ACITRETIN

Products Affected

- acitretin

PA Criteria	Criteria Details
Exclusion Criteria	Severely impaired liver or kidney function. Chronic abnormally elevated blood lipid values. Concomitant use of methotrexate or tetracyclines. Pregnancy. Females of child-bearing potential who intend to become pregnant during therapy or at any time for at least 3 years after discontinuing therapy. Females of child-bearing potential who will not use reliable contraception while undergoing treatment and for at least 3 years following discontinuation. Females of child-bearing potential who drink alcohol during treatment or for two months after cessation of therapy
Required Medical Information	Diagnosis for severe, recalcitrant psoriasis (including plaque, guttate, erythrodermic palmar- plantar and pustular) AND patient must have tried and failed, contraindication or intolerance to one formulary first line agent (e.g., Topical Corticosteroids (betamethasone, fluocinonide, desoximetasone),Topical Calcipotriene/Calcitriol, Topical Calcipotriene, OR Topical Tazarotene)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ACTIMMUNE

Products Affected

- Actimmune

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Chronic granulomatous disease for use in reducing the frequency and severity of serious infections associated with chronic granulomatous disease, or B.) Severe, malignant osteopetrosis (SMO)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ADCIRCA

Products Affected

- Adcirca
- Alyq

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use of organic nitrate or guanylate cyclase stimulators (includes intermittent use)
Required Medical Information	Diagnosis of Pulmonary arterial hypertension (PAH) confirmed by right heart catheterization or Doppler echocardiogram if patient is unable to undergo a right heart catheterization AND Patient has tried and had an insufficient response to a combination of therapy (e.g. tadalafil and ambrisentan) .
Age Restrictions	18 years and older
Prescriber Restrictions	N/A
Coverage Duration	PAH: Initial: 6 months. Reauth: 12 months.
Other Criteria	PAH (Reauth): Documentation of positive clinical response to therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ADEMPAS

Products Affected

- Adempas

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant administration with nitrates or nitric oxide donors (such as amyl nitrate) in any form, B.) Concomitant administration with phosphodiesterase inhibitors, including specific PDE-5 inhibitors (such as sildenafil, tadalafil, or vardenafil) or non-specific PDE inhibitors (such as dipyridamole or theophylline), C.) Pregnancy, or D.) Patients with pulmonary hypertension associated with idiopathic interstitial pneumonia
Required Medical Information	Diagnosis of one of the following A.) Pulmonary arterial hypertension (WHO group I) and diagnosis was confirmed by right heart catheterization, or B.) Chronic thromboembolic pulmonary hypertension (CTEPH, WHO group 4) and patient has persistent or recurrent disease after surgical treatment (e.g., pulmonary endarterectomy) or has CTEPH that is inoperable (Female patients must be enrolled in the ADEMPAS REMS program)
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AFINITOR

Products Affected

- 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, Nexavar).Tuberous sclerosis complex (TSC) Associated subependymal

PA Criteria	Criteria Details
	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

AFINITOR DISPERZ

Products Affected

- Afinitor Disperz oral tablet for suspension
  2 mg, 3 mg, 5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC) that requires therapeutic intervention but patient is not a candidate for curative surgical resection, or B.) Diagnosis of tuberous sclerosis complex- associated partial-onset seizures
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AJOVY

Products Affected

- Ajovy Autoinjector
- Ajovy Syringe

PA Criteria	Criteria Details
Exclusion Criteria	Combination therapy with Aimovig, Vyepti or Emgality
Required Medical Information	Diagnosis, number of migraine headaches per month, prior therapies tried
Age Restrictions	18 years and older
Prescriber Restrictions	N/A
Coverage Duration	1 year
Other Criteria	Approve if the patient meets the following criteria (A and B): A) Patient has greater than or equal to 4 migraine headache days per month (prior to initiating a migraine-preventative medication), AND B) Patient has tried at least two standard prophylactic pharmacologic therapies, at least one drug each from a different pharmacologic class (e.g., anticonvulsant, beta-blocker), and has had inadequate responses to those therapies or the patient has a contraindication to other prophylactic pharmacologic therapies according to the prescribing physician.
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	Diagnosis of metastatic anaplastic lymphoma kinase positive non-small cell lung cancer as detected by an FDA approved test.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ALPHA1 PROTEINASE INH

Products Affected

- Prolastin-C intravenous recon soln

PA Criteria	Criteria Details
Exclusion Criteria	Not covered for patients with IgA deficiency
Required Medical Information	Diagnosis of alpha-1 proteinase inhibitor (alpha-1 antitrypsin) deficiency in adult patients with emphysema
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ALUNBRIG

