ABIRATERONE

Products Affected

- abiraterone oral tablet 250 mg
- Zytiga oral tablet 500 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Node-positive (N1), non-metastatic (M0) prostate cancer

ACITRETIN

Products Affected

- acitretin

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Prevention of non-melanoma skin cancers in high risk individuals, Lichen planus, Keratosis follicularis (Darier Disease).

ACTIMMUNE

Products Affected

- Actimmune

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Mycosis fungoides, Sezary syndrome, atopic dermatitis.

ADEMPAS

Products Affected

- Adempas

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For pulmonary arterial hypertension (PAH) (WHO Group 1): PAH was confirmed by right heart catheterization. For chronic thromboembolic pulmonary hypertension (CTEPH) (WHO Group 4): Patient has persistent or recurrent CTEPH after pulmonary endarterectomy (PEA), OR patient has inoperable CTEPH with the diagnosis confirmed by right heart catheterization AND by computed tomography (CT), magnetic resonance imaging (MRI), or pulmonary angiography. For new starts only (excluding recurrent/persistent CTEPH after PEA): 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, AND 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

AFINITOR

Products Affected

- Afinitor Disperz oral tablet for suspension 2 mg, 3 mg, 5 mg
- Afinitor oral tablet 10 mg
- everolimus (antineoplastic)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Breast Cancer-HER2 status, hormone receptor (HR) status.
Age Restrictions	Relapsed or refractory classical Hodgkin lymphoma-18 years and older
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Breast CA-approve if the patient meets ALL the following criteria(A, B, C, D, E, and F):A)patient has recurrent or Stage IV, hormone receptor positive (HR+) [i.e.,estrogen receptor positive (ER+) and/or progesterone receptor positive (PR+)] disease AND B)patient has human epidermal growth factor receptor 2 (HER2)-negative breast cancer AND C)patient has tried at least one prior endocrine therapy (e.g., anastrozole, letrozole, or tamoxifen) AND D)patient meets ONE of the following conditions (i or ii): i.patient is a postmenopausal female or a male OR ii. patient is premenopausal or perimenopausal AND is receiving ovarian suppression/ablation with a gonadotropin-releasing hormone (GnRH) agonist (e.g., Lupron [leuprolide], Trelstar [triptorelin], Zoladex (goserelin]), or has had surgical bilateral oophorectomy or ovarian irradiation AND E) The patient meets ONE of the following conditions (i or ii): i. If patient is a male AND if everolimus will be used in combination with exemestane, the patient is receiving a gonadotropin-releasing hormone (GnRH) agonist (e.g., Lupron [leuprolide], Trelstar [triptorelin], Zoladex (goserelin]) OR ii. everolimus will be used in combination with exemestane, Faslodex (fulvestrant intramuscular), or tamoxifen AND F) The patient has not had disease progression while on everolimus. Renal Cell Carcinoma (Clear Cell or Non-clear cell histology)-approve if the patient has relapsed or Stage IV disease and if using for clear cell disease, the patient has tried one prior systemic therapy (e.g., Inlyta, Votrient, Sutent, Cabometyx,

PA Criteria	Criteria Details
	Nexavar).Tuberous sclerosis complex (TSC) Associated subependymal giant cell astrocytoma (SEGA)-approve if the patient requires therapeutic intervention but cannot be curatively resected. Thymomas and Thymic Carcinomas-approve if the patient has tried one prior chemotherapy (e.g., cisplatin plus doxorubicin, cisplatin plus etoposide, carboplatin plus paclitaxel). TSC associated renal angiomyolipoma -approve. WM/LPL - approve if patient has progressive or relapsed disease or if the patient has not responded to ONE primary therapy (e.g., Velcade with dexamethasone with or without Rituxan, Treanda with Rituxan, Rituxan with cyclophosphamide and dexamethasone, Treanda, Velcade with or without Rituxan, Velcade with dexamethasone, Kyprolis with Rituxan and dexamethasone, Imbruvica Rituxan). Differentiated (i.e. papillary, follicular, and Hurthle cell) Thyroid Carcinoma-approve if the patient is refractory to radioactive iodine therapy. Endometrial Carcinoma-approve if everolimus will be used in combination with letrozole and the patient has recurrent, metastatic or high-risk disease. GIST-approve if the patient has tried TWO of the following drugs: Sutent, Stivarga, or imatinib AND there is confirmation that everolimus will be used in combination with one of these drugs (Sutent, Stivarga, or imatinib) in the treatment of GIST.Tuberous sclerosis complex (TSC)-associated partial-onset seizures-approve.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Advanced, unresectable or metastatic neuroendocrine tumors of the thymus (Carcinoid tumors). Perivascular Epitheloid Cell Tumors (PEComa), Recurrent Angiomyolipoma, Lymphangioleiomyomatosis, relapsed or refractory classical Hodgkin lymphoma, Waldenstrom's Macroglobulinemia/Lymphoplasmacytic Lymphoma (WM/LPL), Thymomas and Thymic carcinomas, Differentiated (i.e. papillary, follicular, and Hurthle cell) Thyroid Carcinoma, Endometrial Carcinoma, Gastrointestinal Stromal Tumors (GIST) and Recurrent or progressive Meningioma, men with breast cancer

AIMOVIG

Products Affected

- Aimovig Autoinjector

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The patient received at least 3 months of treatment with the requested drug, and the patient had a reduction in migraine days per month from baseline, OR 2) The patient experienced an inadequate treatment response with a 4-week trial of any of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants, OR 3) The patient experienced an intolerance or has a contraindication that would prohibit a 4-week trial of any of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial 3 months, Reauthorization Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ALECENSA

Products Affected

- Alecensa

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Recurrent anaplastic lymphoma kinase (ALK)-positive non-small cell lung cancer, brain metastases from ALK-positive non-small cell lung cancer.

ALOSETRON

Products Affected

- alosetron

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for a biological female or a person that self-identifies as a female with a diagnosis of severe diarrhea-predominant irritable bowel syndrome (IBS) AND 2) Chronic IBS symptoms lasting at least 6 months AND 3) Gastrointestinal tract abnormalities have been ruled out AND 4) Inadequate response to conventional therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ALPHA1-PROTEINASE INHIBITOR

Products Affected

- Aralast NP intravenous recon soln 1,000 mg
- Prolastin-C intravenous recon soln
- Zemaira

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For alpha1-proteinase inhibitor deficiency: Patient must have 1) clinically evident emphysema, 2) pretreatment serum alpha1-proteinase inhibitor level less than 11 micromol/L (80 mg/dL by radial immunodiffusion or 50 mg/dL by nephelometry), and 3) pretreatment post-bronchodilation forced expiratory volume in 1 second (FEV1) greater than or equal to 25 percent and less than or equal to 80 percent of predicted.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ALUNBRIG

Products Affected

- Alunbrig

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For metastatic or recurrent ALK-positive NSCLC: patient must have progressed on or experienced intolerance to crizotinib. For brain metastases from NSCLC: disease is ALK-positive. Treatment of adult patients with tyrosine kinase inhibitor-na+ve advanced anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC) isn't excluded from coverage.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
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 NSCLC.

ANADROL

Products Affected

- Anadrol-50

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Cachexia associated with AIDS (HIV-wasting)

APOKYN

Products Affected

- APOKYN

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ARCALYST

Products Affected

- Arcalyst

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For prevention of gout flares in members initiating or continuing urate-lowering therapy (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 non-steroidal anti-inflammatory drugs and colchicine, AND 3) concurrent use with urate-lowering therapy. For prevention of gout flares in members initiating or continuing urate-lowering therapy (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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	For prevention of gout flares: 4 months. Other: Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Prevention of gout flares in patients initiating or continuing urate-lowering therapy.

ARMODAFINIL

Products Affected

- armodafinil oral tablet 150 mg, 200 mg, 250 mg, 50 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Diagnosis is narcolepsy confirmed by sleep lab evaluation OR 2) Diagnosis is Shift Work Disorder (SWD) OR 3) Diagnosis is obstructive sleep apnea (OSA) confirmed by polysomnography
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ATYPICAL ANTIPSYCHOTICS

Products Affected

- Fanapt oral tablet
- Fanapt oral tablets,dose pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following: lurasidone, aripiprazole, olanzapine, paliperidone, quetiapine, risperidone, or ziprasidone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

AURYXIA

Products Affected

- Auryxia

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Coverage will be denied if request is for an indication excluded from Part D.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

AUSTEDO

Products Affected

- Austedo oral tablet 12 mg, 6 mg, 9 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

AYVAKIT

Products Affected

- Ayvakit

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BALVERSA

Products Affected

- Balversa

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BANZEL

Products Affected

- Banzel

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BENLYSTA

Products Affected

- Benlysta subcutaneous

PA Criteria	Criteria Details
Exclusion Criteria	Severe active lupus nephritis. Severe active central nervous system lupus.
Required Medical Information	For systemic lupus erythematosus (SLE): 1) Patient is currently receiving standard therapy (e.g., corticosteroids, azathioprine, leflunomide, methotrexate, mycophenolate mofetil, hydroxychloroquine, non-steroidal anti-inflammatory drugs) for SLE OR 2) patient is not currently receiving standard therapy for SLE because patient tried and had an inadequate response or intolerance to standard therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BERINERT

Products Affected

- Berinert intravenous kit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For hereditary angioedema (HAE): patient has hereditary angioedema with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has hereditary angioedema with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, EITHER 1) Patient tested positive for an F12, angiopoietin-1, or plasminogen gene mutation OR 2) Patient has a family history of angioedema and the angioedema was refractory to a trial of antihistamine for at least one month.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BETASERON

Products Affected

- Betaseron subcutaneous kit

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BEXAROTENE

Products Affected

- bexarotene
- Targretin topical

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Mycosis fungoides, Sezary syndrome (capsules only), primary cutaneous CD30-positive T-cell lymphoproliferative disorder types: primary cutaneous anaplastic large cell lymphoma (capsules only) and lymphomatoid papulosis (capsules only), chronic or smoldering adult T-cell leukemia/lymphoma (gel only), primary cutaneous B-cell lymphoma types: primary cutaneous marginal zone lymphoma (gel only) and primary cutaneous follicle center lymphoma (gel only).

BOSENTAN

Products Affected

- bosentan oral tablet 125 mg, 62.5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For pulmonary arterial hypertension (PAH) (WHO Group 1): Diagnosis was confirmed by right heart catheterization. For PAH new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BOSULIF

Products Affected

- Bosulif

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic myelogenous leukemia (CML) or acute lymphoblastic leukemia (ALL): Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML: 1) Patient received a hematopoietic stem cell transplant, OR 2) Patient has accelerated or blast phase CML, OR 3) Patient has chronic phase CML (includes newly diagnosed) and meets one of the following conditions: a) high or intermediate risk for disease progression, or b) low risk for disease progression and has experienced resistance, intolerance or toxicity to imatinib or an alternative tyrosine kinase inhibitor. If patient experienced resistance to imatinib or an alternative tyrosine kinase inhibitor for CML, patient is negative for T315I mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Relapsed or refractory Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL).

BRAFTOVI

Products Affected

- Braftovi oral capsule 75 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BRIVIACT

Products Affected

- Briviact oral

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	4 years of age or older (tablets and oral solution).
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BRUKINSA

Products Affected

- Brukinsa

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

BUPRENORPHINE

Products Affected

- buprenorphine HCl sublingual

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for the treatment of opioid dependence AND 2) The patient is pregnant or breastfeeding, and the requested drug is being prescribed for induction therapy and/or subsequent maintenance therapy for opioid dependence treatment OR 3) The requested drug is being prescribed for induction therapy for transition from opioid use to opioid dependence treatment OR 4) The requested drug is being prescribed for maintenance therapy for opioid dependence treatment in a patient who is intolerant to naloxone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CABOMETYX

Products Affected

- Cabometyx

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma: The disease is relapsed, unresectable, or metastatic. For non-small cell lung cancer: The disease is rearranged during transfection (RET) positive. Treatment of patients with hepatocellular carcinoma who have been previously treated with sorafenib are not excluded from coverage.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Non-small cell lung cancer

CALCIPOTRIENE

Products Affected

- calcipotriene scalp
- calcipotriene topical cream
- calcipotriene topical ointment
- Enstilar

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for the treatment of psoriasis AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to a generic topical steroid.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CALQUENCE

Products Affected

- Calquence

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CAPRELSA

Products Affected

- Caprelsa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For NSCLC: the requested medication is used for NSCLC with RET gene rearrangements.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Non-small cell lung cancer (NSCLC), differentiated thyroid carcinoma: papillary, follicular, and Hurthle cell.

