA or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	For hidradenitis suppurativa (new starts only): Member has severe, refractory disease. For uveitis (new starts only): Inadequate treatment response or intolerance or has a contraindication to a trial of immunosuppressive therapy for uveitis.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

REPATHA

Products Affected

- Repatha
- Repatha Sureclick

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RETACRIT

Products Affected

- Retacrit

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

RETEVMO

Products Affected

- Retevmo TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent rearranged during transfection (RET)-rearrangement positive non-small cell lung cancer (NSCLC), brain metastases from RET fusion-positive NSCLC, Langerhans Cell Histiocytosis with a RET gene fusion, symptomatic or relapsed/refractory Erdheim-Chester Disease with a RET gene fusion, symptomatic or relapsed/refractory Rosai-Dorfman Disease with a RET gene fusion, occult primary cancer with RET gene fusion, solid tumors with RET-gene fusion for recurrent disease
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC), member must meet all of the following: 1) The disease is recurrent, advanced or metastatic, AND 2) The tumor is rearranged during transfection (RET) fusion-positive or RET rearrangement positive. For solid tumors, member must meet all of the following: 1) The disease is recurrent, persistent, progressive, unresectable, locally advanced, or metastatic, 2) The member has progressed on or following prior systemic treatment or has no satisfactory alternative treatment options, AND 3) The tumor is RET fusion-positive.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

REVUFORJ

Products Affected

- Revuforj

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

REZDIFFRA

Products Affected

- Rezdiffra

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

REZLIDHIA

Products Affected

- Rezlidhia

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

RHAPSIDO

Products Affected

- Rhapsido

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an allergists, dermatologists or immunologists.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RINVOQ

Products Affected

- Rinvoq
- Rinvoq Lq

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For new starts of the following conditions: moderate to severe active rheumatoid arthritis, active psoriatic arthritis, moderate to severe active ulcerative colitis, moderately to severely active Crohn's disease, active ankylosing spondylitis, OR non-radiographic axial spondyloarthritis. The member must have experienced an inadequate treatment response, intolerance or contraindication to at least one tumor necrosis factor (TNF) inhibitor (e.g., Yuflyma [adalimumab-aaty], Enbrel [etanercept], Hulio [adalimumab-fkjp]). For Crohn's disease and UC only: If TNF blockers are clinically inadvisable, member has had an inadequate response to treatment with at least one other systemic drug product or use of these therapies are inadvisable. For atopic dermatitis (new starts only): 1) The member has refractory, moderate to severe disease, AND 2) Member has had an inadequate response to treatment with at least one other systemic drug product, including biologics, or use of these therapies are inadvisable. For atopic dermatitis (continuation of therapy): The member has achieved or maintained positive clinical response.
Age Restrictions	Atopic dermatitis: 12 years of age or older, active psoriatic arthritis, aPJIA: 2 years of age or older, All other indications: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	AD Initial: 4 months, AD Continuation: Plan Year, All others: Plan Year

Other Criteria	For active polyarticular juvenile idiopathic arthritis and with giant cell arteritis(new starts only): member has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one tumor necrosis factor (TNF) inhibitor (e.g., Enbrel [etanercept], Hulio [adalimumab-fkjp], Yuflyma [adalimumab-aaty]).
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

RIVFLOZA

Products Affected

- Rivfloza

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For primary hyperoxaluria type 1 (PH1): diagnosis has been confirmed by a molecular genetic test showing a mutation in the alanine:glyoxylate aminotransferase (AGXT) gene or liver enzyme analysis demonstrating absent or significantly reduced alanine:glyoxylate aminotransferase (AGT) activity. For PH1 (continuation): the member has experienced decreased or normalized levels of urinary oxalate since initiating therapy.
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ROMVIMZA

Products Affected

- Romvimza

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ROZLYTREK

Products Affected

- Rozlytrek

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent ROS1-positive non-small cell lung cancer (NSCLC), Non-metastatic neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, first-line treatment of NTRK gene fusion-positive solid tumors, ROS1-gene fusion-positive cutaneous melanoma
Exclusion Criteria	N/A
Required Medical Information	For all neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors: the disease is without a known acquired resistance mutation. For ROS1-positive non-small cell lung cancer: the member has recurrent, advanced, or metastatic disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

RUFINAMIDE

Products Affected

- Rufinamide SUSP 40MG/ML
- Rufinamide TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RYDAPT

Products Affected

- Rydapt

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Re-induction in residual disease for AML, maintenance therapy for AML, myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FGFR1 or FLT3 rearrangements
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML): AML is FMS-like tyrosine kinase 3 (FLT3) mutation-positive. For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and Fibroblast growth factor receptor type 1 (FGFR1) or FLT3 rearrangements: the disease is in chronic or blast phase.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

RYZNEUTA

Products Affected

- Ryzneuta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For prophylaxis of myelosuppressive chemotherapy-induced febrile neutropenia, the member must meet all of the following: 1) Member has non-myeloid cancer, AND 2) Member is currently receiving or will be receiving treatment with myelosuppressive anti-cancer therapy, AND 3) The requested drug will be administered at least 24 hours after chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SAPROPTERIN

Products Affected

- Javygtor
- Sapropterin Dihydrochloride
- Zelvysia

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For phenylketonuria (PKU): For members who have not yet received a therapeutic trial of the requested drug, the member's pretreatment (including before dietary management) phenylalanine level is greater than 6 mg/dL (360 micromol/L). For members who completed a therapeutic trial of the requested drug, the member must have experienced improvement (e.g., reduction in blood phenylalanine levels, improvement in neuropsychiatric symptoms).
Age Restrictions	1 month of age or older
Prescriber Restrictions	N/A
Coverage Duration	Initial: 2 months, All others: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SAVELLA

Products Affected

- Milnacipran Hydrochloride
- Milnacipran Hydrochloride Titration Pack
- Savella
- Savella Titration Pack

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For fibromyalgia: The member has experienced an inadequate treatment response, intolerance, or has a contraindication to duloxetine or pregabalin.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SCEMBLIX

Products Affected

- Scemblix

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in chronic phase or blast phase.
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML) in the chronic phase: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, AND 2) Member meets one of the following: A) Member has newly diagnosed CML and has resistance or intolerance to imatinib, dasatinib, or nilotinib OR B) Member has previously treated CML AND at least one of the prior treatments was imatinib, dasatinib, or nilotinib OR C) Member is positive for the T315I mutation, AND 3) Member is negative for the following mutations: A337T, P465S, and F359V/I/C.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SEMAGLUTIDE ORAL