Products Affected

- Alunbrig oral tablet 180 mg, 30 mg, 90 mg
- Alunbrig oral tablets,dose pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC) following failure (secondary to resistance or intolerance) of prior crizotinib therapy.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AMBRISENTAN

Products Affected

- ambrisentan

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Pregnancy, or B.) Idiopathic pulmonary fibrosis (IPF), including those with pulmonary hypertension
Required Medical Information	Diagnosis of pulmonary arterial hypertension that was confirmed by right heart catheterization or Doppler echocardiogram if patient is unable to undergo a right heart catheterization (e.g., patient is frail, elderly, etc.) and patient has WHO Group I PAH
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

APOKYN

Products Affected

- APOKYN

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with 5-HT(3) receptor antagonists (eg. ondansetron, granisetron, dolasetron, palonosetron, alosetron etc)
Required Medical Information	Diagnosis of Parkinson's disease (PD) and patient is experiencing acute intermittent hypomobility (defined as off episodes characterized by muscle stiffness, slow movements, or difficulty starting movements)
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ARCALYST

Products Affected

- Arcalyst

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cryopyrin-associated periodic syndromes (CAPS), including familial cold autoinflammatory syndrome (FCAS) and Muckle-Wells Syndrome (MWS)
Age Restrictions	12 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ARIKAYCE

Products Affected

- Arikayce

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of pulmonary Mycobacterium avium complex (MAC) infection and used as part of a combination antibacterial regimen in treatment refractory patients
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist or pulmonologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AURYXIA

Products Affected

- Auryxia

PA Criteria	Criteria Details
Exclusion Criteria	A.) Iron overload syndrome (e.g. hemochromatosis), or B.) Iron replacement in patients with iron deficiency anemia with chronic kidney disease not on dialysis.
Required Medical Information	Diagnosis of hyperphosphatemia in patients with chronic kidney disease (CKD) on dialysis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a hematologist or nephrologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AUSTEDO

Products Affected

- Austedo

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Suicidal ideation and/or untreated or inadequately treated depression, B.) Hepatic impairment, C.) Taking MAOIs, reserpine, or tetrabenazine
Required Medical Information	Diagnosis of one of the following A.) Chorea associated with Huntington's disease (Huntington's chorea), or B.) Tardive dyskinesia
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a psychiatrist or neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

AYVAKIT

Products Affected

- Ayvakit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of unresectable or metastatic gastrointestinal stromal tumor, with a platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation, including PDGFRA D842V mutations
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	This criteria applies to new starts only
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

BALVERSA

Products Affected

- Balversa oral tablet 3 mg, 4 mg, 5 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of locally advanced or metastatic urothelial carcinoma with susceptible FGFR3 or FGFR2 genetic alterations and patient has progressed during or following at least one line of prior platinum-containing chemotherapy, including within 12 months of neoadjuvant or adjuvant platinum-containing chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

BETASERON

Products Affected

- Betaseron subcutaneous kit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., relapsing-remitting MS, clinically isolated syndrome, progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis, or C.)Secondary progressive disease with relapses or evidence of new brain lesions
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

BOSULIF

Products Affected

- Bosulif oral tablet 100 mg, 400 mg, 500 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Philadelphia chromosome-positive (Ph+) CML with resistance, relapse, or inadequate response to prior therapy with one of the following tyrosine kinase inhibitors (TKI): Gleevec [imatinib], Tasigna [nilotinib], or Sprycel [dasatinib], or B.) newly diagnosed chronic phase Philadelphia chromosome-positive chronic myelogenous leukemia (Ph + CML).
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

BRAFTOVI

Products Affected

- Braftovi oral capsule 75 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	One of the following: A.) Diagnosis of unresectable or metastic melanoma with documented BRAF V600E or V600K mutation as detected by an FDA-approved test AND used in combination with binimetinib or B.) Diagnosis of metastatic colorectal cancer with documented BRAF V600E mutation as detected by an FDA-approved test, after prior therapy. Used in combination with cetuximab.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

BRUKINSA

Products Affected

- Brukinsa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of A.) Mantle Cell Lymphoma (MCL) and patient has tried one prior therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CABLIVI

Products Affected

- Cablivi injection kit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acquired thrombotic thrombocytopenic purpura (aTTP) and used in combination with plasma exchange and immunosuppression therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a hematologist or oncologist
Coverage Duration	3 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CABOMETYX

Products Affected

- Cabometyx

PA Criteria	Criteria Details
Exclusion Criteria	Patients who have or are at risk for severe hemorrhage and/or patients with a recent history of bleeding or hemoptysis
Required Medical Information	Diagnosis of one of the following A.) Advanced renal cell carcinoma, or B.) Advanced hepatocellular carcinoma AND patient has been previously treated with sorafenib.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CALQUENCE