CARBAGLU

Products Affected

- Carbaglu

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For N-acetylglutamate synthase (NAGS) deficiency: Diagnosis of NAGS deficiency was confirmed by enzymatic or genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CAYSTON

Products Affected

- Cayston

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For treatment of respiratory symptoms in cystic fibrosis patients: 1) Pseudomonas aeruginosa is present in the patient's airway cultures OR 2) The patient 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CERDELGA

Products Affected

- Cerdelga

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Gaucher disease was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing. The patient's CYP2D6 metabolizer status has been established using an FDA-cleared test. The patient 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CLOBAZAM

Products Affected

- clobazam

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CLOMIPRAMINE

Products Affected

- clomipramine

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for one of the following: the treatment of Obsessive-Compulsive Disorder (OCD) or Panic Disorder AND 2) The patient has experienced an inadequate treatment response, intolerance, or contraindication to one of the following: a generic selective serotonin reuptake inhibitor (SSRI), a generic serotonin and norepinephrine reuptake inhibitor (SNRI), mirtazapine OR 3) The requested drug is being prescribed for the treatment of Depression AND 4) The patient has experienced an inadequate treatment response, intolerance, or contraindication to one of the following: a generic selective serotonin reuptake inhibitor (SSRI), a generic serotonin and norepinephrine reuptake inhibitor (SNRI) , mirtazapine, bupropion
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Depression, Panic Disorder

CLORAZEPATE

Products Affected

- clorazepate dipotassium oral tablet 15 mg, 3.75 mg, 7.5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) For the management of anxiety disorders, the requested drug is being used with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the antidepressant becomes effective for the symptoms of anxiety OR the patient has experienced an inadequate treatment response, intolerance or contraindication to a selective serotonin reuptake inhibitor (SSRI) or a serotonin-norepinephrine reuptake inhibitor (SNRI) OR 2) For adjunctive therapy in the management of partial seizures OR 3) Symptomatic relief in acute alcohol withdrawal OR 4) For the short-term relief of the symptoms of anxiety AND 5) The benefit of therapy with the prescribed medication outweighs the potential risk in a patient 65 years of age or older.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Short-term relief anxiety-1 Month, Anxiety Disorders-4 Months, All other Diagnoses-Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 65 years of age or older. The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CLOZAPINE ODT

Products Affected

- clozapine oral tablet,disintegrating 100 mg, 12.5 mg, 25 mg
- clozapine oral tablet,disintegrating 150 mg, 200 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

COMETRIQ

Products Affected

- Cometriq

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For NSCLC: The requested medication is used for NSCLC with RET gene rearrangements.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Non-small cell lung cancer (NSCLC), differentiated thyroid carcinoma: papillary, follicular, and Hurthle cell

COPIKTRA

Products Affected

- Copiktra

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

COSENTYX

Products Affected

- Cosentyx (2 Syringes)
- Cosentyx Pen (2 Pens)

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent Use with other Biologics or Targeted Synthetic Disease-Modifying Antirheumatic Drugs (DMARDs)
Required Medical Information	Diagnosis and previous medications use
Age Restrictions	PP/AS/PSA initial - 18 years of age and older
Prescriber Restrictions	PP initial - prescribed by or in consultation with a dermatologist or rheumatologist. AS initial- by or in consultation with rheumatologist, PsA initial- by or in consultation with rheumatologist or dermatologist.
Coverage Duration	PP/AS/nr-axSpA - initial tx 3 mos, PsA-initial tx 3 mos, cont tx 3 years
Other Criteria	PP initial-approve if the patient has tried TWO of the following: Enbrel, Humira, Skyrizi, Stelara SC, Otezla or Taltz. PsA initial-approve if the patient has tried TWO of the following: Enbrel, Humira, Stelara SC, Otezla, Orencia, Xeljanz/XR or Taltz. (Note: if the patient has not tried TWO of these drugs listed, previous trial(s) with the following drugs can count towards meeting the try TWO requirement: Cimzia, an infliximab product, golimumab SC/IV). AS-approve if the patient has tried TWO of the following-Enbrel, Humira or Taltz. PP/AS/PsA cont - patient must have responded, as determined by the prescriber.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

COTELLIC

Products Affected

- Cotellic

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For melanoma (including brain metastases): 1) The disease is unresectable or metastatic, 2) The disease is positive for the BRAF V600E or V600K mutation, AND 3) The requested medication will be used in combination with vemurafenib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Brain metastases from melanoma

CRESEMBA (ORAL)

Products Affected

- Cresemba oral

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Candidiasis of the esophagus - HIV infection, sepsis

CYSTAGON

Products Affected

- Cystagon

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For nephropathic cystinosis: Diagnosis was confirmed by the presence of increased cystine concentration in leukocytes or by genetic testing.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

CYSTARAN

Products Affected

- Cystaran

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For treatment of corneal cystine crystal accumulation in patients with cystinosis: 1) Diagnosis of cystinosis was confirmed by the presence of increased cystine concentration in leukocytes or by genetic testing, and 2) The patient has corneal cystine crystal accumulation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DALFAMPRIDINE

Products Affected

- dalfampridine

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For multiple sclerosis new starts: Prior to initiating therapy, patient demonstrates sustained walking impairment. For multiple sclerosis continuation of therapy: Patient must have experienced an improvement in walking speed or other objective measure of walking ability since starting the requested medication.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DAURISMO

Products Affected

- Daurismo

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DEFERASIROX

Products Affected

- deferasirox oral granules in packet
- deferasirox oral tablet 360 mg, 90 mg
- deferasirox oral tablet, dispersible

PA Criteria	Criteria Details
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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DEMSER

Products Affected

- metyrosine

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DESVENLAFAXINE

Products Affected

- desvenlafaxine succinate

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Patient experienced an inadequate treatment response, intolerance, or contraindication to any of the following: a generic serotonin and norepinephrine reuptake inhibitor (SNRI), a generic selective serotonin reuptake inhibitor (SSRI), mirtazapine, bupropion
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DHE NASAL

Products Affected

- dihydroergotamine nasal

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The patient experienced an inadequate treatment response, intolerance, or contraindication to one triptan 5-HT1 receptor agonist
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DIAZEPAM

Products Affected

- diazepam oral concentrate
- diazepam oral solution 5 mg/5 mL (1 mg/mL)
- diazepam oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) For the management of anxiety disorders, the requested drug is being used with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the antidepressant becomes effective for the symptoms of anxiety OR the patient has experienced an inadequate treatment response, intolerance or contraindication to a selective serotonin reuptake inhibitor (SSRI) or a serotonin-norepinephrine reuptake inhibitor (SNRI) OR 2) For symptomatic relief in acute alcohol withdrawal OR 3) For use as an adjunct for the relief of skeletal muscle spasms OR 4) For adjunctive therapy in the treatment of convulsive disorders OR 5) For the short-term relief of the symptoms of anxiety AND 6) The benefit of therapy with the prescribed medication outweighs the potential risk in a patient 65 years of age or older.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Short-term relief anxiety-1 Month, Anxiety Disorders-4 Months, All other Diagnoses-Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 65 years of age or older. The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

DRIZALMA

Products Affected

- Drizalma Sprinkle oral capsule, delayed rel sprinkle 20 mg, 30 mg, 40 mg, 60 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The patient has tried duloxetine capsules or the patient is unable to take duloxetine capsules for any reason (e.g., difficulty swallowing capsules, requires nasogastric administration)
Age Restrictions	GAD - 7 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Cancer pain, chemotherapy-induced neuropathic pain

DUPIXENT

Products Affected

- Dupixent Pen
- Dupixent Syringe subcutaneous syringe 200 mg/1.14 mL, 300 mg/2 mL

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with Xolair or another Anti-interleukin (IL) Monoclonal Antibody.
Required Medical Information	Diagnosis, prescriber specialty, other medications tried and length of trials
Age Restrictions	AD-6 years and older, asthma-12 years of age and older. Chronic Rhinosinusitis-18 years of age and older
Prescriber Restrictions	Atopic Dermatitis-Prescribed by or in consultation with an allergist, immunologist or dermatologist, asthma-prescribed by or in consultation with an allergist, immunologist or pulmonologist. Rhinosinusitis-prescribed by or in consultation with an allergist, immunologist or otolaryngologist.
Coverage Duration	AD-Initial-4 months, Cont-1 year, asthma/Rhinosinusitis-initial-6 months, continuation 1 year
Other Criteria	Atopic Dermatitis-Initial-meets both a and b: a.has used at least one medium, medium-high, high, and/or super-high-potency prescription topical corticosteroid OR has atopic dermatitis affecting ONLY the face, eyes/eyelids, skin folds, and/or genitalia and has tried tacrolimus ointment AND b.Inadequate efficacy was demonstrated with these previously tried topical prescription therapies, according to the prescribing physician.Continuation-Approve if the pt has responded to Dupixent therapy as determined by the prescribing physician. Asthma-Initial-approve if pt meets the following criteria (i, ii, and iii):i.Pt meets ONE of the following criteria (a or b):a)has a blood eosinophil level of greater than or equal to 150 cells per microliter within the previous 6 weeks or within 6 weeks prior to treatment with any anti-interleukin (IL) therapy or Xolair OR b)has oral corticosteroid-dependent asthma, per the prescriber AND ii.has received combination therapy with BOTH of the following (a and b): a)An inhaled corticosteroid (ICS) AND b)At least one additional asthma controller/maintenance medication (NOTE:An exception to the requirement for a trial of one additional asthma controller/maintenance medication can be made if the patient has already received anti-IL-5 therapy or Xolair used concomitantly with an ICS. Use of a combination inhaler containing both an ICS and a LABA would fulfil the requirement

PA Criteria	Criteria Details
	for both criteria a and b) AND iii.asthma is uncontrolled or was uncontrolled prior to starting any anti-IL therapy or Xolair as defined by ONE of the following (a, b, c, d or e): a)experienced two or more asthma exacerbations requiring treatment with systemic corticosteroids in the previous year OR b)experienced one or more asthma exacerbation requiring hospitalization or an Emergency Department visit in the previous year OR c)has a forced expiratory volume in 1 second (FEV1) less than 80% predicted OR d)has an FEV1/forced vital capacity (FVC) less than 0.80 OR e)The patient's asthma worsens upon tapering of oral corticosteroid therapy. Continuation-Approve if meets the following criteria (i and ii): i.continues to receive therapy with one inhaled corticosteroid (ICS) or one ICS-containing combination inhaler AND ii.has responded to Dupixent therapy as determined by the prescribing physician. Chronic rhinosinusitis with Nasal Polyposis-Initial-pt is currently receiving therapy with an intranasal corticosteroid AND is experiencing significant rhinosinusitis symptoms such as nasal obstruction, rhinorrhea, or reduction/loss of smell according to the prescriber AND meets ONE of the following (a or b): a)has received treatment with a systemic corticosteroid within the previous 2 years or has a contraindication to systemic corticosteroid therapy OR b)has had prior surgery for nasal polyps. Continuation-approve if the pt continues to receive therapy with an intranasal corticosteroid AND pt has responded to Dupixent therapy as determined by the prescriber.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

EMGALITY

Products Affected

- Emgality Pen
- Emgality Syringe subcutaneous syringe 120 mg/mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for the preventive treatment of migraine in an adult patient AND 2) The patient received at least 3 months of treatment with the requested drug, and the patient had a reduction in migraine days per month from baseline, OR 3) The patient experienced an inadequate treatment response with a 4-week trial of any of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants, OR 4) The patient experienced an intolerance or has a contraindication that would prohibit a 4-week trial of any of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants OR 1) The requested drug is being prescribed for the treatment of episodic cluster headaches in an adult patient AND 2) The patient received the requested drug for at least 3 weeks of treatment and had a reduction in weekly cluster headache attack frequency from baseline OR 3) The patient experienced an inadequate treatment response, intolerance, or contraindication to a triptan medication (i.e., 5-HT1 receptor agonist).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial 3 months, Reauthorization Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

EMSAM

Products Affected

- Emsam

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Patient experienced an inadequate treatment response, intolerance, or contraindication to any of the following antidepressants: bupropion, trazodone, mirtazapine, serotonin norepinephrine reuptake inhibitors (SNRIs), selective serotonin reuptake inhibitors (SSRIs), tricyclic or tetracyclic antidepressants OR 2) Patient is unable to swallow oral formulations.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ENBREL

Products Affected

- Enbrel
- Enbrel Mini
- Enbrel SureClick

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with biologic therapy or targeted synthetic DMARD
Required Medical Information	Diagnosis, concurrent medications, previous therapies tried.
Age Restrictions	PP-4 years and older (initial therapy)
Prescriber Restrictions	RA/AS/JIA/JRA,prescribed by or in consult w/ rheum. PsA, prescribed by or in consultation w/ rheumatologist or dermatologist.PP, prescribed by or in consult w/ dermatologist.GVHD,prescribed by or in consult w/ oncologist,hematologist,or physician affiliated w/ transplant center.Behcet's disease,prescribed by or in consult w/ rheumatologist,dermatologist,ophthalmologist,gastroenterologist,or neurologist. Uveitis, prescribed by or in consultation with an ophthalmologist.
Coverage Duration	FDA dx-3 mo init, 3 yrs cont, Behcet's/uveitis init-3 mo, cont-12 mo.GVHD-1 mo init/3 mo cont.
Other Criteria	RA initial, patient has tried one conventional synthetic DMARD for at least 3 months (note: patients who have already had a 3-month trial of a biologic for RA are not required to step back and try a conventional synthetic DMARD). JIA/JRA, approve if the pt has aggressive disease, as determined by the prescriber, or the pt has tried one other agent for this condition (eg, MTX, sulfasalazine, leflunomide, NSAID, biologic DMARD or the pt will be started on Enbrel concurrently with MTX, sulfasalazine, or leflunomide or the pt has an absolute contraindication to MTX (eg, pregnancy, breast feeding, alcoholic liver disease, immunodeficiency syndrome, blood dyscrasias), sulfasalazine, or leflunomide.Plaque psoriasis (PP) initial approve if the patient meets one of the following conditions: 1) patient has tried at least one traditional systemic agent for at least 3 months for plaque psoriasis, unless intolerant (eg, MTX, cyclosporine, Soriatane, oral methoxsalen plus PUVA, (note: pts who have already tried a biologic for psoriasis are not required to step back and try a traditional agent first) OR 2) the patient has a contraindication to one oral agent for psoriasis such as MTX. GVHD. Tried or currently is receiving with etanercept 1

PA Criteria	Criteria Details
	conventional GVHD tx (high-dose systemic corticosteroid, CSA, tacrolimus, MM, thalidomide, antithymocyte globulin, etc.). Behcet's. Has tried at least 1 conventional tx (eg, systemic corticosteroid, immunosuppressant, interferon alfa, MM, etc) or adalimumab or infliximab. Uveitis-tried one of the following: periocular, intraocular, or systemic corticosteroid, immunosuppressives, Humira or an infliximab product.RA/AS/JIA/PP/PsA Cont - must have a response to tx according to the prescriber. Behcet's, GVHD, Uveitis Cont-if the patient has had a response to tx according to the prescriber. Clinical criteria incorporated into the Enbrel 25 mg quantity limit edit, approve additional quantity (to allow for 50 mg twice weekly dosing) if one of the following is met: 1) Patient has plaque psoriasis, OR 2) Patient has RA/JIA/PsA/AS and is started and stabilized on 50 mg twice weekly dosing, OR 3) Patient has RA and the dose is being increased to 50 mg twice weekly and patient has taken MTX in combination with Enbrel 50 mg once weekly for at least 2 months, unless MTX is contraindicated or intolerant, OR 4) Patient has JIA/PsA/AS and the dose is being increased to 50 mg twice weekly after taking 50 mg once weekly for at least 2 months.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Graft versus host disease (GVHD), Behcet's disease, Uveitis

ENDARI

Products Affected

- Endari

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	5 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

EPCLUSA

Products Affected

- Epclusa oral tablet 400-100 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic 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, liver and kidney transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current AASLD 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

EPIDIOLEX

Products Affected

- Epidiolex

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

EPO

Products Affected

- Procrit injection solution 10,000 unit/mL, 2,000 unit/mL, 20,000 unit/mL, 3,000 unit/mL, 4,000 unit/mL, 40,000 unit/mL

PA Criteria	Criteria Details
Exclusion Criteria	Patients receiving chemotherapy with curative intent. Patients with myeloid cancer.
Required Medical Information	For all uses except surgery: Pretreatment (no erythropoietin treatment in previous month) hemoglobin (Hgb) is less than 10 g/dL (less than 9 g/dL for anemia in congested heart failure only). Additional requirements for primary MF, post-polycythemia vera MF, post-essential thrombocythemia MF: 1) Patient has symptomatic anemia. 2) For initial therapy, pretreatment serum erythropoietin level is less than 500mU/mL. For surgery: 1) Patient is scheduled for elective, noncardiac, nonvascular surgery. 2) Pretreatment Hgb is greater than 10 but not more than 13 g/dL.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	16 weeks
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 (e.g., used for treatment of anemia for a patient with chronic renal failure who is undergoing dialysis, or furnished from physician's supply incident to a physician service). Coverage includes use in anemia in patients whose religious beliefs forbid blood transfusions. Requirements regarding Hgb values exclude values due to a recent transfusion. For reauthorizations (patient received erythropoietin in previous month): 1) For all uses except surgery, there is an increase in Hgb of at least 1 g/dL after at least 12 weeks of therapy. 2) For anemia in chronic kidney disease, MDS, CHF, RA, HIV, hepatitis C treatment, primary MF, post-polycythemia vera MF, post-essential thrombocythemia MF, or patients whose religious beliefs forbid blood transfusions: current Hgb is less than or equal to 12 g/dL. 3) For anemia due to myelosuppressive cancer chemotherapy: current Hgb is less than 11 g/dL.