Products Affected

- Ozempic TABS
- Rybelsus TABS 14MG, 3MG, 7MG

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

SEPHIENCE

Products Affected

- Sephience

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member meets all of the following criteria: 1) Documented diagnosis of hyperphenylalaninemia (HPA) with sepiapterin-responsive phenylketonuria (PKU) AND 2) the medication is used in conjunction with a phenylalanine (Phe)-restricted diet.
Age Restrictions	1 month of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SEROSTIM

Products Affected

- Serostim INJ 4MG, 5MG, 6MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of human immunodeficiency virus (HIV) members with wasting or cachexia: 1) The requested medication is used in combination with antiretroviral therapy . For continuation of therapy: Member must have demonstrated a response to therapy with the requested medication (i.e., body mass index [BMI] has increased or stabilized).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	48 weeks
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SIGNIFOR

Products Affected

- Signifor

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SILDENAFIL (PAH)

Products Affected

- Sildenafil Citrate SUSR
- Sildenafil Citrate TABS 20MG

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

SIRTURO

Products Affected

- Sirturo

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

SKYCLARYS

Products Affected

- Skyclarys

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

SKYRIZI

Products Affected

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

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

SOFOSBUVIR/VELPATASVIR

Products Affected

- Sofosbuvir/velpatasvir

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases and Infectious Diseases Society of America (AASLD-IDSA) treatment guidelines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
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.

SOGROYA

Products Affected

- Sogroya

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

SOHONOS

Products Affected

- Sohonos

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For reduction in volume of new heterotopic ossification in fibrodysplasia ossificans progressiva (FOP): The member has a confirmed genetic mutation in the activin A receptor type I (ACVR1) gene.
Age Restrictions	8 years of age or older if female and 10 years of age or older if male
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

SORAFENIB

Products Affected

- Sorafenib Tosylate TABS

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Acute myeloid leukemia, soft tissue sarcoma (angiosarcoma, desmoid tumors/aggressive fibromatosis, and solitary fibrous tumor subtypes), gastrointestinal stromal tumor, medullary thyroid carcinoma, osteosarcoma, recurrent chordoma, epithelial ovarian cancer, fallopian tube cancer, primary peritoneal cancer, lymphoid and/or myeloid neoplasms with eosinophilia and FLT3 rearrangement in chronic or blast phase
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia: the disease is FMS-like tyrosine kinase 3-internal tandem duplication (FLT3-ITD) mutation-positive and any of the following is met :1) the requested drug will be used as maintenance therapy after hematopoietic stem cell transplant, OR 2) the requested drug is being used for low-intensity treatment induction, post-induction therapy, or consolidation therapy, OR 3) the disease is relapsed/refractory. For thyroid carcinoma: histology is follicular, papillary, oncocytic, or medullary. For gastrointestinal stromal tumor (GIST): 1) the disease is residual, unresectable, recurrent, or metastatic/tumor rupture, AND 2) the disease has progressed after use of at least two FDA-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

SPEVIGO

Products Affected

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

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

STIVARGA

Products Affected

- Stivarga

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For colorectal cancer: 1) The disease is advanced or metastatic, AND 2) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to Lonsurf (trifluridine/tipiracil). For hepatocellular carcinoma (HCC): the member has been previously treated with sorafenib. For locally advanced, unresectable or metastatic gastrointestinal stromal tumor (GIST): the member has been previously treated with imatinib mesylate and sunitinib malate.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

STRENSIQ

Products Affected

- Strensiq

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must have a diagnosis of perinatal/infantile- and juvenile-onset hypophosphatasia (HPP).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist or metabolic/genetic specialist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SUNITINIB

Products Affected

- Sunitinib Malate

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Thyroid carcinoma (follicular, medullary, papillary, and oncocytic), soft tissue sarcoma (angiosarcoma, solitary fibrous tumor, and alveolar soft part sarcoma subtypes), recurrent chordoma, thymic carcinoma, lymphoid and/or myeloid neoplasms with eosinophilia and FLT3 rearrangement in chronic or blast phase, pheochromocytoma, paraganglioma, well differentiated grade 3 neuroendocrine tumors
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma (RCC): 1) The disease is relapsed, advanced, or stage IV OR 2) the requested drug is being used as adjuvant treatment for members that are at high risk of recurrent RCC following nephrectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

SUNOSI

Products Affected

- Sunosi

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

SYMPAZAN

Products Affected

- Sympazan

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Seizures associated with Dravet syndrome
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Seizures associated with Lennox-Gastaut syndrome (LGS): 2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TABLOID

Products Affected

- Tabloid

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

TABRECTA

Products Affected

- Tabrecta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent non-small cell lung cancer (NSCLC), NSCLC with high-level mesenchymal-epithelial transition (MET) amplification, central nervous system (CNS) brain metastases from MET exon-14 mutated NSCLC
Exclusion Criteria	N/A
Required Medical Information	For recurrent, advanced, or metastatic non-small cell lung cancer (NSCLC): Tumor is positive for mesenchymal-epithelial transition (MET) exon 14 skipping mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TADALAFIL (BPH)

Products Affected

- Tadalafil TABS 2.5MG, 5MG

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

TADALAFIL (PAH)

Products Affected

- Alyq
- Tadalafil TABS 20MG

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

TAFINLAR

Products Affected

- Tafinlar

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Langerhans cell histiocytosis, Erdheim-Chester disease, hairy cell leukemia
Exclusion Criteria	N/A
Required Medical Information	For melanoma: 1) The tumor is positive for a BRAF V600 activating mutation (e.g., V600E or V600K), AND 2) The requested drug will be used as a single agent or in combination with trametinib, AND 3) The requested drug will be used for either of the following: a) unresectable, limited resectable, or metastatic disease, b) adjuvant systemic therapy. For non-small cell lung cancer: 1) The tumor is positive for a BRAF V600E mutation, AND 2) The requested drug will be used as a single agent or in combination with trametinib. For papillary, follicular, and oncocytic thyroid carcinoma: 1) The tumor is BRAF V600E-positive, AND 2) The disease is not amenable to radioactive iodine (RAI) therapy, AND 3) the requested drug will be used in combination with trametinib. For Langerhans Cell Histiocytosis and Erdheim-Chester Disease: The disease is positive for a BRAF V600E mutation. For hairy cell leukemia: 1) the requested drug will be used in combination with trametinib, AND 2) the member has not had previous treatment with BRAF inhibitor therapy. For solid tumors: 1) The tumor is positive for a BRAF V600E mutation, AND 2) The requested drug will be used in combination with trametinib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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	Pending CMS Review
Off-Label Uses	Pending CMS Review
Exclusion Criteria	Pending CMS Review
Required Medical Information	Pending CMS Review
Age Restrictions	Pending CMS Review
Prescriber Restrictions	Pending CMS Review
Coverage Duration	Pending CMS Review
Other Criteria	Pending CMS Review
Prerequisite Therapy Required	Pending CMS Review