Products Affected

- Calquence

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) MANTLE CELL LYMPHOMA (MCL) and patient has tried one other therapy, B.) Chronic lymphocytic leukemia, or C.) Small lymphocytic lymphoma
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CAPRELSA

Products Affected

- Caprelsa oral tablet 100 mg, 300 mg

PA Criteria	Criteria Details
Exclusion Criteria	Congenital long QT syndrome
Required Medical Information	Diagnosis of symptomatic or progressive medullary thyroid cancer in patients with unresectable, locally advanced, or metastatic disease.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CARBAGLU

Products Affected

- Carbaglu

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) N-acetyl glutamate synthase (NAGS) deficiency AND patient has either acute or chronic hyperammonemia
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CAYSTON

Products Affected

- Cayston

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis confirmed by appropriate diagnostic or genetic testing AND confirmation of P. aeruginosa in cultures of the airways
Age Restrictions	7 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CINACALCET

Products Affected

- cinacalcet oral tablet 30 mg, 60 mg, 90 mg

PA Criteria	Criteria Details
Exclusion Criteria	A.) Hypocalcemia (calcium less than 8.0 mg/dL)) or B.) Patients with chronic kidney disease who are not receiving dialysis.
Required Medical Information	Supporting statement from the prescriber that Cinacalet is being used to treat hypercalcemia due to primary hyperparathyroidism, parathyroid carcinoma, or kidney transplant. B vs D coverage determination required for Medicare Part B covered indications (i.e. patients with chronic kidney disease on dialysis).
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

COMETRIQ

Products Affected

- Cometriq oral capsule 100 mg/day(80 mg x1-20 mg x1), 140 mg/day(80 mg x1-20 mg x3), 60 mg/day (20 mg x 3/day)

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following a.) Gastrointestinal perforation, B.) Fistula, or C.) Severe hemorrhage
Required Medical Information	Diagnosis of progressive, metastatic medullary thyroid cancer.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

COPAXONE

Products Affected

- Copaxone subcutaneous syringe 20 mg/mL, 40 mg/mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., relapsing-remitting MS or progressive-relapsing MS, clinically isolated MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis, or C.)active secondary progressive disease with relapses or evidence of new brain lesions
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

COPIKTRA

Products Affected

- Copiktra

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of relapsed or refractory (with history of 2 prior therapies) of one of the following A) chronic lymphocytic leukemia, B) small lymphocytic lymphoma, or C) follicular lymphoma
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

CORLANOR

Products Affected

- Corlanor

PA Criteria	Criteria Details
Exclusion Criteria	Decompensated acute heart failure, hypotension (i.e. blood pressure less than 90/50 mmHg), sick sinus syndrome, sinoatrial block, or 3rd degree AV block, unless a functioning demand pacemaker is present
Required Medical Information	Diagnosis of one of the following A.) stable, symptomatic chronic heart failure with left ventricular ejection fraction 35% or less, who are in sinus rhythm and either are on maximally tolerated doses of beta-blockers or have a contraindication to beta-blocker use, or B.) stable, symptomatic heart failure due to dilated cardiomyopathy in patients who are in sinus rhythm with an elevated heart rate.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

COSENTYX

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Plaque psoriasis (Initial): Diagnosis of moderate to severe plaque psoriasis. One of the following: Failure, contraindication, or intolerance to Enbrel (etanercept) AND Humira (adalimumab), OR for continuation of prior Cosentyx therapy. Psoriatic Arthritis (PsA) (Initial): Diagnosis of active PsA. One of the following: Failure, contraindication, or intolerance to both Enbrel (etanercept) and Humira (adalimumab), OR for continuation of prior Cosentyx therapy. Ankylosing Spondylitis (AS) (Initial): Diagnosis of active AS. One of the following: Failure, contraindication, or intolerance to both Enbrel (etanercept) and Humira (adalimumab), OR for continuation of prior Cosentyx therapy. All indications (Initial, reauth): Patient is not receiving Cosentyx in combination with a biologic DMARD [eg, Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Simponi (golimumab)]. Patient is not receiving Cosentyx in combination with a Janus kinase inhibitor [eg, Xeljanz (tofacitinib)]. For a diagnosis of PsA or plaque psoriasis, Patient is not receiving Cosentyx in combination with a phosphodiesterase 4 (PDE4) inhibitor [e.g., Otezla (apremilast)].
Age Restrictions	N/A
Prescriber Restrictions	Plaque psoriasis (Initial): Prescribed by or in consultation with a dermatologist. PsA (Initial): Prescribed by or in consultation with a rheumatologist or dermatologist. AS (Initial): Prescribed by or in consultation with a rheumatologist.
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