PA Criteria	Criteria Details
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Anemia due to myelodysplastic syndromes (MDS), anemia in congestive heart failure (CHF), anemia in rheumatoid arthritis (RA), anemia due to hepatitis C treatment (ribavirin in combination with either interferon alfa or peginterferon alfa), anemia in primary myelofibrosis (MF), post-polycythemia vera MF, and post-essential thrombocythemia MF. Cancer patients who are undergoing palliative treatment.

ERIVEDGE

Products Affected

- Erivedge

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ERLEADA

Products Affected

- Erleada

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ESBRIET

Products Affected

- Esbriet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For idiopathic pulmonary fibrosis (Initial Review 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FARYDAK

Products Affected

- Farydak oral capsule 10 mg, 15 mg, 20 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FASENRA

Products Affected

- Fasenra
- Fasenra Pen

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For severe asthma with an eosinophilic phenotype: For initial therapy: 1) Patient has baseline blood eosinophil count of at least 150 cells per microliter, and 2) Patient has a history of severe asthma despite current treatment with both of the following medications at optimized doses: a) inhaled corticosteroid and b) additional controller (long acting beta2-agonist, leukotriene modifier, or sustained release theophylline). For continuation of therapy: Asthma control has improved on treatment with the requested drug, as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations or a reduction in the daily maintenance oral corticosteroid dose.
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FENTANYL PATCH

Products Affected

- fentanyl transdermal patch 72 hour 100 mcg/hr, 12 mcg/hr, 25 mcg/hr, 50 mcg/hr, 75 mcg/hr

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) 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 2) The requested drug is being prescribed for pain severe enough to require daily, around-the-clock, long-term treatment in a patient who has been taking an opioid AND 3) The patient 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 4) The patient has been evaluated and the patient will be monitored for the development of opioid use disorder
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FETZIMA

Products Affected

- Fetzima oral capsule,Ext Rel 24hr dose pack
- Fetzima oral capsule,extended release 24 hr

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Patient experienced an inadequate treatment response, intolerance, or contraindication to two generic alternatives from the following drug classes: selective serotonin reuptake inhibitors (SSRIs) or serotonin-norepinephrine reuptake inhibitors (SNRIs).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FINTEPLA

Products Affected

- Fintepla

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	2 years and older (initial therapy)
Prescriber Restrictions	Prescribed by or in consultation with an neurologist (initial therapy)
Coverage Duration	1 year
Other Criteria	Dravet Syndrome-Initial therapy-approve if the patient has tried or is concomitantly receiving at least two other antiepileptic drugs or patient has tried or is concomitantly receiving Epidiolex or Diacomit. Dravet Syndrome-Continuation-approve if the patient is responding to therapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FIRAZYR

Products Affected

- icatibant

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The requested drug is being used for the treatment of acute angioedema attacks. Patient has hereditary angioedema (HAE) with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has hereditary angioedema with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, EITHER a) Patient tested positive for an F12, angiopoietin-1, or plasminogen gene mutation OR b) Patient has a family history of angioedema or the angioedema was refractory to a trial of antihistamine for at least one month.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

FYCOMPA

Products Affected

- Fycompa oral suspension
- Fycompa oral tablet 10 mg, 12 mg, 2 mg, 4 mg, 6 mg, 8 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Partial-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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

GATTEX

Products Affected

- Gattex 30-Vial

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For short bowel syndrome (SBS) initial therapy: Patient was dependent on parenteral support for at least 12 months. For SBS continuation: Requirement for parenteral support has decreased from baseline while on therapy with the requested medication.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

GAVRETO

Products Affected

- Gavreto

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	18 years and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	1 year
Other Criteria	NSCLC-approve if the patient has metastatic disease and rearranged during transfection (RET) fusion-positive disease detected by an Food and Drug Administration (FDA) approved test.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

GILENYA

Products Affected

- Gilenya oral capsule 0.5 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

GILOTRIF

Products Affected

- Gilotrif

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC): Patient meets either of the following: A) Patient has metastatic squamous NSCLC that progressed after platinum-based chemotherapy, or B) Patient has a known sensitizing EGFR mutation. For brain metastases from NSCLC, patient has a known sensitizing EGFR mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Brain metastases from non-small cell lung cancer.

GLATIRAMER

Products Affected

- glatiramer subcutaneous syringe 20 mg/mL, 40 mg/mL
- Glatopa subcutaneous syringe 20 mg/mL, 40 mg/mL

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

GROWTH HORMONES

Products Affected

- Genotropin
- Genotropin MiniQuick

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	HIV-1.wasting/cachexia d/t malabsorption, opportunistic infx, depression and other causes which have been addressed prior to starting tx,2.on antiretroviral or HAART for more than 30 days and will cont throughout Serostim tx,3.not being used for alternations in body fat distribution(abdom girth, liopdystrophy, buffalo hump, excess abdm fat)AND 4.unintentional wt loss greater than 10 percent from baseline, wt less than 90 percent of lower limit of IBW or BMI less than or equal to 20 kg/m2. Cont-must be off therapy for 1 month.GHD in Children/Adolescents.Pt meets one of the following-1-had 2 GH stim tests with the following-levodopa, insulin-induced hypoglycemia, arginine, clonidine, or glucagon and both are inadequate as defined by a peak GH response which is below the normal reference range of the testing laboratory OR had at least 1 GH test and results show inadequate response and has at least one risk factor for GHD (e.g., ht for age curve deviated down across 2 major height percentiles [e.g., from above the 25 percentile to below the 10 percentile], growth rate is less than the expected normal growth rate based on age and gender, low IGF-1 and/or IGFBP-3 levels).2.brain radiation or tumor resection and pt has 1 GH stim test and results is inadequate response or has def in at least 1 other pituitary hormone (that is, ACTH, TSH, gonadotropin deficiency [LH and/or FSH] are counted as 1 def], or prolactin).3. congenital hypopituitarism and has 1 GH stim test with inadequate response OR def in at least one other pituitary hormone and/or the patient has the imaging triad of ectopic posterior pituitary and pituitary hypoplasia with abnormal pituitary stalk 4.pt has panhypopituitarism and has pituitary stalk agenesis, empty sella, sellar or supra-sellar mass lesion, or ectopic posterior pituitary bright spot on MRI or CT or pt has 3 or more pituitary hormone def or pt has had 1 GH test and results were inadequate 5.pt had a hypophysectomy.Cont-pt responding to therapy
Age Restrictions	ISS 5 y/o or older, SGA 2 y/o or older, SBS and HIV wasting/cachexia 18 y/o or older
Prescriber Restrictions	GHD (Initial tx children or adolescents w/o hypophysectomy), GHD adults or transitional adolescents, Noonan (initial), Prader Willi (initial for child/adult and cont tx in adults), SHOX (initial), SGA (initial) - prescribed

PA Criteria	Criteria Details
	by or in consultation with an endocrinologist. CKD (initial) endocrinologist or nephrologist.
Coverage Duration	ISS - 6 mos initial, 12 months cont tx, SBS - 1 month, HIV 6 months, others 12 mos
Other Criteria	GHD initial in adults and adolescents 1. endocrine must certify not prescribed for anti-aging or to enhance athletic performance, 2. has either childhood onset or adult onset resulting from GHD alone, multiple hormone deficiency from pituitary dx, hypothalmic dz, pituitary surgery, cranial radiation tx, tumor treatment, TBI or SAH, AND 3. meets one of the following - A. childhood onset has known mutations, embryonic lesions, congenital defects or irreversible structural hypothalmic pituitary lesion/damage, B. 3 or more pituitary hormone def (ACTH, TSH, LH/FSH, or prolactin, age and gender adjusted IGF1 below the lower limits of the normal reference range, AND other causes of low serum IGF-1 have been excluded, C. Neg response to ONE preferred GH stim test (insulin peak response less than or equal to 5 mcg/L, Glucagon peak less than or equal to 3 mcg/L (BMI less than or equal to 25) or less than or equal to 1 mcg/L (BMI is greater than 25), for transitional adults glucagon peak less than or equal to 3 (BMI is less than 25) or less than or equal to 3 if BMI is greater than or equal to 25 and must also have a second GH stim test with low results, if insulin and glucagon contraindicated then Arginine alone test with peak of less than or equal to 0.4 mcg/L, Macrilen peak less than 2.8 ng/ml if BMI is less than or equal to 40 (adults only) AND if a transitional adoles must be off tx for at least one month before retesting. Cont tx - endocrine must certify not prescribed for anti-aging or to enhance athletic performance. ISS initial - baseline ht less than the 1.2 percentile or a standard deviation score (SDS) less than -2.25 for age and gender, open epiphyses, does not have CDGP and ht velocity is either growth rate (GR) is a. less than 4 cm/yr for pts greater than or equal to 5 or b. growth velocity is less than 10th percentile for age/gender. Cont tx - prescriber confirms response to therapy. CKD initial - CKD defined by abnormal CrCl. Noonan initial - baseline ht less than 5th percentile. PW cont tx in adults or adolescents who don't meet child requir - physician certifies not used for anti-aging or to enhance athletic performance. SHOX initial - SHOX def by chromo analysis, open epiphyses, ht less than 3rd percentile for age/gender. SGA initial -baseline ht less than 5th percentile for age/gender and born SGA (birth wt/length that is more than 2 SD below mean for gestational age/gender and didn't have sufficient catch up growth by 2-4 y/o). Cont tx - prescriber confirms response to therapy. Cont Tx for CKD, Noonan, PW in child/adolescent, SHOX, and TS - prescriber confirms response to therapy. SBS initial pt receiving specialized nutritional support.Cont tx - 2nd course if pt responded to tx with a

PA Criteria	Criteria Details
	decrease in the requirement for specialized nutritional support. If requesting Genotropin, Humatrope, Nutropin, Saizen, Norditropin or Zomacton must have tried Omnitrope prior to approval.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	cachexia associated with AIDS, Noonan's Syndrome, renal function impairment with growth failure, short bowel syndrome, and short-stature homeobox-containing gene (SHOX) deficiency

HAEGARDA

Products Affected

- Haegarda subcutaneous recon soln 2,000 unit, 3,000 unit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For hereditary angioedema (HAE): The requested drug is being used for the prevention of acute angioedema attacks. Patient has HAE with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has HAE with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, either 1) Patient tested positive for an F12, angiopoietin-1, or plasminogen gene mutation OR 2) Patient has a family history of angioedema and the angioedema was refractory to a trial of antihistamine for at least one month.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HARVONI

Products Affected

- Harvoni oral tablet 90-400 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic 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, liver and kidney transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current AASLD 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HIGH RISK MEDICATION

Products Affected

- cyproheptadine
- Digitek oral tablet 250 mcg (0.25 mg)
- Digox oral tablet 250 mcg (0.25 mg)
- digoxin oral solution 50 mcg/mL (0.05 mg/mL)
- digoxin oral tablet 250 mcg (0.25 mg)
- guanfacine oral tablet extended release 24 hr
- scopolamine base

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-ANTICONVULSANTS

Products Affected

- phenobarbital

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-ANTIPARKINSON

Products Affected

- benztropine oral
- trihexyphenidyl

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) EPS (extrapyramidal symptoms): 1) The patient has not tried the non-HRM alternative drug amantadine AND 2) The patient has a contraindication to the non-HRM alternative drug amantadine AND 3) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient OR 4) The patient has tried the non-HRM alternative drug amantadine AND 5) The patient experienced an inadequate treatment response OR intolerance to the non-HRM alternative drug amantadine AND 6) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient. Parkinson's: 1) The patient has tried two of the following non-HRM alternative drugs: amantadine, carbidopa/levodopa, pramipexole, or ropinirole have been tried. AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following non-HRM alternative drugs: amantadine, carbidopa/levodopa, pramipexole, or ropinirole AND 3) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off-Label Uses	N/A

HRM-GLYBURIDE

Products Affected

- glyburide micronized oral tablet 1.5 mg, 3 mg, 6 mg
- glyburide oral tablet 1.25 mg, 2.5 mg, 5 mg
- glyburide-metformin oral tablet 1.25-250 mg, 2.5-500 mg, 5-500 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) 1) The patient has not tried one of the following non-HRM alternative drugs: glipizide or metformin AND 2) The patient has a contraindication to one of the following non-HRM alternative drugs: glipizide or metformin AND 3) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient OR 4) The patient has tried one of the following non-HRM alternative drugs: glipizide or metformin AND 5) The patient experienced an inadequate treatment response OR intolerance to one of the following non-HRM alternative drugs: glipizide or metformin AND 6) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-HYDROXYZINE