TALZENNA

Products Affected

- Talzenna

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent germline breast cancer susceptibility gene (BRCA)-mutated breast cancer
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TARPEYO

Products Affected

- Tarpeyo

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

TASIMELTEON

Products Affected

- Tasimelteon

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For Non-24-Hour Sleep-Wake Disorder: 1) For initial therapy and continuation of therapy the member must meet both of the following: a) diagnosis of total blindness in both eyes (e.g., nonfunctioning retinas) and b) unable to perceive light in either eye, AND 2) If currently on therapy with the requested drug, member must meet at least one of the following: a) increased total nighttime sleep or b) decreased daytime nap duration. For nighttime sleep disturbances in Smith-Magenis Syndrome (SMS): 1) For initial therapy and continuation therapy, the member has a confirmed diagnosis of SMS, AND 2) If currently on therapy with the requested drug, the member experienced improvement in the quality of sleep since starting therapy.
Age Restrictions	Non-24: 18 years of age or older, SMS: 16 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with a sleep disorder specialist, neurologist, or psychiatrist
Coverage Duration	Initiation: 6 months, Renewal: Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For continuation of treatment for severe anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis: the member has experienced benefit from therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TAZAROTENE

Products Affected

- Tazarotene CREA
- Tazarotene GEL
- Trilitas

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

TAZVERIK

Products Affected

- Tazverik

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Epithelioid sarcoma: 16 years of age or older, Follicular lymphoma: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TEMAZEPAM

Products Affected

- Temazepam

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For short-term treatment of insomnia: 1) The prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for the member AND 2) The member has experienced an inadequate treatment response, has a contraindication, or intolerance to doxepin (3 mg or 6 mg) or ramelteon AND 3) If the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TEPMETKO

Products Affected

- Tepmetko

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent non-small cell lung cancer (NSCLC), NSCLC with high level mesenchymal-epithelial transition (MET) amplification, central nervous system (CNS) cancer including brain metastases and leptomeningeal metastases from MET exon-14 mutated NSCLC
Exclusion Criteria	N/A
Required Medical Information	For recurrent, advanced, or metastatic non-small cell lung cancer (NSCLC): Tumor is positive for mesenchymal-epithelial transition (MET) exon 14 skipping mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TERIFLUNOMIDE

Products Affected

- Teriflunomide

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TERIPARATIDE

Products Affected

- Teriparatide

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

TESTOSTERONE CYPIONATE INJ

Products Affected

- Depo-testosterone INJ 100MG/ML, 200MG/ML
- Testosterone Cypionate INJ 100MG/ML, 200MG/ML

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

TESTOSTERONE ENANTHATE INJ

Products Affected

- Testosterone Enanthate INJ

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

TETRABENAZINE

Products Affected

- Tetrabenazine

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

THALOMID

Products Affected

- Thalomid CAPS 100MG, 50MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Acquired immunodeficiency syndrome (AIDS)-related aphthous stomatitis, Kaposi sarcoma, multicentric Castleman's disease, Rosai-Dorfman disease, Langerhans cell histiocytosis, pediatric medulloblastoma
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Conventional (grades 1-3) or dedifferentiated chondrosarcoma, central nervous system (CNS) cancers (astrocytoma, oligodendroglioma)
Exclusion Criteria	N/A
Required Medical Information	Member has disease with a susceptible isocitrate dehydrogenase-1 (IDH1) mutation. For acute myeloid leukemia (AML): 1) member has newly-diagnosed AML and meets one of the following: a) 75 years of age or older, b) member has comorbidities that preclude use of intensive induction chemotherapy, OR 2) the requested drug will be used as post-induction therapy following response to induction therapy with the requested drug, OR 3) member has relapsed or refractory AML. For locally advanced, unresectable, resected gross residual, or metastatic cholangiocarcinoma: the requested drug will be used as subsequent treatment for progression on or after systemic treatment. For CNS cancers: 1) disease is recurrent or progressive, AND 2) member has oligodendroglioma or astrocytoma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TOBI INHALER

Products Affected

- Tobi Podhaler

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-cystic fibrosis bronchiectasis
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis and non-cystic fibrosis bronchiectasis: 1) Pseudomonas aeruginosa is present in the member's airway cultures, OR 2) The member has a history of Pseudomonas aeruginosa infection or colonization in the airways.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TOPICAL TESTOSTERONES

Products Affected

- Testosterone GEL 20.25MG/1.25GM, 25MG/2.5GM, 40.5MG/2.5GM, 50MG/5GM
- Testosterone SOLN
- Testosterone Pump GEL 12.5MG/ACT, 20.25MG/ACT

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

TOPICAL TRETINOIN

Products Affected

- Tretinoin CREA
- Tretinoin GEL

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TOPIRAMATE

Products Affected

- Topiramate SOLN 25MG/ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For treatment of partial-onset seizures (i.e., focal-onset seizures): 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND 2) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to any of the following: Aptiom (if 4 years of age or older), Xcopri (if 18 years of age or older), Spritam (if 4 years of age or older). For monotherapy treatment of primary generalized tonic-clonic seizures: 1) The member has experienced an inadequate treatment response or intolerance to a generic topiramate immediate release product, OR 2) The member has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules). For adjunctive treatment of primary generalized tonic-clonic seizures: 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND 2) If the member is 6 years of age or older, the member has experienced an inadequate treatment response, intolerance, or has a contraindication to Spritam. For the preventative treatment of migraines: 1) The member has experienced an inadequate treatment response or intolerance to a generic topiramate immediate release product, OR 2) The member has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules).
Age Restrictions	Epilepsy: 2 years of age or older, Migraine: 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A

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

TRAMADOL - IR BEFORE ER

Products Affected

- Tramadol Hcl Er CP24 100MG, 200MG, 300MG
- Tramadol Hcl Er TB24
- Tramadol Hydrochloride Er