COTELLIC

Products Affected

- Cotellic

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of unresectable or metastatic malignant melanoma with BRAF V600E OR V600K mutation, and documentation of combination therapy with vemurafenib (Zelboraf)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	Known serious hypersensitivity to penicillamine or cysteamine
Required Medical Information	Diagnosis of nephropathic cystinosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

DALFAMPRIDINE

Products Affected

- dalfampridine

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) History of seizure. B.) Moderate or severe renal impairment (creatinine clearance less than or equal to 50 mL/minute)
Required Medical Information	Diagnosis of multiple sclerosis and patient must demonstrate sustained walking impairment, but with the ability to walk 25 feet (with or without assistance) prior to starting dalfampridine
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

DAURISMO

Products Affected

- Daurismo

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of newly diagnosed acute myeloid leukemia (AML) and used in combination with low-dose cytarabine in patients 75 years of age or older or who have comorbidities that preclude use of intensive induction chemotherapy
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

DEFERASIROX

Products Affected

- deferasirox

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Creatinine clearance less than 40 mL/min, B.) Poor performance status, C.) Platelet count less than 50 x 10(9)/L, D.) Advanced malignancy, E.) High-risk myelodysplastic syndrome (MDS)
Required Medical Information	Diagnosis of one of the following A.) Chronic iron overload in patients with non-transfusion-dependent thalassemia syndromes who have liver iron concentrations of at least 5 mg Fe/g dry weight AND serum ferritin level greater than 300 mcg/L, or B.) Chronic iron overload due to blood transfusions (transfusion hemosiderosis) as evidenced by transfusion of at least 100 mL/kg packed red blood cells AND serum ferritin level greater than 1000 mcg/L
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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

ELIGARD

Products Affected

- Eligard
- Eligard (3 month)
- Eligard (4 month)
- Eligard (6 month)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of advanced or metastatic prostate cancer
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ENBREL

Products Affected

- Enbrel
- Enbrel Mini
- Enbrel SureClick

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe rheumatoid arthritis, B.) Moderate to severe polyarticular juvenile idiopathic arthritis, C.) Psoriatic arthritis, D.) Ankylosing spondylitis, or E.) Moderate to severe chronic plaque psoriasis in patients who are candidates for systemic therapy or phototherapy. Screening for latent tuberculosis infection is required prior to initiation of treatment.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ENDARI

Products Affected

- Endari

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acute sickle cell disease AND patient must have trial history of Hydroxyurea. Otherwise Endari requires documentation of (1) history of inadequate treatment with Hydroxyurea OR (2) history of adverse event with Hydroxyurea OR (3) Hydroxyurea is contraindicated.
Age Restrictions	5 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ENTRESTO

Products Affected

- Entresto

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) History of angioedema related to previous ACE inhibitor or ARB therapy, B.) Concomitant use or use within 36 hours of ACE inhibitors, or C.) Concomitant use of aliskiren in patients with diabetes
Required Medical Information	Diagnosis of one of the following A.) Chronic heart failure, NYHA Class II to IV, or B.) Symptomatic heart failure with systemic left venticular systolic dysfunction
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

EPIDIOLEX

Products Affected

- Epidiolex

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Lennox-Gastaut syndrome, or B.) Severe myoclonic epilepsy in infancy (Dravet syndrome), or C.) Tuberous sclerosis complext (TSC)
Age Restrictions	Patients 1 year and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ERIVEDGE

Products Affected

- Erivedge

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: A.) Metastatic basal cell carcinoma, B.) Locally advanced basal cell carcinoma that has recurred following surgery or the patient is not a candidate for surgery or radiation.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ERLEADA

Products Affected

- Erleada

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of A.) Nonmetastatic, castration-resistant prostate cancer or B.) Metastatic, castration-sensitive prostate cancer.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ERLOTINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) locally advanced, unresectable, or metastatic pancreatic cancer and erlotinib (Tarceva) will be used in combination with gemcitabine, or B.) locally advanced or metastatic non-small cell lung cancer with one of the following: 1.) failure with at least one prior chemotherapy regimen, 2.) no evidence of disease progression after four cycles of first-line platinum-based chemotherapy and Tarceva will be used as maintenance treatment, or 3.) Patient has known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutation as detected by an FDA-approved test or Clinical Laboratory Improvement Amendments-approved facility.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ESRD THERAPY

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pretreatment hemoglobin levels of less than 10g/dL. Dose reduction or interruption if hemoglobin exceeds 10 g/dL (CKD not on dialysis-adult, cancer), 11 g/dL (CKD on dialysis), 12 g/dL (pediatric CKD) in addition to supporting statement of diagnosis from physician.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