Products Affected

- hydroxyzine HCl oral solution 10 mg/5 mL
- hydroxyzine HCl oral tablet
- hydroxyzine pamoate oral capsule 25 mg, 50 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) For anxiety: 1) The patient has tried two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline, or venlafaxine extended-release have been tried AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline, or venlafaxine extended-release AND 3) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient OR 4) If being requested for pruritus, prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-HYPNOTICS

Products Affected

- eszopiclone
- zaleplon
- zolpidem oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) 1) The patient has a contraindication to two of the following non-HRM alternative drugs: doxepin (3mg or 6mg) and trazodone AND 2) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient OR 3) One non-HRM alternative drug doxepin (3mg or 6mg) or trazodone has been tried AND 4) The patient experienced an inadequate treatment response OR intolerance to one of the following non-HRM alternative drugs: doxepin (3mg or 6mg) or trazodone AND 5) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient APPLIES TO GREATER THAN CUMULATIVE 90 DAYS OF THERAPY PER YEAR.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-PROMETHAZINE

Products Affected

- promethazine oral

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) Rhinitis: 1) The patient has tried one of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal AND 2) The patient experienced an inadequate treatment response OR intolerance to one of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal AND 3) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient OR 4) The requested drug is being prescribed for urticaria AND 5) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient OR 6) The drug is being requested for antiemetic therapy in postoperative patients or motion sickness AND 7) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient OR 8) The requested drug is being prescribed for any of the following: allergic conjunctivitis, dermatographism, allergic reaction to blood or plasma, sedation, adjunct therapy with analgesics for postoperative pain, angioedema, or adjunct therapy with epinephrine for anaphylaxis after acute symptoms are controlled AND 9) Prescriber must

PA Criteria	Criteria Details
	acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HRM-SKELETAL MUSCLE RELAXANTS

Products Affected

- cyclobenzaprine oral tablet 10 mg, 5 mg
- methocarbamol oral

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweigh the potential risks for this patient.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

HUMIRA

Products Affected

- Humira Pen
- Humira Pen Crohns-UC-HS Start
- Humira Pen Psor-Uveits-Adol HS
- Humira subcutaneous syringe kit 10 mg/0.2 mL, 20 mg/0.4 mL, 40 mg/0.8 mL
- Humira(CF) Pedi Crohns Starter
- Humira(CF) Pen Crohns-UC-HS
- Humira(CF) Pen Psor-Uv-Adol HS
- Humira(CF) subcutaneous syringe kit 10 mg/0.1 mL, 20 mg/0.2 mL, 40 mg/0.4 mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): 1) Inadequate response, intolerance or contraindication to methotrexate (MTX) OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD (e.g., tofacitinib). For moderately to severely active polyarticular juvenile idiopathic arthritis (new starts only): 1) Inadequate response, intolerance or contraindication to MTX OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD). For active ankylosing spondylitis and axial spondyloarthritis (new starts only): Inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial OR intolerance or contraindication to NSAIDs. For moderate to severe chronic plaque psoriasis (new starts only): 1) At least 5% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) Patient meets any of the following: a) Patient has experienced an inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, b) Pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, c) Patient has severe psoriasis that warrants a biologic DMARD as first-line therapy. For moderately to severely active Crohn's disease (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 2) Intolerance or contraindication to conventional therapy. For moderately to severely active ulcerative colitis (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids, aminosalicylates), OR 2) Intolerance or contraindication to conventional therapy.
Age Restrictions	N/A
Prescriber	N/A

PA Criteria	Criteria Details
Restrictions	
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Axial spondyloarthritis.

IBRANCE

Products Affected

- Ibrance

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 years
Other Criteria	Breast cancer - approve advanced (metastatic) hormone receptor positive (HR+) [i.e., estrogen receptor positive- (ER+) and/or progesterone receptor positive (PR+)] disease, and HER2-negative breast cancer for patients who have not had disease progression while on Ibrance, Kisqali or Verzenio when the pt meets ONE of the following 1. Pt is postmenopausal and Ibrance will be used in combination with anastrozole, exemestane, or letrozole 2, pt is premenopausal or perimenopausal and is receiving ovarian suppression/ablation with GnRH agonists, or has had surgical bilateral oophorectomy, or ovarian irradiation AND meets one of the following conditions: Ibrance will be used in combination with anastrozole, exemestane, or letrozole or Ibrance will be used in combination with fulvestrant 3. pt is a man (a man is defined as an individual with the biological traits of a man, regardless of the individual's gender identity or gender expression) who is receiving GnRH agonist AND Ibrance with be used in combination with anastrozole, exemestane, or letrozole or Ibrance will be used in combination with fulvestrant 4. Pt is postmenopausal and Ibrance will be used in combination with fulvestrant
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Liposarcoma

ICLUSIG

Products Affected

- Iclusig

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Follow-up therapy after hematopoietic stem cell transplant (HSCT) for CML and ALL patients.

IDHIFA

Products Affected

- Idhifa

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

IMATINIB

Products Affected

- imatinib oral tablet 100 mg, 400 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML, patient did not fail (excluding failure due to intolerance) prior therapy with a tyrosine kinase inhibitor. For melanoma, c-Kit mutation is positive.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Desmoid tumors, pigmented villonodular synovitis/tenosynovial giant cell tumor (PVNS/TGCT), chordoma, melanoma, and AIDS-related Kaposi sarcoma.

IMBRUVICA

Products Affected

- Imbruvica

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For mantle cell lymphoma: 1) the requested drug will be used in a patient who has received at least one prior therapy, OR 2) the requested drug will be used in combination with rituximab as pretreatment to induction therapy with RHyperCVAD (cyclophosphamide, vincristine, doxorubicin, and dexamethasone) regimen. For gastric MALT lymphoma and non-gastric MALT lymphoma: 1) disease is recurrent, refractory, or progressive, AND 2) 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 central nervous system lymphoma: the disease is relapsed or refractory disease. For nodal marginal zone lymphoma or splenic marginal zone lymphoma: 1) disease is refractory or progressive, AND 2) the requested drug will be used as second-line or subsequent therapy. For histologic transformation of marginal zone lymphoma to diffuse large B-cell lymphoma: the requested drug will be used in patients who have received prior chemoimmunotherapy. For diffuse large B-cell lymphoma: 1) disease is progressive or refractory AND 2) the requested drug will be used as second-line or subsequent therapy. For AIDS-related B-cell lymphoma: the requested drug will be used as second-line or subsequent therapy for relapsed disease. For post-transplant lymphoproliferative disorders: 1) the disease is partially responsive, persistent, or progressive AND 2) the requested drug will be used in patients who have received prior chemoimmunotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.

PA Criteria	Criteria Details
Off-Label Uses	Gastric mucosa-associated lymphoid tissue (MALT) lymphoma, non-gastric MALT lymphoma, hairy cell leukemia, and lymphoplasmacytic lymphoma, follicular lymphoma, primary central nervous system lymphoma, AIDS-related B-cell lymphoma, histologic transformation of marginal zone lymphoma to diffuse large B-cell lymphoma, diffuse large B-cell lymphoma, post-transplant lymphoproliferative disorders.

INCRELEX

Products Affected

- Increlex

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	Patients 2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	1 year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

INLYTA

Products Affected

- Inlyta oral tablet 1 mg, 5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For renal cell carcinoma, the disease is relapsed, metastatic, or unresectable.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Papillary, Hurthle cell, or follicular thyroid carcinoma.

INQOVI

Products Affected

- Inqovi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

INREBIC

Products Affected

- Inrebic

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

IR BEFORE ER

Products Affected

- Hysingla ER
- methadone oral solution
- methadone oral tablet
- morphine oral tablet extended release
- Nucynta ER

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) 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 2) The requested drug is being prescribed for pain severe enough to require daily, around-the-clock, long-term treatment in a patient who has been taking an opioid AND 3) The patient 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 4) The patient has been evaluated and the patient will be monitored for the development of opioid use disorder AND 5) The request is for continuation of therapy for a patient who has been receiving an extended-release opioid agent for at least 30 days OR the patient has severe continuous pain and the patient has received 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

IRESSA

Products Affected

- Iressa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For non-small cell lung cancer (NSCLC) (including brain metastases from NSCLC), patient has a known sensitizing EGFR mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Brain metastases from non-small cell lung cancer.

ISOTRETINOIN

Products Affected

- Amnesteem
- Claravis
- isotretinoin
- Myorisan
- Zenatane

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Refractory acne, 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.

ITRACONAZOLE

Products Affected

- itraconazole oral capsule

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	If for the treatment of onychomycosis due to tinea, the diagnosis has been confirmed by a fungal diagnostic test.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Coccidioidomycosis, Cryptococcosis, Microsporidiosis, Penicilliosis, Sporotrichosis, Pityriasis versicolor/Tinea versicolor, Tinea corporis/Tinea cruris, Tinea capitis, Tinea manuum/Tinea pedis.

IVIG

Products Affected

- Gammagard Liquid
- Gammagard S-D (IgA < 1 mcg/mL)
- Gammaked injection solution 1 gram/10 mL (10 %)
- Gammaplex
- Gammaplex (with sorbitol)
- Gamunex-C injection solution 1 gram/10 mL (10 %)
- Octagam
- Panzyga
- Privigen

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 12 months.
Other Criteria	Part B versus D determination per CMS guidance to establish if drug used for PID in pt's home.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

JAKAFI

Products Affected

- Jakafi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	ALL-less than 21 years of age
Prescriber Restrictions	N/A
Coverage Duration	GVHD/ALL-1 year, all others-3 years.
Other Criteria	For polycythemia vera patients must have tried hydroxyurea. ALL-approve if the mutation/pathway is Janus associated kinase (JAK)-related. GVHD, chronic-approve if the patient has tried one conventional systemic treatment for graft versus host disease.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Acute lymphoblastic leukemia, graft versus host disease, chronic

JUXTAPID

Products Affected

- Juxtapid oral capsule 10 mg, 20 mg, 30 mg, 5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For initiation of therapy to treat homozygous familial hypercholesterolemia: 1) Patient has a diagnosis of homozygous familial hypercholesterolemia (HoFH) confirmed by genetic analysis or clinical criteria (see Other Criteria), AND 2) Prior to initiation of treatment with the requested drug, patient is/was receiving a combination lipid-lowering regimen consisting of at least 2 of the following treatment options: high-intensity statin, fibrate, bile acid sequestrant, ezetimibe, or niacin, at maximally tolerated doses or at the maximum doses approved by the Food and Drug Administration (FDA), (unless patient has intolerance to Statin) AND 3) Prior to initiation of treatment with the requested drug, patient is/was experiencing an inadequate response to such combination regimen as demonstrated by treated low-density lipoprotein cholesterol (LDL-C) greater than 100 mg/dL (or greater than 70 mg/dL with clinical atherosclerotic cardiovascular disease). For renewal of therapy to treat HoFH: 1) Patient meets all initial criteria AND 2) Has responded to therapy as demonstrated by a reduction in LDL-C.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Diagnosis of HoFH must be confirmed by one of the following: 1) Genetic diagnosis: Mutations in both alleles at LDL receptor, apolipoprotein B (ApoB), proprotein convertase subtilisin/kexin type 9 (PCSK9), or LDL receptor adaptor protein/ARH gene locus, OR 2) Clinical diagnosis: Untreated total cholesterol concentration greater than 500 mg/dL, triglyceride concentration less than 300 mg/dL, and documented untreated total cholesterol concentration greater than 250 mg/dL in both parents. Per AHFS https://www.ahfscdi.com/drugs/314017 a) Tendon or cutaneous xanthomas at age 10 or younger, or b) Diagnosis of familial

PA Criteria	Criteria Details
	hypercholesterolemia (FH) by genetic analysis, Simon-Broome Diagnostic Criteria or Dutch Lipid Clinic Network Criteria in both parents, or c) Evidence of FH in both parents with a history including any of the following: Total cholesterol greater than or equal to 310 mg/dL, premature atherosclerotic cardiovascular disease (ASCVD) [before 55 years in men and 60 years in women], tendon xanthoma, or sudden premature cardiac death. Diagnosis of FH must be confirmed by one of the following: 1) Genetic diagnosis: An LDL-receptor mutation, familial defective apo B-100, or a PCSK9 gain-of-function mutation, or 2) Simon-Broome Diagnostic Criteria for FH: Total cholesterol greater than 290 mg/dL or LDL-C greater than 190 mg/dL, plus tendon xanthoma in patient, first-degree (parent, sibling or child) or second-degree relative (grandparent, uncle or aunt), or family history of myocardial infarction in a first degree relative before the age 60 or in a second degree relative before age 50, or total cholesterol greater than 290 mg/dL in an adult first or second degree relative, or total cholesterol greater than 260 mg/dL in a child, brother, or sister aged younger than 16 years, or 3) Dutch Lipid Clinic Network Criteria for FH: Total score greater than 5 points.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

KALYDECO

Products Affected

- Kalydeco

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis: The patient has one mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene that is responsive to ivacaftor potentiation based on clinical and/or in vitro assay data. If the patient's genotype is unknown, an FDA-cleared CF mutation test should be used to detect the presence of a CFTR mutation.
Age Restrictions	6 months of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	The requested drug will not be used in combination with lumacaftor/ivacaftor or tezacaftor/ivacaftor.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

KETOCONAZOLE

Products Affected

- ketoconazole oral

PA Criteria	Criteria Details
Exclusion Criteria	Acute or chronic liver disease. Current use with dofetilide, quinidine, pimozide, cisapride, methadone, disopyramide, dronedarone, ranolazine, ergot alkaloids, alprazolam or simvastatin.
Required Medical Information	1) Patient has one of the following diagnoses: blastomycosis, coccidioidomycosis, histoplasmosis, chromomycosis, or paracoccidioidomycosis, OR 2) The requested drug is being prescribed for a patient with Cushing's syndrome who cannot tolerate surgery or surgery has not been curative.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Cushing's syndrome.