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the member meets all of the following: 1) The requested drug is being prescribed for pain severe and persistent enough to require an extended treatment period with a daily opioid analgesic in a member who has been taking an opioid AND 2) The member can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The member has been evaluated and will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a member who has been receiving an extended-release opioid agent for at least 30 days OR the member has taken an immediate-release opioid for at least one week.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TRAZIMERA

Products Affected

- Trazimera

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer (including appendiceal adenocarcinoma), HER2-positive recurrent salivary gland tumor, HER2-positive unresectable or metastatic hepatobiliary carcinoma (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma), HER2 overexpression positive locally advanced, unresectable, or recurrent gastric adenocarcinoma, HER2-postiive endometrial cancer.
Exclusion Criteria	N/A
Required Medical Information	For colorectal cancer (including appendiceal adenocarcinoma): 1) the disease is HER2-amplified and RAS and BRAF wild-type and 2) the requested drug is used in combination with pertuzumab, tucatinib or lapatinib and 3) the member has not had previous treatment with a HER2 inhibitor. For hepatobiliary carcinoma: 1) the disease is HER2 positive and 2) the requested drug is used in combination with pertuzumab. For endometrial cancer: 1) the requested drug is being used in combination with carboplatin and paclitaxel and 2) continued as a single agent for maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year

Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRELSTAR

Products Affected

- Trelstar Mixject

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Gender dysphoria, ovarian suppression in breast cancer
Exclusion Criteria	N/A
Required Medical Information	For gender dysphoria, member meets ONE of the following): 1) the requested drug is used to suppress puberty and the member is at Tanner stage 2 or greater, OR 2) member is undergoing gender transition, and the member will receive the requested drug concomitantly with gender-affirming hormones. For breast cancer, member meets ALL of the following: 1) the requested drug is being used for ovarian suppression in premenopausal members, and 2) the requested drug will be used in combination with endocrine therapy, and 3) the disease is hormone receptor positive, and 4) the disease is at a higher risk of recurrence (e.g., young age, high-grade tumor, lymph-node involvement).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TREMFYA

Products Affected

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

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

TRIKAFTA

Products Affected

- Trikafta

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis: The requested medication will not be used in combination with other medications containing ivacaftor.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRULICITY

Products Affected

- Trulicity

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

TRUQAP

Products Affected

- Truqap TABS 200MG
- Truqap TBPK

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TRUXIMA

Products Affected

- Truxima

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-Hodgkin's lymphoma subtypes [small lymphocytic lymphoma (SLL), mantle cell lymphoma, marginal zone lymphomas (nodal, splenic, extranodal marginal zone lymphoma (EMZL) of the stomach, EMZL of nongastric sites (noncutaneous)), Burkitt lymphoma, high-grade B-cell lymphoma, histological transformation of indolent lymphomas to diffuse large B-cell lymphoma, histological transformation of chronic lymphocytic leukemia (CLL)/SLL to diffuse large B-cell lymphoma, primary cutaneous B-cell lymphoma, Castleman disease, human immunodeficiency virus (HIV)-related B-cell lymphoma, hairy cell leukemia, post-transplant lymphoproliferative disorder (PTLD), B-cell lymphoblastic lymphoma], refractory immune or idiopathic thrombocytopenic purpura (ITP), autoimmune hemolytic anemia, Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma, chronic graft-versus-host disease (GVHD), Sjogren syndrome, thrombotic thrombocytopenic purpura, refractory myasthenia gravis, nodular lymphocyte-predominant Hodgkin lymphoma, primary central nervous system (CNS) lymphoma, leptomeningeal metastases from lymphomas, acute lymphoblastic leukemia, prevention of Epstein-Barr virus (EBV)-related PTLD, relapsing remitting multiple sclerosis, immune checkpoint inhibitor-related toxicities, Rosai-Dorfman disease, pediatric aggressive mature B-cell lymphomas (including Burkitt-like lymphoma, primary mediastinal large B-cell lymphoma), and pediatric mature B-cell acute leukemia, neuromyelitis optica spectrum disorder
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): 1) member meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) OR b) member has intolerance or contraindication to MTX, AND 2) member meets ANY of the following: a) inadequate treatment response, intolerance, or contraindication to MTX OR b) inadequate treatment response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. Hematologic malignancies must be CD20-positive
Age Restrictions	N/A

Prescriber Restrictions	N/A
Coverage Duration	Immune checkpoint inhibitor-related toxicities: 3 months, All other: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

TRYNGOLZA

Products Affected

- Tryngolza

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

TRYVIO

Products Affected

- Tryvio

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hypertension: 1) the member is currently taking at least two of the following: a) angiotensin converting enzyme (ACE) inhibitor (e.g., captopril, enalapril), b) angiotensin II receptor blocker (ARB) (e.g., candesartan, valsartan), c) diuretic (e.g., hydrochlorothiazide, chlorthalidone), c) calcium channel blocker (e.g., amlodipine, nifedipine), or e) mineralocorticoid receptor antagonist (MRA) (e.g., eplerenone, spironolactone) at maximally tolerated doses AND 2) for initial therapy, the member's blood pressure is not adequately controlled with their current regimen. For continuation: the member has demonstrated a positive response to therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent human epidermal growth factor receptor 2 (HER2)-positive breast cancer, HER2-positive biliary tract cancer (gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma)
Exclusion Criteria	N/A
Required Medical Information	For colorectal cancer (including appendiceal adenocarcinoma): 1) the member has advanced, unresectable, or metastatic disease, AND 2) the member has human epidermal growth factor receptor 2 (HER2)-positive disease, AND 3) the member has RAS wild-type disease, AND 4) the requested drug will be used in combination with trastuzumab, AND 5) the member has not previously been treated with a HER2 inhibitor. For biliary tract cancer: 1) the member has unresectable or metastatic disease, AND 2) the member has human epidermal growth factor receptor 2 (HER2)-positive disease, AND 3) the requested drug will be used in combination with trastuzumab.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TURALIO

Products Affected

- Turalio CAPS 125MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Langerhans cell histiocytosis, Erdheim-Chester disease, Rosai-Dorfman disease
Exclusion Criteria	N/A
Required Medical Information	For Langerhans cell histiocytosis: 1) disease has colony stimulating factor 1 receptor (CSF1R) mutation. For Erdheim-Chester disease and Rosai-Dorfman disease: 1) disease has CSF1R mutation AND member has any of the following: a) symptomatic disease OR b) relapsed/refractory disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