FARYDAK

Products Affected

- Farydak

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Multiple Myeloma and used in combination with Velcade (bortezomib) and dexamethasone. Patient has received at least two prior treatment regimens which included both of the following: Velcade (bortezomib) and an immunomodulatory agent [eg, Revlimid (lenalidomide), Thalomid (thalidomide)].
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

FENTANYL ORAL

Products Affected

- fentanyl citrate buccal lozenge on a handle

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Management of acute or postoperative pain (including headache/migraine, dental pain, and use in the emergency room), B.) Use in opioid non-tolerant patients
Required Medical Information	Patient meets the following: A) Diagnosis of cancer and use is for breakthrough cancer pain, B) Must have tried and failed at least two of the following alts: MORPHINE, HYDROMORPHONE, OXYMORPHONE, APAP/CODEINE, OXYCODODONE/APAP, OXYCODONE, HYDROCODONE/APAP), C) other formulary short-acting strong narcotic analgesic alternatives (other than fentanyl) have been ineffective, not tolerated, or contraindicated, D) prescriber and patient are registered in the Transmucosal Immediate Release Fentanyl (TIRF) Risk Evaluation and Mitigation Strategy Access program
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

FERRIPROX

Products Affected

- deferiprone
- Ferriprox oral solution
- Ferriprox oral tablet 1,000 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Serum ferritin level
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hematologist
Coverage Duration	1 year
Other Criteria	Initial therapy - approve if prior to starting therapy the serum ferritin level was greater than 2,500 mcg/L. Continuation therapy - approve is the patient is benefiting from therapy as confirmed by the prescribing physician.
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

FIRDAPSE

Products Affected

- Firdapse

PA Criteria	Criteria Details
Exclusion Criteria	History of seizures
Required Medical Information	Diagnosis of Lambert-Eaton myasthenic syndrome
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GALAFOLD

Products Affected

- Galafold

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Fabry disease with an amenable galactosidase alpha gene (GLA) mutation
Age Restrictions	16 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GATTEX

Products Affected

- Gattex 30-Vial

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of short bowel syndrome and patient is dependent on parenteral support.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GAVRETO

Products Affected

- Gavreto

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	NSCLC-18 years and older, MTC/thyroid cancer-12 years and older
Prescriber Restrictions	N/A
Coverage Duration	3 years
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. Medullary thyroid cancer (MTC)-approve if the patient has advanced or metastatic rearranged during transfection (RET)-mutant disease and the disease requires treatment with systemic therapy. Thyroid cancer (other than MTC)-approve if the patient has advanced or metatstatic rearranged during transfection (RET) fusion-positive disease, the disease is radioactive iodine-refractory AND the disease requires treatment with systemic therapy.
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	Any of the following A.) Recent (within the last 6 months) occurrence of: myocardial infarction, unstable angina, stroke, transient ischemic attack, decompensated heart failure requiring hospitalization, or Class III/IV heart failure, B.) History or presence of Mobitz Type II 2nd degree or 3rd degree AV block or sick sinus syndrome, unless patient has a pacemaker, C.) Baseline QTC interval greater than or equal to 500 milliseconds, D.) Receiving concurrent treatment with Class Ia or Class III anti-arrhythmic drugs (quinidine, procainamide, amiodarone, sotalol)
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., relapsing-remitting MS, clinically isolated syndrome, progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis, or C.) Active secondary progressive disease with relapses or evidence of new brain lesions
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GILOTRIF

Products Affected

- Gilotrif

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Metastatic non-small cell lung cancer (NSCLC) whose tumors have non-resistant epidermal growth factor receptor (EGFR) mutations as detected by an FDA-approved test, or B.) Metastatic squamous NSCLC, progressing after platinum-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GOCOVRI

Products Affected

- Gocovri oral capsule,extended release 24hr 137 mg, 68.5 mg

PA Criteria	Criteria Details
Exclusion Criteria	Patients with end-stage reanl disease (ESRD, CrCl below 15 ml/min/1.73 m2)
Required Medical Information	Diagnosis of one of the following A.) Parkinsons disease and patient is experiencing dyskinesia, receiving levodopa based therapy, and has documented trial and failure to amantadine immediate release, or B.) Extrapyramidal disease and has documented trial and failure to amantadine immediate release
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

GROWTH HORMONES

Products Affected

- Norditropin FlexPro

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

PA Criteria	Criteria Details
	child/adult and cont tx in adults), SHOX (initial), SGA (initial) - prescribed 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

PA Criteria	Criteria Details
	nutritional support.Cont tx - 2nd course if pt responded to tx with a 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, renal function impairment with growth failure, short bowel syndrome, and short-stature homeobox-containing gene (SHOX) deficiency