KISQALI

Products Affected

- Kisqali
- Kisqali Femara Co-Pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 years
Other Criteria	Breast cancer - approve advanced or metastatic hormone receptor positive (HR+) [i.e., estrogen receptor positive (ER+) and/or progesterone receptor positive (PR+)]disease, and HER2-negative breast cancer in patients who have not had disease progression while on Kisqali, Ibrance or Verzenio when the pt meets ONE of the following 1. Pt is postmenopausal and Kisqali will be used in combination with anastrozole, exemestane, or letrozole 2. pt is premenopausal or perimenopausal and is receiving ovarian suppression/ablation with GnRH agonist, or has had surgical bilateral oophorectomy, or ovarian irradiation AND Kisqali will be used in combination with anastrozole, exemestane, or letrozole 3. pt is a man (a man is defined as an individual with the biological traits of a man, regardless of the individual's gender identity or gender expression) who is receiving GnRH agonist AND Kisqali with be used in combination with anastrozole, exemestane or letrozole. 4. Patient is postmenopausal, pre/perimenopausal or a man, and Kisqali (not Co-Pack) will be used in combination with Faslodex 5. Patient is pre/perimenopausal and Kisqali (not Co-Pack) will be used in combination with tamoxifen as first line therapy. If the request is for Kisqali Femara, patients do not need to use in combination with anastrozole, exemestane, or letrozole. Patients must have a trial of Ibrance prior to approval of Kisqali/Kisqali Femara Co-Pack unless the patient meets one of the following-a) Patient has been taking Kisqali or Kisqali Femara Co-Pack and is continuing therapy OR b) Patient is pre/perimenopausal and will be using Kisqali or Kisqali Femara Co-Pack

PA Criteria	Criteria Details
	in combination with an aromatase inhibitor as initial endocrine-based therapy OR c) Kisqali will be used in combination with Faslodex in postmenopausal female or male patients as initial endocrine-based therapy OR d) The patient is peri/premenopausal and Kisqali will be used in combination with tamoxifen.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Men with breast cancer

KORLYM

Products Affected

- Korlym

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

KUVAN

Products Affected

- sapropterin

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with Palynziq
Required Medical Information	Diagnosis, Phe concentration
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a specialist who focuses in the treatment of metabolic diseases (initial therapy)
Coverage Duration	Initial-12 weeks, Continuation-1 year
Other Criteria	Initial - approve. Continuation - approve if the patient has had a clinical response (e.g., cognitive and/or behavioral improvements) as determined by the prescribing physician OR patient had a 20% or greater reduction in blood Phe concentration from baseline.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LENVIMA

Products Affected

- Lenvima

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Medullary thyroid carcinoma

LETAIRIS

Products Affected

- ambrisentan

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (WHO Group 1) was confirmed by right heart catheterization. For new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LIDOCAINE PATCHES

Products Affected

- lidocaine topical adhesive patch,medicated 5 %

PA Criteria	Criteria Details
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
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]).

LONG ACTING OPIOIDS

Products Affected

- OxyContin oral tablet,oral only,ext.rel.12 hr

PA Criteria	Criteria Details
Exclusion Criteria	Acute (ie, non-chronic) pain
Required Medical Information	Pain type (chronic vs acute), prior pain medications/therapies tried, concurrent pain medications/therapies
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 12 months.
Other Criteria	For pain severe enough to require daily, around-the-clock, long-term opioid treatment, approve if all of the following criteria are met: 1) patient is not opioid naive, AND 2) non-opioid therapies have been optimized and are being used in conjunction with opioid therapy according to the prescribing physician, AND 3) the prescribing physician has checked the patient's history of controlled substance prescriptions using state prescription drug monitoring program (PDMP), AND 4) the prescribing physician has discussed risks (eg, addiction, overdose) and realistic benefits of opioid therapy with the patient, AND 5) according to the prescriber physician there is a treatment plan (including goals for pain and function) in place and reassessments are scheduled at regular intervals. Patients with cancer, sickle cell disease, in hospice or who reside in a long term care facility are not required to meet above criteria. Clinical criteria incorporated into the quantity limit edits for all oral long-acting opioids require confirmation that the indication is intractable pain (ie, FDA labeled use) prior to reviewing for quantity exception.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LONSURF

Products Affected

- Lonsurf

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For colorectal cancer: The disease is unresectable advanced or metastatic. Patient has progressed on treatment with EITHER a) FOLFOXIRI (fluorouracil, leucovorin, oxaliplatin, and irinotecan) regimen OR b) irinotecan- AND oxaliplatin-based regimens.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LORBRENA

Products Affected

- Lorbrena

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LUPRON

Products Affected

- leuprolide subcutaneous kit
- Lupron Depot (3 month) intramuscular syringe kit 11.25 mg
- Lupron Depot intramuscular syringe kit 3.75 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For central precocious puberty (CPP), patients not currently receiving therapy must meet all of the following criteria: 1) Diagnosis of CPP confirmed by: a) a pubertal response to a gonadotropin releasing hormone (GnRH) agonist test OR a pubertal level of a third generation luteinizing hormone (LH) assay AND b) Assessment of bone age versus chronological age, and 2) The onset of secondary sexual characteristics occurred prior to 8 years of age for female patients OR prior to 9 years of age for male patients. For uterine fibroids, patient must meet one of the following: 1) Diagnosis of anemia (eg, hematocrit less than or equal to 30 percent and/or hemoglobin less than or equal to 10g/dL), OR 2) the requested medication will be used prior to surgery for uterine fibroids.
Age Restrictions	CPP: Patient must be less than 12 years old if female and less than 13 years old if male.
Prescriber Restrictions	N/A
Coverage Duration	Fibroids: 3 months (mo), max 6 mo total. Endometriosis: 6 mo, max 12 mo total. Others: Plan Year
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LYNPARZA

Products Affected

- Lynparza oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For HER2-negative, recurrent or metastatic breast cancer, patient must have a deleterious or suspected deleterious germline BRCA mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

LYRICA CR

Products Affected

- Lyrica CR oral tablet extended release 24 hr 165 mg, 330 mg, 82.5 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

MAVYRET

Products Affected

- Mavyret

PA Criteria	Criteria Details
Exclusion Criteria	Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh class B or C).
Required Medical Information	For chronic 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, liver and kidney transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current AASLD 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

MEGESTROL

Products Affected

- megestrol oral suspension 625 mg/5 mL (125 mg/mL)

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

MEKINIST

Products Affected

- Mekinist

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Mekinist is being used. For melanoma, thyroid cancer, colon or rectal cancer and NSCLC must have documentation of BRAF V600 mutations
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Melanoma must be used in patients with BRAF V600 mutation, and patient has unresectable, advanced (including Stage III or Stage IV disease), or metastatic melanoma. Note-This includes adjuvant treatment in patients with Stage III disease with no evidence of disease post-surgery.For NSCLC requires BRAF V600E Mutation and use in combination with Tafinlar. Thyroid cancer, anaplastic-patient has locally advanced or metastatic anaplastic disease AND Mekinist will be taken in combination with Tafinlar, unless intolerant AND the patient has BRAF V600-positive disease. Colon or Rectal cancer-approve if the patient meets the following (A, B, and C): A) The patient has BRAF V600E mutation-positive disease AND B) The patient has previously received a chemotherapy regimen for colon or rectal cancer AND C) The agent is prescribed as part of a combination regimen for colon or rectal cancer.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Colon or rectal cancer

MEKTOVI

Products Affected

- Mektovi

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

MEMANTINE

Products Affected

- memantine oral capsule,sprinkle,ER 24hr
- memantine oral solution
- memantine oral tablet
- memantine oral tablets,dose pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	This edit only applies to patients less than 30 years of age.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

MIGLUSTAT

Products Affected

- miglustat

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For Gaucher disease, 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

NATPARA

Products Affected

- Natpara

PA Criteria	Criteria Details
Exclusion Criteria	Acute postsurgical hypoparathyroidism (within 6 months of surgery) and expected recovery from the hypoparathyroidism.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

NERLYNX

Products Affected

- Nerlynx

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Brain metastases.

NEXAVAR

Products Affected

- Nexavar

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Osteosarcoma, approve if the patient has tried standard chemotherapy and have relapsed/refractory or metastatic disease. GIST, approve if the patient has tried TWO of the following: imatinib mesylate (Gleevec), sunitinib (Sutent), or regorafenib (Stivarga). Differentiated (ie, papillary, follicular, Hurthle) thyroid carcinoma (DTC), approve if the patient is refractory to radioactive iodine treatment. Medullary thyroid carcinoma, approve if the patient has tried vandetanib (Caprelsa) or cabozantinib (Cometriq). AML - Approve if disease is FLT3-ITD mutation positive as detected by an approved test. Renal cell carcinoma (RCC)-approve if the patient has relapsed or Stage IV clear cell histology and the patient has tried at least one prior systemic therapy (e.g., Inlyta, Votrient, Sutent Cabometyx). Ovarian, fallopian tube, primary peritoneal cancer-approve if the patient has platinum resistant disease and Nexavar is used in combination with topotecan.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Osteosarcoma, angiosarcoma, desmoids tumors (aggressive fibromatosis), gastrointestinal stromal tumors (GIST), medullary thyroid carcinoma, Acute Myeloid Leukemia, Chordoma with recurrent disease, solitary fibrous tumor and hemangiopericytoma, ovarian, fallopian tube, primary peritoneal cancer

NINLARO

Products Affected

- Ninlaro

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	MM - be used in combination with Revlimid and dexamethasone OR pt had received at least ONE previous therapy for multiple myeloma OR the agent will be used following autologous stem cell transplantation (ASCT). Systemic light chain amyloidosis-approve if the patient has tried at least one other regimen for this condition. Waldenstrom Macroglobulinemia/lymphoplasmacytic lymphoma-approve if used in combination with a rituximab product and dexamethasone.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Patients with systemic light chain amyloidosis, Waldenstrom Macroglobulinemia/lymphoplasmacytic lymphoma

NUBEQA

Products Affected

- Nubeqa

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

NUCALA

Products Affected

- Nucala

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For initial therapy for severe asthma with an eosinophilic phenotype: 1) Patient has baseline blood eosinophil count of at least 150 cells per microliter, and 2) Patient has a history of severe asthma despite current treatment with both of the following medications at optimized doses: a) inhaled corticosteroid and b) additional controller (long acting beta2-agonist, leukotriene modifier, or sustained-release theophylline). For continuation therapy for severe asthma with an eosinophilic phenotype: Asthma control has improved on treatment with the requested drug, demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations or a reduction in the daily maintenance oral corticosteroid dose. For initial therapy for eosinophilic granulomatosis with polyangiitis (EGPA): Patient has a history or the presence of an eosinophil count of more than 1000 cells per microliter or a blood eosinophil level greater than 10 percent. For continuation of therapy for EGPA: Patient has a beneficial response to treatment with the requested drug, demonstrated by any of the following: 1) a reduction in the frequency of relapses, 2) a reduction in the daily oral corticosteroid dose, or 3) no active vasculitis.
Age Restrictions	Asthma: 6 years of age or older, EGPA: 18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

NUEDEXTA

Products Affected

- Nuedexta

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

NUPLAZID

Products Affected

- Nuplazid oral capsule
- Nuplazid oral tablet 10 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OCALIVA

Products Affected

- Ocaliva

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Prescriber specialty, lab values, prior medications used for diagnosis and length of trials
Age Restrictions	18 years and older (initial)
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, hepatologist, or liver transplant physician (initial)
Coverage Duration	6 months initial, 1 year cont.
Other Criteria	Initial treatment of PBC-Patient must meet both 1 and 2-1. Patient has a diagnosis of PBC as defined by TWO of the following:a)Alkaline phosphatase (ALP) elevated above the upper limit of normal as defined by normal laboratory reference values b)Positive anti-mitochondrial antibodies (AMAs) or other PBC-specific auto-antibodies, including sp100 or gp210, if AMA is negative c)Histologic evidence of primary biliary cholangitis (PBC) from a liver biopsy 2. Patient meets ONE of the following: a) Patient has been receiving ursodiol therapy for greater than or equal to 1 year and has had an inadequate response. b) Patient is unable to tolerate ursodiol therapy. Cont tx - approve if the patient has responded to Ocaliva therapy as determined by the prescribing physician (e.g., improved biochemical markers of PBC (e.g., alkaline phosphatase [ALP], bilirubin, gamma-glutamyl transpeptidase [GGT], aspartate aminotransferase [AST], alanine aminotransferase [ALT] levels)).
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OCTREOTIDE

Products Affected

- octreotide acetate injection solution

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For acromegaly (initial): 1) Patient 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) Patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy. For meningiomas: patient has unresectable disease.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	For acromegaly (continuation of therapy): patient's IGF-1 level has decreased or normalized since initiation of therapy.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Meningiomas, thymomas and thymic carcinomas, and neuroendocrine tumors (NETs) of the gastrointestinal (GI) tract, thymus, lung, and pancreas.

ODOMZO

Products Affected

- Odomzo

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OFEV

Products Affected

- Ofev

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For idiopathic pulmonary fibrosis (Initial Review 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OPSUMIT

Products Affected

- Opsumit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (WHO Group 1) was confirmed by right heart catheterization. For new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ORAL-INTRANASAL FENTANYL

Products Affected

- fentanyl citrate buccal lozenge on a handle

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is indicated for the treatment of breakthrough CANCER-related pain only. The requested drug is being prescribed for the management of breakthrough pain in a CANCER patient who is currently receiving around-the-clock opioid therapy for underlying CANCER pain. [Note: Ensure that the patient is opioid tolerant. Patients considered opioid tolerant are those who are taking around-the-clock medicine consisting of at least 60 mg of oral morphine per day, at least 25 mcg per hour of transdermal fentanyl, at least 30 mg of oral oxycodone per day, at least 60 mg oral hydrocodone per day, at least 8 mg of oral hydromorphone per day, at least 25 mg oral oxymorphone per day, or an equianalgesic dose of another opioid medication daily for a week or longer.] AND 2) The International Classification of Diseases (ICD) diagnosis code provided supports the CANCER-RELATED diagnosis. [Note: For drug coverage approval, ICD diagnosis code provided MUST support the CANCER-RELATED diagnosis.]
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ORFADIN

Products Affected

- nitisinone
- Orfadin oral capsule 20 mg
- Orfadin oral suspension

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For hereditary tyrosinemia type 1: Diagnosis of hereditary tyrosinemia type 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ORKAMBI

Products Affected

- Orkambi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis: the patient is positive for the F508del mutation on both alleles of the cystic fibrosis transmembrane conductance regulator (CFTR) gene.
Age Restrictions	2 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	The requested drug will not be used in combination with ivacaftor or tezacaftor/ivacaftor.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OSPHENA

Products Affected

- Osphena

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OXANDROLONE

Products Affected

- oxandrolone

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	Coverage will be denied if request is for an indication excluded from Part D.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Cachexia associated with AIDS (HIV-wasting) or to enhance growth in patients with Turner's Syndrome.