TYENNE

Products Affected

- Tyenne INJ 162MG/0.9ML

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Rheumatoid Arthritis (RA) (Initial): Must meet all the following: 1) The member must have a documented diagnosis of moderately to severely active RA AND 2) an inadequate treatment response, intolerance or a contraindication to one or more DMARDs (e.g., methotrexate, leflunomide, sulfasalazine, hydroxychloroquine) OR trial and failure, intolerance or contraindication to a targeted immunomodulator FDA-approved for the treatment of RA (e.g., Enbrel, adalimumab, Xeljanz, Rinvoq). Giant Cell Arteritis (GCA) (Initial): The member must have a documented diagnosis of GCA. Systemic Juvenile Idiopathic Arthritis (SJIA) (Initial): Must meet all the following: 1) The member must have a documented diagnosis of active SJIA. Polyarticular Juvenile Idiopathic Arthritis (PJIA) (Initial): Must meet all the following: 1) The member must have a documented diagnosis of active PJIA AND 2) an inadequate treatment response, intolerance or a contraindication to one DMARD (e.g., methotrexate, sulfasalazine or leflunodmide), or one NSAID OR b) for continuation of prior therapy.
Age Restrictions	SJIA, PJIA: 2 years of age or older
Prescriber Restrictions	All indications: prescribed by or in consultation with a rheumatologist
Coverage Duration	Plan Year
Other Criteria	All indications for continuation of therapy: documentation of a 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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For acute treatment of migraine: The member has experienced an inadequate treatment response, intolerance, or the member has a contraindication to at least one triptan 5-HT1 receptor agonist.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

UDENYCA

Products Affected

- Udenyca
- Udenyca Onbody

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Stem cell transplantation-related indications
Exclusion Criteria	N/A
Required Medical Information	If receiving chemotherapy, the requested drug will be administered at least 24 hours after chemotherapy. For prophylaxis of myelosuppressive chemotherapy-induced febrile neutropenia: the member must meet both of the following: 1) Member has a non-myeloid cancer, AND 2) Member is currently receiving or will be receiving treatment with myelosuppressive anti-cancer therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

UPTRAVI

Products Affected

- Uptravi TABS
- Uptravi Titration Pack

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

USTEKINUMAB

Products Affected

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

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

VALCHLOR

Products Affected

- Valchlor

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Smoldering adult T-cell leukemia/lymphoma (ATLL), Stage 2 or higher mycosis fungoides (MF)/Sezary syndrome (SS), primary cutaneous marginal zone lymphoma, primary cutaneous follicle center lymphoma, CD30-positive lymphomatoid papulosis (LyP), unifocal Langerhans cell histiocytosis (LCH) with isolated skin disease
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT 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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the acute treatment of intermittent, stereotypic episodes of frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) that are distinct from a member's usual seizure pattern in members with documented epilepsy
Age Restrictions	2 years of age or older
Prescriber Restrictions	Must be prescribed by a neurologist or must be prescribed after consultation with a neurologist.
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Re-induction in members with residual disease for AML
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML): 1) AML is FMS-like tyrosine kinase 3 (FLT3) internal tandem duplication (ITD)-positive and 2) medication will be used for induction, re-induction, consolidation, or maintenance therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VANRAFIA

Products Affected

- Vanrafia

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For reduction of proteinuria in members with primary immunoglobulin A nephropathy (IgAN) at risk of rapid disease progression: 1) The member had an inadequate response to therapy with a maximally tolerated dose of a renin-angiotensin system (RAS) inhibitor (e.g., angiotensin-converting enzyme [ACE] inhibitor or angiotensin-receptor blocker [ARB]) OR 2) The member experienced an intolerance or has a contraindication to RAS inhibitors.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VENCLEXTA

Products Affected

- Venclexta
- Venclexta Starting Pack

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Mantle cell lymphoma, blastic plasmacytoid dendritic cell neoplasm (BPDCN), multiple myeloma, relapsed or refractory acute myeloid leukemia (AML), poor-risk AML, therapy related AML, post-induction therapy for AML following response to previous lower intensity therapy with the same regimen, Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma, relapsed or refractory systemic light chain amyloidosis with translocation t(11:14), accelerated or blast phase myeloproliferative neoplasms, B-cell acute lymphoblastic leukemia/T-cell acute lymphoblastic leukemia (B-ALL/T-ALL), hairy cell leukemia, higher risk myelodysplastic syndromes, chronic myelomonocytic leukemia (CMML)-2.
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML): 1) member has newly-diagnosed AML and meets one of the following: a) 75 years of age or older, b) member has comorbidities that preclude use of intensive induction chemotherapy, OR 2) will be used for induction or consolidation therapy in members with poor-risk or therapy related AML, OR 3) member has relapsed or refractory AML, OR 4) will be used for post-induction therapy for AML following response to previous lower intensity therapy with the same regimen. For blastic plasmacytoid dendritic cell neoplasm (BPDCN): 1) member has systemic disease being treated with palliative intent, OR 2) member has relapsed or refractory disease. For multiple myeloma: 1) the disease is relapsed or progressive, AND 2) the requested drug will be used in combination with one of the following: a) dexamethasone, b) dexamethasone and daratumumab c) dexamethasone with bortezomib, carfilzomib, or ixazomib AND 3) member has t(11:14) translocation. For Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma: 1) member has previously treated disease that did not respond to primary therapy, OR 2) member has progressive or relapsed disease.
Age Restrictions	N/A

Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VEOZAH

Products Affected

- Veozah

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VEPPANU

Products Affected

- Veppanu

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must meet all of the following requirements: 1) have a documented diagnosis of estrogen receptor (ER)-positive, human epidermal growth factor receptor 2 (HER2)-negative, estrogen receptor-1 (ESR1)-mutated advanced or metastatic breast cancer AND 2) the disease progressed following at least one line of endocrine therapy in combination with a a CDK4/6i (i.e. Ibrance (palbociclib), Kisqali (ribociclib) or Verzenio (abemaciclib)).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VERSACLOZ

Products Affected

- Versacloz

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of a severely ill member with schizophrenia who failed to respond adequately to standard antipsychotic treatment (i.e., treatment-resistant schizophrenia): 1) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following generic products: aripiprazole, asenapine, lurasidone, olanzapine, quetiapine, risperidone, ziprasidone, AND 2) the member has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following brand products: Caplyta, Lybalvi, Rexulti, Secuado, Vraylar. For reducing the risk of recurrent suicidal behavior in schizophrenia or schizoaffective disorder: member is determined to be at chronic risk for re-experiencing suicidal behavior
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VERZENIO