HEPATITIS B

Products Affected

- Vemlidy

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of chronic hepatitis B and all of the following: 1.) Patient has evidence of viral replication, 2.) Patient has evidence of persistent elevations in serum aminotransferase (ALT or AST) or histologically active disease, and 3.) Patient is receiving anti-retroviral therapy if the patient has HIV co-infection
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, hepatologist, or infectious disease specialist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

HEPATITIS C

Products Affected

- Epclusa oral tablet 400-100 mg
- Harvoni oral tablet 90-400 mg
- Vosevi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Must submit documentation of HCV genotype, subtype and quantitative HCV RNA (viral load) testing any time prior to therapy. Must document the following within 12 weeks of starting therapy, (1) CBC, INR, hepatic function panel and GFR. Must document cirrhosis status, prior treatment history (if any), and planned duration of treatment. FOR GENOTYPE 1,4,5,6 : Must include, trial/failure, contraindication to, or intolerance to Harvoni prior to approval of Epclusa or Vosevi. FOR GENOTYPE 2,3 : Must include, trial/failure, contraindication to, or intolerance to Epclusa prior to approval of Vosevi.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with gastroenterologist, hepatologist, or infectious disease specialist
Coverage Duration	Duration of approval per AASLD Guidelines
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

HRM - ONCOLOGY

Products Affected

- megestrol oral suspension 400 mg/10 mL (40 mg/mL), 625 mg/5 mL (125 mg/mL)
- megestrol oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The drug is being prescribed for an FDA-approved indication AND If formulary non HRM alternatives for diagnosis of cachexia secondary to chronic illness (oxandrolone) considered safe and effective in the elderly are available, then the member had an inadequate response, intolerable side effect, or contraindication to at least 2 of the alternative(s) AND the prescribing physician attests to the medical necessity for using this high risk medication and intent to monitor for side effects, AND anticipated treatment course/duration. For treatment of cancer related diagnosis or endometrial hyperplasia, or endometriosis, requests will be automatically approved.
Age Restrictions	Automatic approval if member is less than 65 years of age. Prior Auth required for age 65 or older.
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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 40 mg/0.8 mL
- Humira(CF)
- Humira(CF) Pedi Crohns Starter subcutaneous syringe kit 80 mg/0.8 mL, 80 mg/0.8 mL-40 mg/0.4 mL
- Humira(CF) Pen Crohns-UC-HS
- Humira(CF) Pen Psor-Uv-Adol HS
- Humira(CF) Pen subcutaneous pen injector kit 80 mg/0.8 mL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe rheumatoid arthritis, B.) Moderate to severe polyarticular juvenile idiopathic arthritis, C.) Psoriatic arthritis, D.) Ankylosing spondylitis, E.) Moderate to severe chronic plaque psoriasis in patients who are candidates for systemic therapy or phototherapy and when other systemic therapies are medically less appropriate, F.) Moderate to severe Crohn's disease in patients who have had an inadequate response to conventional therapy (e.g. corticosteroids, methotrexate, mercaptopurine), G.) Moderate to severe ulcerative colitis in patients who have had an inadequate response to immunosuppressants or 5-ASA (e.g. corticosteroids, azathioprine, sulfasalazine, mesalamine), H.) Non-infectious uveitis (including intermediate, posterior, and panuveitis), or I.) Moderate to severe hidradenitis suppurativa. Screening for latent tuberculosis infection is required prior to initiation of treatment.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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 oral tablet 10 mg, 15 mg, 30 mg, 45 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Chronic phase, accelerated phase, or blast phase chronic myeloid leukemia (CML) in adult patients who are T315I-positive or for whom no other tyrosine kinase inhibitor therapy is indicated, or B.) Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL) in adult patients who are T315I-positive or for whom no other tyrosine kinase inhibitor therapy is indicated.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

IDHIFA

Products Affected

- Idhifa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of relapsed or refractory acute myeloid leukemia with an isocitrate dehydrogenase 2 mutation as detected by an FDA approved test
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

IMATINIB

Products Affected

- imatinib

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Philadelphia chromosome-positive chronic myelogenous leukemia (Ph+ CML), B.) Ph+ acute lymphoblastic leukemia (ALL), C.) Gastrointestinal tumor (GIST) where patient has documented c-KIT (CD117) positive unresectable or metastatic malignant GIST or patient had resection of c-KIT positive GIST and imatinib will be used as an adjuvant therapy, D.) Dermatofibrosarcoma protuberans that is unresectable, recurrent, or metastatic, E.) hypereosinophilic syndrome or chronic eosinophilic leukemia, F.) myelodysplastic syndrome or myeloproliferative disease associated with platelet-derived growth factor receptor gene re-arrangements, or G.) aggressive systemic mastocytosis without the D816V c-KIT mutation or with c-KIT mutational status unknown. Diagnosis of one of the following in a pediatric patient: A) Ph+ CML that is newly diagnosed in the chronic phase B) newly diagnosed Ph+ ALL
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