PEGASYS

Products Affected

- Pegasys

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic hepatitis C (CHC): CHC infection confirmed by presence of HCV RNA in 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, liver and kidney transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current AASLD-IDSA treatment guidelines.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	HCV=Criteria will be applied consistent with current AASLD-IDSA guidance. HBV=48 wks. Other=Plan Yr
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Myeloproliferative neoplasm (essential thrombocythemia, polycythemia vera, primary myelofibrosis and post-polycythemia vera or post-essential thrombocythemia myelofibrosis)

PEMAZYRE

Products Affected

- Pemazyre

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis, prior therapies
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years
Other Criteria	Cholangiocarcinoma-approve if the patient has unresectable locally advanced or metastatic disease with a fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement, as detected by an approved test AND the patient has been previously treated with at least one systemic therapy regimen
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

PHENYLBUTYRATE

Products Affected

- sodium phenylbutyrate

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For urea cycle disorder: Diagnosis of urea cycle disorder (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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

PIQRAY

Products Affected

- Piqray

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

POMALYST

Products Affected

- Pomalyst

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years
Other Criteria	AIDS-Related Kaposi Sarcoma-approve if the patient has relapsed or refractory disease. CNS Lymphoma-approve if the patient has relapsed or refractory disease
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Systemic Light Chain Amyloidosis, Central Nervous System (CNS) Lymphoma

PRALUENT

Products Affected

- Praluent Pen

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

PREGABALIN

Products Affected

- pregabalin oral capsule 100 mg, 150 mg, 200 mg, 225 mg, 25 mg, 300 mg, 50 mg, 75 mg
- pregabalin oral solution

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for the management of postherpetic neuralgia, neuropathic pain associated with diabetic peripheral neuropathy, cancer-related neuropathic pain or cancer treatment related neuropathic pain AND 2) The patient has experienced an inadequate treatment response, intolerance, or contraindication to gabapentin OR 3) The requested drug is being prescribed as adjunctive therapy for partial onset seizures OR 4) The requested drug is being prescribed for the management of fibromyalgia or management of neuropathic pain associated with spinal cord injury.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Cancer-related neuropathic pain, cancer treatment related neuropathic pain.

PROMACTA

Products Affected

- Promacta oral powder in packet 12.5 mg
- Promacta oral tablet 12.5 mg, 25 mg, 50 mg, 75 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Cause of thrombocytopenia. Thrombocytopenia due to HCV-related cirrhosis, platelet counts. Severe aplastic anemia, platelet counts and prior therapy. MDS-platelet counts.
Age Restrictions	N/A
Prescriber Restrictions	Thrombocytopenia due to chronic ITP or Aplastic Anemia, approve if prescribed by, or after consultation with, a hematologist. Thrombocytopenia due to HCV-related cirrhosis, approve if prescribed by, or after consultation with, a gastroenterologist, hematologist, hepatologist, or a physician who specializes in infectious disease. MDS-presc or after consult with heme/onc.
Coverage Duration	Chronic ITP - 3 years, others 12 months.
Other Criteria	Thrombocytopenia in patients with chronic immune (idiopathic) thrombocytopenia purpura, approve if the patient has tried ONE other therapy or has undergone a splenectomy. Treatment of thrombocytopenia due to HCV-related cirrhosis, approve to allow for initiation of antiviral therapy if the patient has low platelet counts (eg, less than 75,000 mm3) and the patient has chronic HCV infection and is a candidate for hepatitis C therapy. Aplastic anemia - has low platelet counts at baseline/pretreatment (e.g., less than 30,000 mm3) AND tried one immunosuppressant therapy (e.g., cyclosporine, mycophenolate moefetil, sirolimus) OR patient will be using Promacta in combination with standard immunosuppressive therapy. MDS-approve if patient has low- to intermediate-risk MDS AND according to the prescriber the patient has clinically-significant thrombocytopenia (e.g., low platelet counts [pretreatment], is platelet transfusion-dependent, active bleeding, and/or a history of bleeding at low platelet counts).
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Thrombocytopenia in Myelodysplastic Syndrome (MDS)

PULMOZYME

Products Affected

- Pulmozyme

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis: Diagnosis of cystic fibrosis was confirmed by appropriate diagnostic or genetic testing.
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.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

QINLOCK

Products Affected

- Qinlock

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis, other therapies tried
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years
Other Criteria	Gastrointestinal stromal tumor (GIST), advanced-approve if, according to labeling, the patient has been previously treated with imatinib and at least two other kinase inhibitors, in addition to imatinib.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

QUETIAPINE XR

Products Affected

- quetiapine oral tablet extended release 24 hr 150 mg, 200 mg, 300 mg, 400 mg, 50 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For schizophrenia, acute treatment of manic or mixed episodes associated with bipolar I disorder, both as monotherapy and as an adjunct to lithium or divalproex, the acute treatment of depressive episodes associated with bipolar disorder, maintenance treatment of bipolar I disorder, as an adjunct to lithium or divalproex, adjunctive treatment of major depressive disorder, or maintenance monotherapy treatment in bipolar I disorder: The patient has had an inadequate treatment response, intolerance or contraindication to one of the following: aripiprazole, lurasidone, olanzapine, paliperidone, quetiapine immediate-release, risperidone, or ziprasidone
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Maintenance monotherapy treatment in bipolar I disorder, monotherapy treatment of generalized anxiety disorder, monotherapy treatment of major depressive disorder

QUININE SULFATE

Products Affected

- quinine sulfate

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 month
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Babesiosis, uncomplicated Plasmodium vivax malaria.

REGRANEX

Products Affected

- Regranex

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For the treatment of lower extremity diabetic neuropathic ulcers that extend into the subcutaneous tissue or beyond and have an adequate blood supply
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	20 weeks
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

RELISTOR INJ

Products Affected

- Relistor subcutaneous solution
- Relistor subcutaneous syringe

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for opioid-induced constipation in an adult patient with advanced illness or pain caused by active cancer who requires opioid dosage escalation for palliative care OR 2) The requested drug is being prescribed for opioid-induced constipation in an adult patient 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 AND 3) The patient is unable to tolerate oral medications OR 4) An oral drug indicated for opioid-induced constipation in an adult patient with chronic non-cancer pain has been tried. (Note: Examples are Amitiza or Movantik) AND 5) The patient experienced an inadequate treatment response or intolerance to an oral drug indicated for opioid-induced constipation in an adult patient with chronic non-cancer pain. (Note: Examples are Amitiza or Movantik) OR 6) The patient has a contraindication to an oral drug indicated for opioid-induced constipation in an adult patient with chronic non-cancer pain (Note: Examples are Amitiza or Movantik).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	4 months
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

RETEVMO

Products Affected

- Retevmo oral capsule 40 mg, 80 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	Medullary Thyroid Cancer/Thyroid Cancer-12 years and older
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years
Other Criteria	Non-Small Cell Lung Cancer (NSCLC)-Approve if the patient has metastatic disease AND the tumor is RET fusion-positive. Medullary Thyroid Cancer-approve if the patient has advanced or metastatic RET-mutant disease and the disease requires treatment with systemic therapy.Thyroid Cancer-approve if the patient has advanced or metastatic RET fusion positive disease, the disease is radioactive iodine-refractory (if radioactive iodine is appropriate) and the disease requires treatment with systemic therapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

REVLIMID

Products Affected

- Revlimid

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For myelodysplastic syndrome (MDS): Low- to intermediate-1 risk MDS with symptomatic anemia
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
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, myelofibrosis-associated anemia, POEMS syndrome, non-Hodgkin's lymphoma with the following subtypes: AIDS-related diffuse large B-cell lymphoma, primary central nervous system (CNS) lymphoma, monomorphic post-transplant lymphoproliferative disorder, chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL), diffuse large B-cell lymphoma, nongastric/gastric mucosa associated lymphoid tissue (MALT) lymphoma, primary cutaneous B-cell lymphoma, multicentric Castleman's disease, adult T-cell leukemia/lymphoma, mycosis fungoides (MF)/Sezary syndrome (SS), angioimmunoblastic T-cell lymphoma (AITL), peripheral T-cell lymphoma not otherwise specified (PTCL NOS), enteropathy-associated T-cell lymphoma, monomorphic epitheliotropic intestinal T-cell lymphoma, nodal peripheral T-cell lymphoma, primary cutaneous anaplastic large cell lymphoma (ALCL).

RINVOQ

Products Affected

- Rinvoq

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): 1) inadequate response, intolerance or contraindication to methotrexate (MTX) OR 2) inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD (e.g. tofacitinib).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ROZLYTREK

Products Affected

- Rozlytrek

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

RUBRACA

Products Affected

- Rubraca

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

RYDAPT

Products Affected

- Rydapt

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For acute myeloid leukemia (AML), AML must be FLT3 mutation-positive.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Relapsed or refractory acute myeloid leukemia

SAMSCA

Products Affected

- Samsca oral tablet 15 mg
- tolvaptan oral tablet 30 mg

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with Jynarque.
Required Medical Information	Serum sodium less than 125 mEq/L at baseline or less marked hyponatremia, defined as serum sodium less than 135 mEq/L at baseline, that is symptomatic (eg, nausea, vomiting, headache, lethargy, confusion).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 30 days
Other Criteria	Hyponatremia - Pt must meet ONE of the following: 1. serum sodium less than 125 mEq/L at baseline, OR 2. marked hyponatremia, defined as less than 135 mEq/L at baseline, that is symptomatic (eg, nausea, vomiting, headache, lethargy, confusion), OR 3. patient has already been started on Samsca and has received less than 30 days of therapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SIGNIFOR

Products Affected

- Signifor

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SILDENAFIL

Products Affected

- sildenafil (Pulmonary Arterial Hypertension) oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For pulmonary arterial hypertension (WHO Group 1), the diagnosis was confirmed by right heart catheterization. For new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SIRTURO

Products Affected

- Sirturo oral tablet 100 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	The requested drug is not being prescribed for the treatment of latent infection due to Mycobacterium tuberculosis, drug-sensitive tuberculosis, extra-pulmonary tuberculosis, or infection caused by the non-tuberculous mycobacteria
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SKYRIZI

Products Affected

- Skyrizi subcutaneous syringe kit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For moderate to severe plaque psoriasis (new starts only): 1) at least 5% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) patient meets any of the following: a) patient has experienced an inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, or b) pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, or c) patient has severe psoriasis that warrants a biologic disease-modifying antirheumatic drug (DMARD) as first-line therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SOMATULINE DEPOT

Products Affected

- Somatuline Depot subcutaneous syringe 120 mg/0.5 mL, 60 mg/0.2 mL, 90 mg/0.3 mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For acromegaly (initial): 1) patient 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) patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	For acromegaly continuation of therapy: patient's IGF-1 level has decreased or normalized since initiation of therapy.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Neuroendocrine tumors (NETs) of the gastrointestinal (GI) tract, thymus, lung, and pancreas

SOMAVERT

Products Affected

- Somavert

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For acromegaly (initial): 1) patient 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) patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	For acromegaly continuation of therapy: patient's IGF-1 level has decreased or normalized since initiation of therapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SOVALDI

Products Affected

- Sovaldi oral tablet 400 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Genotype 1 and 4 -18 years or older, Genotype 2 and 3-3 years or older
Prescriber Restrictions	Prescribed by or in consultation w/ GI, hepatologist, ID, or a liver transplant MD
Coverage Duration	Will be c/w AASLD guidance and inclusive of treatment already received for the requested drug
Other Criteria	Criteria will be applied consistent with current AASLD/IDSA guidance. And, patients with genotype 1 and 4 must have a trial with Harvoni, Vosevi or Epclusa prior to approval of Sovaldi, unless Harvoni, Vosevi and Epclusa are not specifically listed as an alternative therapy for a specific patient population in the guidelines. Patients with Genotype 2 and 3 must have a trial of Epclusa or Vosevi prior to approval of Sovaldi, unless Epclusa and Vosevi are not specifically listed as an alternative therapy for a specific patient population in the guidelines.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Indications consistent with current AASLD/IDSA guidance

SPRYCEL

Products Affected

- Sprycel

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Sprycel is being used. For indications of CML and ALL, the Philadelphia chromosome (Ph) status of the leukemia must be reported.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	For CML, new patient must have Ph-positive CML for approval of Sprycel. For ALL, new patient must have Ph-positive ALL for approval of Sprycel. GIST - has D842V mutation AND previously tried Sutent and Gleevec.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	GIST, chondrosarcoma, chordoma

STELARA

Products Affected

- Stelara subcutaneous solution
- Stelara subcutaneous syringe 45 mg/0.5 mL, 90 mg/mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For moderate to severe plaque psoriasis (new starts only): 1) at least 5% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) the patient had an inadequate response, intolerance, or contraindication to Humira. For active psoriatic arthritis (PsA) (new starts only): the patient had an inadequate response, intolerance, or contraindication to Humira or Xeljanz/Xeljanz XR. For moderately to severely active Crohn's disease (new starts only): patient had an inadequate response, intolerance, or contraindication to Humira. For moderately to severely active ulcerative colitis (new starts only): patient had an inadequate response, intolerance, or contraindication to Humira or Xeljanz.
Age Restrictions	Plaque psoriasis: 6 years of age or older. All other indications: 18 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

STIVARGA

Products Affected

- Stivarga

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Stivarga is being used. Prior therapies tried. For metastatic CRC, KRAS/NRAS mutation status.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	For GIST, patient must have previously been treated with imatinib (Gleevec) and sunitinib (Sutent). For HCC, patient must have previously been treated with at least one tyrosine kinase inhibitor (e.g., Nexavar, Lenvima). Soft tissue sarcoma-approve if the patient has non-adipocytic extremity/superficial trunk, head/neck or retroperitoneal/intra-abdominal sarcoma OR pleomorphic rhabdomyosarcoma. Osteosarcoma-approve if the patient has relapsed/refractory or metastatic disease and the requested medication is being used as subsequent therapy.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Soft tissue Sarcoma, Osteosarcoma

SUTENT

Products Affected

- Sutent

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Gastrointestinal stromal tumors (GIST), approve if the patient has tried imatinib (Gleevec). Chordoma, approve if the patient has recurrent disease. Differentiated thyroid carcinoma, approve if the patient is refractory to radioactive iodine therapy. Medullary thyroid carcinoma, approve if the patient has tried vandetanib (Caprelsa) or cabozantinib (Cometriq). Meningioma, approve if the patient has recurrent or progressive disease. Thymic carcinoma - has tried chemotherapy or radiation therapy. Renal Cell Carcinoma (RCC), clear cell or non-clear cell histology-approve if the patient is at high risk of recurrent clear cell RCC following nephrectomy and Sutent is used for adjuvant therapy or if the patient has relapsed or Stage IV disease. Neuroendocrine tumors of the pancreas-approve for advanced or metastatic disease.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Chordoma, angiosarcoma, solitary fibrous tumor/hemangiopericytoma, alveolar soft part sarcoma (ASPS), differentiated (ie, papillary, follicular, and Hurthle) thyroid carcinoma, medullary thyroid carcinoma, meningioma, thymic carcinoma.