Products Affected

- Verzenio

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent breast cancer, endometrial cancer, in combination with letrozole for estrogen receptor positive tumor
Exclusion Criteria	N/A
Required Medical Information	For breast cancer: 1) the disease is either: a) advanced, recurrent, or metastatic, OR b) early breast cancer AND 2) the member has hormone receptor (HR)-positive and human epidermal growth factor receptor 2 (HER2)-negative disease, AND 3) the requested drug will be used in combination with endocrine therapy or as a single agent.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

V-GO

Products Affected

- V-go 20
- V-go 30
- V-go 40

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Initial: 1) The member has diabetes requiring insulin management with multiple daily injections AND 2) The member is self-testing glucose levels 4 or more times per day OR the member is using a continuous glucose monitor AND 3) The member has experienced any of the following with the current diabetes regimen: inadequate glycemic control, recurrent hypoglycemia, wide fluctuations in blood glucose, dawn phenomenon with persistent severe early morning hyperglycemia, severe glycemic excursions.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VIGABATRIN

Products Affected

- Vigabatrin
- Vigadrone
- Vigafyde

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For complex partial seizures (i.e., focal impaired awareness seizures): member has experienced an inadequate treatment response to at least two antiepileptic drugs for complex partial seizures (i.e., focal impaired awareness seizures).
Age Restrictions	Infantile Spasms: 1 month to 2 years of age. Complex partial seizures (i.e., focal impaired awareness seizures): 2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VIJOICE

Products Affected

- Vijoice

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-metastatic neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, first-line treatment of NTRK gene fusion-positive solid tumors.
Exclusion Criteria	N/A
Required Medical Information	For all neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, the disease is without a known acquired resistance mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Recurrent non-small cell lung cancer (NSCLC)
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): 1) the disease is recurrent, advanced, or metastatic, and 2) the member has sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Accelerated or blast phase myeloproliferative neoplasms
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOQUEZNA

Products Affected

- Voquezna

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Helicobacter pylori (H. pylori): Must meet all the following: 1) The member must have a documented diagnosis of H. pylori infection 2) the requested drug will be used in combination with amoxicillin and clarithromycin or amoxicillin alone AND 3) member has had a trial and failure, contraindication, or intolerance to bismuth quadruple therapy (e.g., bismuth and metronidazole and tetracycline and proton pump inhibitor [PPI]). For both Healing and Maintenance, Relief of Heartburn associated with Erosive Esophagitis (HRH and MHRH): Must meet all the following: 1) The member must have a documented diagnosis of erosive esophagitis AND 2) member has had a 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. Non-Erosive Gastrointestinal Reflux Disease (NERD): Must meet all the following: 1) The member must have a documented diagnosis of non-erosive gastroesophageal reflux disease AND 2) member has had a 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: 14 days NERD: 1 month HRH: 2 months MHRH: 6 months
Other Criteria	N/A

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

VORANIGO

Products Affected

- Voranigo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VORICONAZOLE

Products Affected

- Voriconazole INJ

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

VOSEVI

Products Affected

- Vosevi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh class B or C)
Required Medical Information	For hepatitis C: Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of HIV coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases and Infectious Diseases Society of America (AASLD-IDSA) treatment guidelines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
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.

VOWST

Products Affected

- Vowst

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the prevention of recurrence of Clostridioides difficile infection (CDI): 1) The diagnosis of CDI has been confirmed by a positive stool test for C. difficile toxin, AND 2) The requested drug will be administered at least 48 hours after the last dose of antibiotics used for the treatment of recurrent CDI.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	1 month
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOYDEYA

Products Affected

- Voydeya

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For paroxysmal nocturnal hemoglobinuria (PNH) (initial): 1) the diagnosis of PNH was confirmed by detecting a deficiency of glycosylphosphatidylinositol-anchored proteins (GPI-APs) AND 2) flow cytometry is used to demonstrate GPI-AP deficiency AND 3) the requested drug is being used as add-on therapy to ravulizumab or eculizumab for the treatment of extravascular hemolysis (EVH). For PNH (continuation): 1) there is no evidence of unacceptable toxicity or disease progression while on the current regimen AND 2) the member has demonstrated a positive response to therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VOYXACT

Products Affected

- Voyxact

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For reduction of proteinuria in members with primary immunoglobulin A nephropathy (IgAN) at risk of disease progression: 1) The member had an inadequate response to therapy with a maximally tolerated dose of a renin-angiotensin system (RAS) inhibitor (e.g., angiotensin-converting enzyme [ACE] inhibitor or angiotensin-receptor blocker [ARB]) OR 2) The member experienced an intolerance or has a contraindication to RAS inhibitors.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a nephrologist.
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

VYKAT XR

Products Affected

- Vykat Xr

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Member must have a documented diagnosis of Prader-Willi syndrome (PWS).
Age Restrictions	4 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

VYNDAMAX

Products Affected

- Vyndamax

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

VYVGART HYTRULO

Products Affected

- Vyvgart Hytrulo

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

WAINUA

Products Affected

- Wainua

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For polyneuropathy of hereditary transthyretin (TTR)-mediated amyloidosis, initial therapy: Member is positive for a mutation of the TTR gene and exhibits clinical manifestation of disease (for example, amyloid deposition in biopsy specimens, TTR protein variants in serum, progressive peripheral sensory-motor polyneuropathy). For polyneuropathy of hereditary TTR-mediated amyloidosis, continuation: Member demonstrates a beneficial response to therapy (for example, improvement of neuropathy severity and rate of disease progression).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

WEGOVY

Products Affected

- Wegovy

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

WELIREG

Products Affected

- Welireg

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Pheochromocytoma or Paraganglioma (PPGL): 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

WINREVAIR

Products Affected

- Winrevair

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

XALKORI

Products Affected

- Xalkori

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

XATMEP

Products Affected

- Xatmep

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The member must have a diagnosis of acute lymphoblastic leukemia (ALL) or polyarticular juvenile idiopathic arthritis (pJIA).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Subject to Part B vs D review.
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

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

XERMELO

Products Affected

- Xermelo

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XGEVA

Products Affected

- Osenvelt
- Wyost

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For hypercalcemia of malignancy: condition is refractory to intravenous (IV) bisphosphonate therapy or there is a clinical reason to avoid IV bisphosphonate therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