IMBRUVICA

Products Affected

- Imbruvica oral capsule 140 mg, 70 mg
- Imbruvica oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Mantle cell lymphoma (MCL) who have received at least one prior therapy, B.) Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL), C.) Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL) with 17p deletion, D.) Waldenstrom's macroglobulinemia (WM), E.) Marginal zone lymphoma who require systemic therapy and have received at least one prior anti-CD20-based therapy, or F.) Graft vs host disease after failure of a least one first-line corticosteroid therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

INCRELEX

Products Affected

- Increlex

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) active or suspected malignancy, B.) use for growth promotion in patients with closed epiphyses, C.) Intravenous administration
Required Medical Information	Diagnosis of one of the following A.) growth failure in children with severe primary IGF-1 deficiency, or B.) growth hormone (GH) gene deletion in children who have developed neutralizing antibodies to GH
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

INHALED TOBRAMYCIN

Products Affected

- Tobi Podhaler inhalation capsule, w/inhalation device

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Must meet all of the following A.) Diagnosis of cystic fibrosis and B.) Patient has evidence of P. aeruginosa in the lungs
Age Restrictions	6 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Diagnosis of (A) advanced renal cell carcinoma and patient failed one or more systemic therapies for renal cell carcinoma (e.g., sunitinib-, bevacizumab-, temsirolimus-, or cytokine-containing regimens) (B) for the first-line treatment of patients with advanced renal cell carcinoma (RCC), axitinib in combination with aveluma (C) for the first-line treatment of patients with advanced RCC axitinib in combination with pembrolizumab
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	Diagnosis of intermediate-2 or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis (MF).
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

INTRAROSA

Products Affected

- Intrarosa

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Vaginal bleeding or dysfunctional uterine bleeding of an undetermined origin, B.) Known or suspected estrogen-dependent neoplasia
Required Medical Information	Diagnosis of moderate to severe dyspareunia, due to vulvar and vaginal atrophy associated with menopause. Patient has tried and failed, has a contraindication or intolerance to a low dose vaginal estrogen preparation (e.g. Premarin vaginal cream, Yuvafem)
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	Initial: 3 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

INTRON-A

Products Affected

- Intron A injection

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Autoimmune hepatitis, B.) Decompensated liver disease
Required Medical Information	Diagnosis of one of the following A.) Hairy cell leukemia, B.) Diagnosis of condylomata acuminata involving external surfaces to the genital or perianal areas, C.) Diagnosis of AIDS-related Kaposi's sarcoma, D.) Clinically aggressive follicular lymphoma and the medication will be used concurrently with anthracycline-containing chemotherapy or is not a candidate for anthracycline-containing chemotherapy, E.)Malignant melanoma and the request for coverage is within 56 days of surgery and the patient is at high risk of disease recurrence, F.) Diagnosis of chronic hepatitis B with compensated liver disease and patient has evidence of hepatitis B viral replication and patient has been serum hepatitis B surface antigen-positive for at least 6 months, or G.) Diagnosis of chronic hepatitis C with compensated liver disease and is receiving combination therapy with ribavirin, unless ribavirin is contraindicated, and the medication will not be used as part of triple therapy with a protease inhibitor and patient has a clinical reason for not using peginterferon
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Condylomata: 3 mos, HBV: E antigen pos: 16 wks, E antigen neg: 48 wks, KS: 16 wks, Other: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

IRESSA

Products Affected

- Iressa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of metastatic Non-small cell lung cancer (NSCLC) and used as first- line therapy and patient has known active epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutations as detected by an FDA-approved test or Clinical Laboratory Improvement Amendments-approved facility
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ITRACONAZOLE

Products Affected

- itraconazole

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Ventricular dysfunction (e.g., congestive heart failure (CHF) or history of CHF), B.) Concurrent therapy with a CYP3A4 substrate (e.g., methadone, lovastatin, simvastatin, etc.)
Required Medical Information	Diagnosis of one of the following A.) Systemic fungal infection (e.g., aspergillosis, histoplasmosis, blastomycosis), B.) Onychomycosis confirmed by one of the following: positive potassium hydroxide (KOH) preparation, fungal culture, or nail biopsy, or C.) Candidiasis (esophageal or oropharyngeal) that is refractory to treatment with fluconazole (ORAL SOLUTION ONLY)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
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	Diagnosis of one of the following A.) Intermediate or high-risk myelofibrosis, including primary myelofibrosis, post-polycythemia vera myelofibrosis and post-essential thrombocythemia myelofibrosis, B.) Polycythemia vera in patients with an inadequate response to or are intolerant of hydroxyurea, C.) Acute graft versus host disease (GVHD), AND disease is refractory to steroid therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