SYMDEKO

Products Affected

- Symdeko

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis (CF): The patient is positive for the F508del mutation on both alleles of the cystic fibrosis transmembrane conductance regulator (CFTR) gene OR the patient has a mutation in the CFTR gene that is responsive to tezacaftor/ivacaftor potentiation based on clinical and/or in vitro assay data. If the patient's genotype is unknown, an FDA-cleared CF mutation test should be used to detect the presence of a CFTR mutation
Age Restrictions	6 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	Symdeko will not be used in combination with Orkambi or Kalydeco.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SYMPAZAN

Products Affected

- Sympazan

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

SYNRIBO

Products Affected

- Synribo

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Follow-up therapy for CML patients after hematopoietic stem cell transplant (HSCT), treatment of chronic CML patients with a T315I mutation.

TABRECTA

Products Affected

- Tabrecta

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Non-Small Cell Lung Cancer (NSCLC)-Approve if the patient has metastatic disease AND the tumor is positive for a mutation that leads to mesenchymal-epithelial transition (MET) exon 14 skipping, as detected by an approved test.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TAFINLAR

Products Affected

- Tafinlar

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Tafinlar is being used. BRAF V600 mutations
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Melanoma with BRAF V600 mutation AND patient has unresectable, advanced (including Stage III or Stage IV disease) or metastatic melanoma. Note-This includes adjuvant treatment in patients with Stage III disease with no evidence of disease post-surgery. For NSCLC, must have BRAF V600E mutation. Thyroid Cancer, anaplastic-must have BRAF V600-positive disease AND Tafinlar will be taken in combination with Mekinist, unless intolerant AND the patient has locally advanced or metastatic anaplastic disease. Thyroid Cancer, differentiated (i.e., papillary, follicular, or Hurthle cell) AND the patient has disease that is refractory to radioactive iodine therapy AND the patient has BRAF-positive disease. Colon or Rectal cancer-approve if the patient meets the following (A, B, and C): A) The patient has BRAF V600E mutation-positive disease AND B) The patient has previously received a chemotherapy regimen for colon or rectal cancer AND C) The agent is prescribed as part of a combination regimen for colon or rectal cancer.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Patients with Differentiated Thyroid Cancer, Colon or rectal cancer

TAGRISSO

Products Affected

- Tagrisso

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For metastatic or recurrent non-small cell lung cancer (NSCLC), patient must have sensitizing EGFR mutation-positive NSCLC (including brain metastases from non-small cell lung cancer).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Sensitizing epidermal growth factor receptor (EGFR) mutation-positive recurrent or metastatic non-small cell lung cancer, brain metastases from non-small cell lung cancer.

TALZENNA

Products Affected

- Talzenna

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TARCEVA

Products Affected

- erlotinib oral tablet 100 mg, 150 mg, 25 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For NSCLC (including brain metastases from NSCLC), patient has a known sensitizing EGFR mutation. For pancreatic cancer, the disease is locally advanced, unresectable, or metastatic.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Chordoma, renal cell carcinoma (RCC), brain metastases from non-small cell lung cancer (NSCLC).

TASIGNA

Products Affected

- Tasigna

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For chronic myelogenous leukemia (CML) or acute lymphoblastic leukemia (ALL), diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML, 1) patient has received a hematopoietic stem cell transplant, OR 2) patient has accelerated or blast phase CML, OR 3) For chronic phase CML, the patient has one of the following: a) patient is 18 years of age or younger, b) high or intermediate risk for disease progression, or c) low risk for disease progression and has experienced resistance, intolerance or toxicity to imatinib or an alternative tyrosine kinase inhibitor. If patient experienced resistance to imatinib or an alternative tyrosine kinase inhibitor for CML, patient is negative for T315I mutation. For GIST, patient must have progressed on imatinib, sunitinib or regorafenib.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), gastrointestinal stromal tumor (GIST).

TAZAROTENE

Products Affected

- tazarotene
- Tazorac topical cream 0.05 %

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For plaque psoriasis, the requested drug is being prescribed to treat less than 20 percent of the patient's body surface area.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TAZVERIK

Products Affected

- Tazverik

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	16 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TECFIDERA

Products Affected

- dimethyl fumarate oral capsule,delayed release(DR/EC) 120 mg, 240 mg
- Tecfidera oral capsule,delayed release(DR/EC) 120 mg, 120 mg (14)- 240 mg (46), 240 mg

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with other disease-modifying agents used for multiple sclerosis (MS).
Required Medical Information	Relapsing form of Multiple Sclerosis (MS), to include, clinically-isolated syndrome, relapsing-remitting disease, and active secondary progressive disease.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist or MS specialist.
Coverage Duration	Authorization will be for 1 year.
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TERIPARATIDE

Products Affected

- teriparatide

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with other medications for osteoporosis (eg, denosumab [Prolia], bisphosphonates, raloxifene, calcitonin nasal spray [Fortical], abaloparatide), except calcium and Vitamin D.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Auth for 2 yr total therapy between Tymlos, Bonsity and teriparatide over a pt's lifetime
Other Criteria	Treatment of PMO, approve if pt has tried one oral bisphosphonate OR pt cannot take an oral bisphosphonate because the pt cannot swallow or has difficulty swallowing or the pt cannot remain in an upright position post oral bisphosphonate administration or pt has a pre-existing GI medical condition (eg, patient with esophageal lesions, esophageal ulcers, or abnormalities of the esophagus that delay esophageal emptying [stricture, achalasia]), OR pt has tried an IV bisphosphonate (ibandronate or zoledronic acid), OR pt has severe renal impairment (creatinine clearance less than 35 mL/min) or CKD or pt has had an osteoporotic fracture or fragility fracture. Increase bone mass in men (a man is defined as an individual with the biological traits of a man, regardless of the individual's gender identity or gender expression) with primary or hypogondal osteoporosis/Treatment of GIO, approve if pt tried one oral bisphosphonate OR pt cannot take an oral bisphosphonate because the patient cannot swallow or has difficulty swallowing or the patient cannot remain in an upright position post oral bisphosphonate administration or has a pre-existing GI medical condition (eg, patient with esophageal lesions, esophageal ulcers, or abnormalities of the esophagus that delay esophageal emptying [stricture, achalasia]), OR pt has tried zoledronic acid (Reclast), OR pt has severe renal impairment (CrCL less than 35 mL/min) or has CKD or has had an osteoporotic fracture or fragility fracture. If the request

PA Criteria	Criteria Details
	is for brand name Forteo, patients must have a trial of teriparatide first.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TESTOSTERONE CYPIONATE INJ

Products Affected

- testosterone cypionate intramuscular oil 100 mg/mL, 200 mg/mL, 200 mg/mL (1 ML)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Request is for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who has had a confirmed low testosterone level according to current practice guidelines or your standard male lab reference values before starting testosterone therapy OR 2) Request is not for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who has at least two confirmed low testosterone levels according to current practice guidelines or your standard male lab reference values OR 3) Requested drug is being prescribed for gender dysphoria in a transgender male patient who is able to make an informed, mature decision to engage in therapy
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Gender Dysphoria in transgender male patients

TESTOSTERONE ENANTHATE INJ

Products Affected

- testosterone enanthate

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Request is for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who had a confirmed low testosterone level according to current practice guidelines or your standard male lab reference values before starting testosterone therapy OR 2) Request is not for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who has at least two confirmed low testosterone levels according to current practice guidelines or your standard male lab reference values OR 3) Requested drug is being prescribed for inoperable metastatic breast cancer in a patient who is 1 to 5 years postmenopausal and who has had an incomplete response to other therapy for metastatic breast cancer OR 4) Requested drug is being prescribed for a premenopausal patient with breast cancer who has benefited from oophorectomy and is considered to have a hormone-responsive tumor OR 5) Requested drug is being prescribed for delayed puberty in a male patient OR 6) Requested drug is being prescribed for gender dysphoria in a transgender male patient who is able to make an informed, mature decision to engage in therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Gender Dysphoria in transgender male patients.

TETRABENAZINE

Products Affected

- tetrabenazine oral tablet 12.5 mg, 25 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For treatment of chorea associated with Huntington's disease and tardive dyskinesia: The patient must have a prior inadequate response or intolerable adverse event with deutetrabenazine therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Chronic tics, tardive dyskinesia, hemiballismus, chorea not associated with Huntington's disease.

THALOMID

Products Affected

- Thalomid oral capsule 100 mg, 150 mg, 200 mg, 50 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cachexia: Cachexia must be due to cancer or human immunodeficiency virus (HIV) infection. For Kaposi's sarcoma: The patient has human immunodeficiency virus (HIV) infection.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Myelofibrosis-related anemia, recurrent aphthous stomatitis, recurrent HIV-associated aphthous ulcers, cachexia, human immunodeficiency virus (HIV)-associated diarrhea, Kaposi's sarcoma, Behcet's syndrome, chronic graft-versus-host disease, Crohn's disease, multicentric Castleman's disease.

TIBSOVO

Products Affected

- Tibsovo

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TOBRAMYCIN

Products Affected

- tobramycin in 0.225 % NaCl

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis and non-cystic fibrosis bronchiectasis, the patient must meet one of the following: 1) Pseudomonas aeruginosa is present in the patient's airway cultures, OR 2) the patient 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	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.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Non-cystic fibrosis bronchiectasis

TOPICAL LIDOCAINE

Products Affected

- lidocaine HCl mucous membrane jelly
- lidocaine HCl mucous membrane solution 4 % (40 mg/mL)
- lidocaine topical ointment
- lidocaine-prilocaine topical cream

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being used for topical anesthesia, 2) If the requested drug will be used as part of a compounded product, then all the active ingredients in the compounded product are FDA-approved for topical use
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
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.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TOPICAL TESTOSTERONES

Products Affected

- Androderm
- testosterone transdermal gel in metered-dose pump 12.5 mg/ 1.25 gram (1 %)
- testosterone transdermal gel in packet 1 % (25 mg/2.5gram), 1 % (50 mg/5 gram)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Request is for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who had a confirmed low testosterone level according to current practice guidelines or your standard male lab reference values before starting testosterone therapy OR 2) Request is not for continuation of testosterone therapy and requested drug is being prescribed for hypogonadism in a male patient or a patient that self-identifies as male who has at least two confirmed low testosterone levels according to current practice guidelines or your standard male lab reference values OR 3) Requested drug is being prescribed for gender dysphoria in a transgender male patient who is able to make an informed, mature decision to engage in therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Gender Dysphoria in transgender male patients.

TOPICAL TRETINOIN

Products Affected

- Avita topical cream
- Avita topical gel
- tretinoin topical cream
- tretinoin topical gel 0.01 %, 0.025 %

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TRELSTAR

Products Affected

- Trelstar intramuscular suspension for reconstitution 11.25 mg, 3.75 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TRIENTINE

Products Affected

- trientine

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TRIKAFTA

Products Affected

- Trikafta

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For cystic fibrosis (CF): The patient has at least one F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene.
Age Restrictions	12 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	The requested medication will not be used in combination with other medications containing ivacaftor.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TUKYSA

Products Affected

- Tukysa oral tablet 150 mg, 50 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis, prior therapies
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Breast Cancer-approve if the patient has advanced unresectable or metastatic human epidermal growth factor receptor 2 (HER2)-positive disease, has received at least one prior anti-HER2-based regimen in the metastatic setting and Tukysa is used in combination with trastuzumab and capecitabine.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TURALIO

Products Affected

- Turalio

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

TYKERB

Products Affected

- lapatinib
- Tykerb

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Tykerb is being used. Metastatic breast cancer, HER2 status or hormone receptor (HR) status.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	HER2-positive advanced or metastatic breast cancer, approve if Tykerb will be used in combination with Xeloda or Herceptin and the patient has received prior therapy with Herceptin. HER2-positive HR positive metastatic breast cancer, approve if the patient is a man receiving a GnRH agonist, a premenopausal or perimenopausal woman receiving ovarian suppression/ablation with a GnRH agonist, surgical bilateral oophorectomy or ovarian radiation, or a postmenopausal woman and Tykerb will be used in combination with an aromatase inhibitor, that is letrozole (Femara), anastrozole, or exemestane. In this criteria, man/woman is defined as an individual with the biological traits of a man/woman, regardless of the individual's gender identity or expression. Colon or rectal cancer-approve if the patient has unresectable advanced or metastatic disease that is human epidermal receptor 2 (HER2) amplified and with wild-type RAS and the medication is used as subsequent therapy in combination with trastuzumab and the patient has not been previously treated with a HER2-inhibitor.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Bone cancer-chordoma, colon or rectal cancer

UPTRAVI

Products Affected

- Uptravi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Confirmation of right heart catheterization, medication history.
Age Restrictions	N/A
Prescriber Restrictions	PAH must be prescribed by, or in consultation with, a cardiologist or a pulmonologist.
Coverage Duration	1 year
Other Criteria	Must have PAH (WHO Group 1) and had a right heart catheterization to confirm the diagnosis of PAH (WHO Group 1). Patient new to therapy must meet a) OR b): a) tried TWO or is currently taking TWO oral therapies for PAH (either alone or in combination) each for 30 days, unless patient has experienced treatment failure, intolerance, or oral therapy is contraindicated: PDE5 inhibitor (eg, sildenafil, Revatio), endothelin receptor antagonist (ERA) [eg, Tracleer, Letairis or Opsumit], or Adempas, OR b) receiving or has received in the past one prostacyclin therapy for PAH (eg, Orenitram, Ventavis, or epoprostenol injection).
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VALCHLOR

Products Affected

- Valchlor

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Chronic or smoldering adult T-cell leukemia/lymphoma, Stage 2 or higher mycosis fungoides/Sezary syndrome, primary cutaneous marginal zone lymphoma, primary cutaneous follicle center lymphoma, lymphomatoid papulosis.