XIFAXAN

Products Affected

- Xifaxan TABS 550MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Small intestinal bacterial overgrowth syndrome (SIBO)
Exclusion Criteria	N/A
Required Medical Information	For irritable bowel syndrome with diarrhea (IBS-D): 1) The member has not previously received treatment with the requested drug, OR 2) The member has previously received treatment with the requested drug, AND a) the member is experiencing a recurrence of symptoms, AND b) the member has not already received an initial 14-day course of treatment and two additional 14-day courses of treatment with the requested drug. For small intestinal bacterial overgrowth (SIBO): 1) the member is experiencing a recurrence after completing a successful course of treatment with the requested drug OR 2) diagnosis has been confirmed by one of the following: a) quantitative culture of upper gut aspirate, b) breath testing (e.g., lactulose hydrogen or glucose hydrogen breath test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Reduction in risk of overt HE recurrence: 6 months, IBS-D and SIBO: 14 days
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XIFAXAN 200MG

Products Affected

- Xifaxan TABS 200MG

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Must have a documented diagnosis of Traveler's diarrhea caused by Noninvasive strains of E coli.
Age Restrictions	Traveler's diarrhea: 12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XOLAIR

Products Affected

- Xolair

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

XOLREMDI

Products Affected

- Xolremdi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For WHIM syndrome (warts, hypogammaglobulinemia, infections and myelokathexis), initial: 1) Diagnosis has been confirmed via testing to detect mutations in the CXCR4 gene AND 2) The member exhibits at least one clinical manifestation of the disease (such as warts, hypogammaglobulinemia, infections, myelokathexis) AND 3) The member has a confirmed low neutrophil count based on the reference laboratory range or current practice guidelines. For WHIM syndrome, continuation: The member has demonstrated a positive response to therapy.
Age Restrictions	12 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

XOSPATA

Products Affected

- Xospata

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FLT3 rearrangement, acute myeloid leukemia (AML) post allogeneic hematopoietic cell transplantation (HCT), in remission.
Exclusion Criteria	N/A
Required Medical Information	For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FMS-like tyrosine kinase 3 (FLT3) rearrangement: the disease is in chronic or blast phase. For AML with FLT3 mutation: The requested drug will be used for one of the following: a) relapsed or refractory disease, b) induction therapy, c) post-induction therapy following response to induction therapy with the requested drug, d) consolidation therapy, e) maintenance therapy in members who are in remission after allogeneic hematopoietic cell transplantation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Histologic transformation of indolent lymphomas to diffuse large B-cell lymphoma, Human Immunodeficiency Virus (HIV)-related B-cell lymphoma, high-grade B-cell lymphoma, post-transplant lymphoproliferative disorders
Exclusion Criteria	N/A
Required Medical Information	For multiple myeloma: Member must have been treated with at least one prior therapy. For B-cell lymphomas: Member must have been treated with at least two lines of systemic therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of castration-resistant prostate cancer or metastatic castration-sensitive prostate cancer: The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

YONSA

Products Affected

- Yonsa

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

YORVIPATH

Products Affected

- Yorvipath

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	Acute post-surgical hypoparathyroidism (within 6 months of surgery) and expected recovery from hypoparathyroidism.
Required Medical Information	For hypoparathyroidism, initial: prior to initiation, the member's albumin-corrected serum calcium has been or will be confirmed to be greater than or equal to 7.8 mg/dL. For hypoparathyroidism, continuation: the member is experiencing benefit from therapy (for example, maintenance or normalization of serum calcium levels compared to baseline).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Continuation: Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZALEPLON

Products Affected

- Zaleplon

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For insomnia: 1) The member has had an inadequate treatment response, contraindication, or intolerance to non-HRM (high-risk medication) alternative drug doxepin (3 mg or 6 mg) and ramelteon AND 2) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 3) If the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZARXIO

Products Affected

- Zarxio

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Neutropenia in myelodysplastic syndromes (MDS), agranulocytosis, neutropenia in aplastic anemia, human immunodeficiency virus (HIV)-related neutropenia
Exclusion Criteria	N/A
Required Medical Information	If receiving chemotherapy, the requested drug will be administered at least 24 hours after chemotherapy. For prophylaxis or treatment of myelosuppressive chemotherapy-induced febrile neutropenia (FN) member must meet both of the following: 1) Member has a solid tumor or non-myeloid cancer, AND 2) Member has received, is currently receiving, or will be receiving treatment with myelosuppressive anti-cancer therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZEJULA

Products Affected

- Zejula TABS

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

ZELBORAF

Products Affected

- Zelboraf

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Non-small cell lung cancer, hairy cell leukemia, central nervous system cancer (i.e., glioma, glioblastoma, pediatric diffuse high-grade glioma), adjuvant systemic therapy for cutaneous melanoma, Langerhans cell histiocytosis.
Exclusion Criteria	N/A
Required Medical Information	For central nervous system (CNS) cancer (i.e., glioma, astrocytoma, glioblastoma, pediatric diffuse high-grade glioma): 1) The tumor is positive for BRAF V600E mutation, AND 2) The requested drug will be used in combination with cobimetinib OR the requested drug is being used for the treatment of pediatric diffuse high-grade glioma. For melanoma: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), AND 2) the requested drug will be used as a single agent, or in combination with cobimetinib, AND 3) The requested drug will be used for either of the following: a) unresectable, limited resectable, or metastatic disease, or b) adjuvant systemic therapy. For Erdheim-Chester Disease and Langerhans Cell Histiocytosis: Tumor is positive for BRAF V600 mutation. For non-small cell lung cancer: 1) The tumor is positive for the BRAF V600E mutation, AND 2) The member has recurrent, advanced, or metastatic disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZELSUVMI

Products Affected

- Zelsuvmi

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	1 year of age or older
Prescriber Restrictions	N/A
Coverage Duration	12 weeks
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZENBEXUS

Products Affected

- Zenbexus

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

ZEPOSIA

Products Affected

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

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

ZILBRYSQ

Products Affected

- Zilbrysq

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

ZIRABEV

Products Affected

- Zirabev

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Ampullary adenocarcinoma, appendiceal adenocarcinoma, central nervous system (CNS) cancers (including pediatric diffuse high-grade gliomas), pleural mesothelioma, peritoneal mesothelioma, pericardial mesothelioma, tunica vaginalis testis mesothelioma, soft tissue sarcomas, uterine neoplasms, endometrial carcinoma, vulvar cancers, vaginal cancer, small bowel adenocarcinoma, and ophthalmic-related disorders: diabetic macular edema, neovascular (wet) age-related macular degeneration including polypoidal choroidopathy and retinal angiomatous proliferation subtypes, macular edema following retinal vein occlusion, proliferative diabetic retinopathy, choroidal neovascularization, neovascular glaucoma and retinopathy of prematurity.
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZOLINZA

Products Affected

- Zolinza

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Mycosis fungoides (MF)/Sezary syndrome (SS)
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES NOT require use of a prerequisite Part D drug.