VELTASSA

Products Affected

- Veltassa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The patient has experienced an inadequate treatment response or intolerance to Lokelma OR 2) The patient has a contraindication that would prohibit a trial of Lokelma.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VENCLEXTA

Products Affected

- Venclexta
- Venclexta Starting Pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For AML, patient meets any of the following: 1) the patient is 60 years of age or older, OR 2) the requested drug will be used as a component of repeating the initial successful induction regimen if late relapse, OR 3) the patient has comorbidities that preclude use of intensive induction chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Mantle cell lymphoma

VENTAVIS

Products Affected

- Ventavis

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For pulmonary arterial hypertension (WHO Group 1), the diagnosis was confirmed by right heart catheterization. For new starts only, the patient must meet all of the following: 1) pretreatment mean pulmonary arterial pressure is greater than or equal to 25 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND 3) pretreatment pulmonary vascular resistance is greater than 3 Wood units.
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.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VERSACLOZ

Products Affected

- Versacloz

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VERZENIO

Products Affected

- Verzenio

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VIGABATRIN

Products Affected

- vigabatrin
- Vigadrone

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For complex partial seizures (CPS): patient had an inadequate response to at least 2 alternative therapies for CPS.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VITRAKVI

Products Affected

- Vitrakvi

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VIZIMPRO

Products Affected

- Vizimpro

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VORICONAZOLE

Products Affected

- voriconazole intravenous
- voriconazole oral suspension for reconstitution

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	The patient will be using the requested drug orally or intravenously.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Prophylaxis of invasive aspergillosis in a high-risk patient, empiric antifungal therapy for febrile neutropenia in a high-risk patient, pulmonary aspergillosis, oropharyngeal candidiasis, mycosis due to Scedosporium prolificans

VOSEVI

Products Affected

- Vosevi

PA Criteria	Criteria Details
Exclusion Criteria	Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh class B or C)
Required Medical Information	For chronic 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, liver and kidney transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current AASLD 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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

VOTRIENT

Products Affected

- Votrient

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3 years.
Other Criteria	Soft tissue sarcoma other than GIST [angiosarcoma, Pleomorphic rhabdomyosarcoma, retroperitoneal/intra-abdominal soft tissue sarcoma that is unresectable or progressive, soft tissue sarcoma of the extremity/superficial trunk or head/neck, including synovial sarcoma, or solitary fibrous tumor/hemangiopericytoma or alveolar soft part sarcoma], approve. Differentiated (ie, papillary, follicular, Hurthle) thyroid carcinoma, approve if the patient is refractory to radioactive iodine therapy. Uterine sarcoma, approve if the patient has recurrent, advanced or metastatic disease. Advanced Renal Cell Carcinoma, Clear Cell or non-Clear Cell histology-approved if the patient has relapsed or stage IV disease. Ovarian Cancer (ie, Epithelial Ovarian, Fallopian Tube, or Primary Peritoneal Cancer) - approve if the patient has persistent or recurrent disease. GIST - approve if the patient has tried TWO of the following: Gleevec, Sutent, or Stivarga. Medullary Thyroid Carcinoma, approve if the patient has tried vandetanib (Caprelsa) or cabozantinib (Cometriq).
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Differentiated (ie, papillary, follicular, Hurthle cell) thyroid carcinoma. Uterine sarcoma, Epithelial Ovarian, Fallopian Tube, or Primary Peritoneal Cancer, Gastrointestinal Stromal Tumor (GIST), Medullary thyroid carcinoma.

VRAYLAR

Products Affected

- Vraylar oral capsule
- Vraylar oral capsule,dose pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following: lurasidone, aripiprazole, olanzapine, paliperidone, quetiapine, risperidone, or ziprasidone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XALKORI

Products Affected

- Xalkori

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Recurrent anaplastic lymphoma kinase (ALK)-positive or ROS1-positive non-small cell lung cancer (NSCLC), NSCLC with high-level MET amplification or MET exon 14 skipping mutation, ALK- or ROS1-postive brain metastases from NSCLC, ALK-positive inflammatory myofibroblastic tumors (IMT), ALK-positive anaplastic large cell lymphoma (ALCL).

XELJANZ

Products Affected

- Xeljanz
- Xeljanz XR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For moderately to severely active rheumatoid arthritis (new starts only): Patient meets at least one of the following criteria: 1) Inadequate response, intolerance or contraindication to methotrexate (MTX), OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD). For active psoriatic arthritis (new starts only): Patient meets BOTH of the following criteria: 1) Inadequate response to MTX or other nonbiologic DMARDs OR a prior biologic DMARD, AND 2) The requested drug is used in combination with a nonbiologic DMARD. For moderately to severely active ulcerative colitis (new starts only): Patient meets at least one of the following criteria: 1) Inadequate response, intolerance or contraindication to at least one conventional therapy option (e.g., aminosalicylates), or 2) Inadequate response or intolerance to a prior biologic DMARD.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XGEVA

Products Affected

- Xgeva

PA Criteria	Criteria Details
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.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Systemic mastocytosis related osteopenia or osteoporosis

XIFAXAN

Products Affected

- Xifaxan oral tablet 550 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed to reduce the risk of overt hepatic encephalopathy (HE) recurrence OR 2) The patient has the diagnosis of irritable bowel syndrome with diarrhea (IBS-D) AND 3) If the patient has previously received treatment with the requested drug, the patient has experienced a recurrence of symptoms AND 4) The patient has not already received an initial 14-day course of treatment and two additional 14-day courses of treatment with the requested drug OR 5) The patient has not previously received treatment with the requested drug
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Reduction in risk of overt HE recurrence: 6 Months, IBS-D: 14 Days
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XOLAIR

Products Affected

- Xolair

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For allergic asthma initial therapy: 1) Patient has positive skin test (or blood test) to at least 1 perennial aeroallergen, 2) Patient has baseline IgE level greater than or equal to 30 IU/mL, and 3) Patient has inadequate asthma control despite current treatment with both of the following medications at optimized doses: a) Inhaled corticosteroid, and b) Additional controller (long acting beta2-agonist, leukotriene modifier, or sustained-release theophylline) unless patient has an intolerance or contraindication to such therapies. For allergic asthma continuation therapy only: Patient's asthma control has improved on treatment with the requested drug since initiation of therapy. For chronic idiopathic urticaria (CIU) initial therapy: 1) Patient has been evaluated for other causes of urticaria, including bradykinin-related angioedema and IL-1-associated urticarial syndromes (auto-inflammatory disorders, urticarial vasculitis), and 2) Patient has experienced a spontaneous onset of wheals, angioedema, or both, for at least 6 weeks. For CIU continuation therapy: Patient has experienced a response (e.g., improved symptoms) since initiation of therapy.
Age Restrictions	For CIU: 12 years of age or older. For allergic asthma: 6 years of age or older.
Prescriber Restrictions	N/A
Coverage Duration	Allergic asthma: Plan Year. CIU initial: 6 months. CIU continuation: Plan Year.
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XOSPATA

Products Affected

- Xospata

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XPOVIO

Products Affected

- Xpovio

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XTANDI

Products Affected

- Xtandi

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

XYREM

Products Affected

- Xyrem

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) The requested drug is being prescribed for the treatment of excessive daytime sleepiness in a patient 7 years of age or older with narcolepsy and 2) If the patient is 18 years of age or older, the patient experienced an inadequate treatment response, intolerance, or contraindication to at least one central nervous system (CNS) wakefulness promoting drug and at least one central nervous system (CNS) stimulant drug OR 3) If the patient is less than 18 years of age, the patient experienced an inadequate treatment response, intolerance, or contraindication to at least one central nervous system (CNS) stimulant drug (NOTE: Examples of a central nervous system (CNS) stimulant drug are amphetamine, dextroamphetamine, or methylphenidate. Example of a central nervous system (CNS) wakefulness promoting drug is armodafinil. Coverage of armodafinil or amphetamines may require prior authorization). OR 4) The requested drug is being prescribed for the treatment of cataplexy in a patient 7 years of age or older with narcolepsy
Age Restrictions	7 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	If the request is for a continuation of therapy, then the patient experienced a decrease in daytime sleepiness with narcolepsy or a decrease in cataplexy episodes with narcolepsy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ZARXIO

Products Affected

- Zarxio

PA Criteria	Criteria Details
Exclusion Criteria	Use of the requested product within 24 hours prior to or following chemotherapy or radiotherapy.
Required Medical Information	For prophylaxis or treatment of myelosuppressive chemotherapy-induced FN patients must meet all of the following: 1) Patient has a non-myeloid cancer, 2) Patient 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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Treatment of chemotherapy-induced febrile neutropenia (FN), following chemotherapy for acute lymphocytic leukemia (ALL), stem cell transplantation related indications, myelodysplastic syndromes (MDS), agranulocytosis, aplastic anemia, HIV-related neutropenia, neutropenia related to renal transplant.

ZEJULA

Products Affected

- Zejula

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ZELBORAF

Products Affected

- Zelboraf

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For brain metastases with melanoma, all of the following criteria must be met: 1) The tumor is positive for BRAF V600 activating mutation (e.g., BRAF V600E or V600K mutation), and 2) The requested drug will be used in combination with cobimetinib. For non-small cell lung cancer, tumor is positive for the BRAF V600E mutation. For thyroid carcinoma, tumor is positive for BRAF mutation. For rectal cancer, tumor is positive for the BRAF V600E mutation. For colon cancer, tumor is positive for the BRAF V600E mutation.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Brain metastases with melanoma, non-small cell lung cancer, hairy cell leukemia, thyroid carcinoma (papillary carcinoma, follicular carcinoma, and Hurthle cell carcinoma), rectal cancer, and colon cancer.

ZIEXTENZO

Products Affected

- Ziextenzo

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	Cancer patients receiving chemotherapy, if prescribed by or in consultation with an oncologist or hematologist.
Coverage Duration	Cancer pts receiving chemo-6 mo
Other Criteria	Cancer patients receiving chemotherapy, approve if-the patient is receiving myelosuppressive anti-cancer medications that are associated with a high risk of febrile neutropenia (the risk is at least 20% based on the chemotherapy regimen), OR the patient is receiving myelosuppressive anti-cancer medications that are associated with a risk of febrile neutropenia but the risk is less than 20% based on the chemotherapy regimen and the patient has one or more risk factors for febrile neutropenia according to the prescribing physician (eg, aged greater than or equal to 65 years, prior chemotherapy or radiation therapy, persistent neutropenia, bone marrow involvement by tumor, recent surgery and/or open wounds, liver and/or renal dysfunction, poor performance status or HIV infection, OR the patient has had a neutropenic complication from prior chemotherapy and did not receive prophylaxis with a colony stimulating factor and a reduced dose or frequency of chemotherapy may compromise treatment.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ZOLINZA

Products Affected

- Zolinza

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Mycosis fungoides, Sezary syndrome.

ZYDELIG

Products Affected

- Zydelig

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Relapsed or refractory chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL), refractory or progressive follicular lymphoma, and marginal zone lymphomas [nodal marginal zone lymphoma, gastric mucosa associated lymphoid tissue (MALT) lymphoma, non-gastric MALT lymphoma, and splenic marginal zone lymphoma].

ZYKADIA

Products Affected

- Zykadia oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For NSCLC, patient has recurrent or metastatic ALK-positive or ROS1-positive disease. For inflammatory myofibroblastic tumor, the tumor is ALK-positive. For brain metastases, patient has ALK-positive NSCLC.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Plan Year
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Anaplastic lymphoma kinase (ALK)-positive inflammatory myofibroblastic tumor, recurrent ALK-positive non-small cell lung cancer (NSCLC), metastatic or recurrent ROS1-positive NSCLC, brain metastases from NSCLC.

ZYPREXA RELPREVV

Products Affected

- Zyprexa Relprevv intramuscular suspension for reconstitution 210 mg

PA Criteria	Criteria Details
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
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

PART B VERSUS PART D

Products Affected

- Abelcet
- acetylcysteine
- acyclovir sodium intravenous solution
- albuterol sulfate inhalation solution for nebulization 0.63 mg/3 mL, 1.25 mg/3 mL, 2.5 mg /3 mL (0.083 %), 2.5 mg/0.5 mL
- AmBisome
- Aminosyn-PF 7 % (sulfite-free)
- amphotericin B
- aprepitant
- azathioprine
- Brovana
- budesonide inhalation suspension for nebulization 0.25 mg/2 mL, 0.5 mg/2 mL
- calcitriol oral
- cinacalcet oral tablet 30 mg, 60 mg, 90 mg
- Clinimix 5%/D15W Sulfite Free
- Clinimix 4.25%/D10W Sulf Free
- Clinimix 4.25%/D5W Sulfit Free
- Clinimix 5%-D20W(sulfite-free)
- Clinisol SF 15 %
- cromolyn inhalation
- cyclophosphamide oral capsule
- cyclosporine modified
- cyclosporine oral capsule
- dronabinol
- Emend oral suspension for reconstitution
- Engerix-B (PF) intramuscular syringe
- Engerix-B Pediatric (PF) intramuscular syringe
- everolimus (immunosuppressive)
- Gengraf oral capsule 100 mg, 25 mg
- Gengraf oral solution
- granisetron HCl oral
- heparin (porcine) injection solution
- Hepatamine 8%
- Intralipid intravenous emulsion 20 %
- Intralipid intravenous emulsion 30 %
- Intron A injection
- ipratropium bromide inhalation
- ipratropium-albuterol
- levalbuterol HCl
- levocarnitine (with sugar)
- levocarnitine oral tablet
- methotrexate sodium (PF) injection solution
- methotrexate sodium injection
- methylprednisolone oral tablet
- mycophenolate mofetil
- mycophenolate sodium
- Nephramine 5.4 %
- Nutrilipid
- ondansetron
- ondansetron HCl oral
- paricalcitol oral
- pentamidine inhalation
- Plenamine
- Prednisone Intensol
- prednisone oral tablet
- Premasol 10 %
- Procalamine 3%
- Prograf oral granules in packet
- Prosol 20 %
- Recombivax HB (PF) intramuscular suspension 10 mcg/mL, 40 mcg/mL
- Recombivax HB (PF) intramuscular syringe
- Sandimmune oral solution
- sirolimus
- tacrolimus oral
- Travasol 10 %
- TrophAmine 10 %
- Xatmep
- Zortress

Details

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

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