ZOLPIDEM

Products Affected

- Zolpidem Tartrate TABS
- Zolpidem Tartrate Er

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For insomnia: 1) The member has had an inadequate treatment response, contraindication, or intolerance to non-HRM (high-risk medication) alternative drug doxepin (3 mg or 6 mg) and ramelteon AND 2) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this member AND 3) If the member is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the member.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization only applies to members 65 years of age or older. Prior Authorization applies to greater than cumulative 90 days of therapy per year.
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZONISADE

Products Affected

- Zonisade

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For adjunctive treatment of partial-onset seizures (i.e., focal-onset seizures): 1) The member has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND the member has experienced an inadequate treatment response, intolerance, or has a contraindication to any of the following: Aptiom, Xcopri, Spritam OR 2) The member has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules).
Age Restrictions	16 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
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 FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	For the treatment of postpartum depression (PPD): diagnosis was confirmed using standardized rating scales that reliably measure depressive symptoms (e.g., Hamilton Depression Rating Scale [HDRS], Edinburgh Postnatal Depression Scale [EPDS], Member Health Questionnaire 9 [PHQ9], Montgomery-Asberg Depression Rating Scale [MADRS], Beck's Depression Inventory [BDI], etc.).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 month
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 FDA-approved Indications, Some Medically accepted Indications.
Off-Label Uses	Small lymphocytic lymphoma (SLL)
Exclusion Criteria	N/A
Required Medical Information	For chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL): the requested drug is used as second-line or subsequent therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

ZYKADIA

Products Affected

- Zykadia TABS

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

ZYMFENTRA

Products Affected

- Zymfentra 2-pen
- Zymfentra 2-syringe

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

ZYPREXA RELPREVV

Products Affected

- Zyprexa Relprevv

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off-Label Uses	N/A
Exclusion Criteria	N/A
Required Medical Information	Tolerability with oral olanzapine has been established.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Prerequisite Therapy Required	Criteria DOES require use of a prerequisite Part D drug.

PART B VERSUS PART D

Products Affected

- Acetylcysteine SOLN
- Albuterol Sulfate NEBU 0.083%, 0.63MG/3ML, 1.25MG/3ML, 2.5MG/0.5ML
- Alyglo
- 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
- Azasan
- Azathioprine TABS
- Bivigam INJ 10%, 5GM/50ML
- Budesonide SUSP
- Clinimix 4.25%/dextrose 10%
- Clinimix 4.25%/dextrose 5%
- Clinimix 5%/dextrose 15%
- Clinimix 5%/dextrose 20%
- Clinimix 6/5
- Clinimix 8/10
- Clinimix 8/14
- Clinisol Sf 15%
- Cromolyn Sodium NEBU
- Cyclophosphamide CAPS
- Cyclophosphamide TABS 50MG
- Cyclosporine CAPS
- Cyclosporine Modified
- Dronabinol CAPS
- Emend SUSR
- Engerix-b
- Envarsus Xr
- Everolimus TABS 0.25MG, 0.5MG, 0.75MG, 1MG
- Flebogamma Dif INJ 10GM/200ML, 20GM/400ML, 5GM/100ML
- Formoterol Fumarate NEBU
- Gammagard Liquid
- Gammagard Liquid Erc

- Gammagard S/d Iga Less Than 1mcg/ml
- Gammaked INJ 10GM/100ML, 1GM/10ML, 20GM/200ML, 5GM/50ML
- Gammaplex INJ 10GM/100ML, 10GM/200ML, 20GM/200ML, 20GM/400ML, 5GM/100ML, 5GM/50ML
- Gamunex-c
- Gengraf CAPS 100MG, 25MG
- Granisetron Hydrochloride TABS
- Heplisav-b
- Imovax Rabies (h.d.c.v.)
- Intralipid INJ 20GM/100ML, 30GM/100ML
- Ipratropium Bromide INHALATION SOLN 0.02%
- Ipratropium Bromide/albuterol Sulfate
- Levalbuterol NEBU
- Levalbuterol Hcl NEBU 0.31MG/3ML, 1.25MG/3ML
- Levalbuterol Hydrochloride NEBU 0.63MG/3ML
- Mycophenolate Mofetil CAPS
- Mycophenolate Mofetil SUSR
- Mycophenolate Mofetil TABS
- Mycophenolic Acid Dr
- Myhibbin
- Nutrilipid
- Octagam INJ 10GM/100ML, 10GM/200ML, 1GM/20ML, 2.5GM/50ML, 20GM/200ML, 2GM/20ML, 30GM/300ML, 5GM/100ML, 5GM/50ML
- Ohtuvayre
- Ondansetron Hcl SOLN 4MG/5ML
- Ondansetron Hydrochloride TABS
- Ondansetron Odt
- Panzyga
- Pentamidine Isethionate INHALATION SOLR
- Plenamine INJ 147.4MEQ/L; 2.17GM/100ML; 1.47GM/100ML; 434MG/100ML; 749MG/100ML; 1.04GM/100ML; 894MG/100ML; 749MG/100ML; 1.04GM/100ML; 1.18GM/100ML; 749MG/100ML; 1.04GM/100ML; 894MG/100ML; 592MG/100ML; 749MG/100ML; 250MG/100ML; 39MG/100ML; 960MG/100ML
- Premasol INJ 52MEQ/L; 1760MG/100ML; 880MG/100ML; 34MEQ/L; 1760MG/100ML; 372MG/100ML; 406MG/100ML; 526MG/100ML; 492MG/100ML; 492MG/100ML; 526MG/100ML; 356MG/100ML; 356MG/100ML; 390MG/100ML; 34MG/100ML; 152MG/100ML
- Privigen
- Prograf PACK
- Prosol
- Pulmozyme SOLN 2.5MG/2.5ML
- Qivigy
- Rabavert
- Recombivax Hb
- Sirolimus SOLN
- Sirolimus TABS
- Tacrolimus CAPS
- Tacrolimus Er
- Tobramycin NEBU
- 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
- Yupelri

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.