KALYDECO

Products Affected

- Kalydeco

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis AND the patient has 1 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.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

KEVZARA

Products Affected

- Kevzara subcutaneous syringe

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of moderate to severe active Rheumatoid Arthritis (RA) and one of the following: A.) Trial and failure, contraindication, or intolerance to Enbrel (etanercept) or Humira (adalimumab) or B.) Attestation demonstrating a trial may be inappropriate or C.) For continuation of prior Kevzara therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	Any of the following A.) pregnancy, B.) coadministration with simvastatin, lovastatin, or CYP3A substrates with narrow therapeutic ranges, C.) concomitant treatment with systemic corticosteroids for serious medical conditions or illnesses, D.) history of unexplained vaginal bleeding, E.) endometrial hyperplasia with atypia or endometrial carcinoma
Required Medical Information	Diagnosis of hyperglycemia secondary to endogenous Cushing syndrome in patients with type 2 diabetes mellitus or glucose intolerance and patient had failed surgery or is not a candidate for surgery.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

KUVAN

Products Affected

- sapropterin

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of hyperphenylalaninemia (HPA) caused by tetrahydrobiopterin (BH4)-responsive phenylketonuria (PKU)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LENVIMA

Products Affected

- Lenvima

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Locally recurrent or metastatic, progressive, radioactive iodine-refractory differentiated thyroid cancer, B.) Advanced renal cell carcinoma following one prior anti-angiogenic therapy in combination with everolimus, C.) Unresectable liver carcinoma, first-line therapy or D.) Advanced endometrial carcinoma that is not microsatellite instability-high (MSI-H) or mismatch repair deficient (dMMR), in combination with pembrolizumab and patient has disease progression following prior systemic therapy and are not candidates for curative surgery or radiation.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LEUKINE

Products Affected

- Leukine injection recon soln

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with myelosuppressive chemotherapy or radiation or excessive (greater than or equal to 10%) leukemic myeloid blasts in bone marrow or peripheral blood
Required Medical Information	Diagnosis of one of the following A.) Patient has undergone allogeneic or autologous bone marrow transplant (BMT) and engraftment is delayed or failed, B.) Patient is undergoing autologous peripheral-blood progenitor cell transplant to mobilize progenitor cells for collection by leukapheresis, C.) Medication will be used for myeloid reconstitution after an autologous or allogeneic BMT, D.) Patient has acute myeloid leukemia and administration will be after completion of induction chemotherapy, E.) Hematopoietic subsyndrome of acute radiation syndrome (H-ARS) or F.) Autologous peripheral blood stem cell transplant, Following myeloablative chemotherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LEUPROLIDE

Products Affected

- leuprolide subcutaneous kit
- Lupron Depot
- Lupron Depot (3 month)
- Lupron Depot (4 month)
- Lupron Depot (6 Month)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Advanced or metastatic prostate cancer and patient has tried and failed Eligard, B.) Diagnosis of central precocious puberty and patient had early onset of secondary sexual characteristics (male: earlier than 9 years of age, female: earlier than 8 years of age) and advanced bone age of at least one year compared with chronological age and has undergone gonadotropin-releasing hormone agonist (GnRHa) testing with peak luteinizing hormone (LH) level above pre-pubertal range or random LH level in pubertal range and patient had the following diagnostic evaluations to rule out tumors, when suspected: diagnostic imaging of the brain (MRI or CT scan), Pelvic/testicular/adrenal ultrasound, Human chorionic gonadotropin levels, Adrenal steroids to rule out congenital adrenal hyperplasia, C.) The medication will be used for stimulation testing to confirm the diagnosis of central precocious puberty, D.) Management of endometriosis, or E.) Preoperative treatment of anemia due to uterine leiomyomata.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LIDOCAINE PATCH

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	Authorization will be for 12 months
Other Criteria	N/A
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Diabetic neuropathic pain, chronic back pain

LINEZOLID

Products Affected

- linezolid
- linezolid in dextrose 5%

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use of MAOIs or within 2 weeks of taking an MAOI, B.) Treatment of Gram-negative infections
Required Medical Information	Diagnosis of one of the following A.) Community acquired pneumonia, B.) Hospital-acquired pneumonia, C.) Vancomycin-resistant Enterococcus faecium infection, D.) Complicated skin and skin structure infections, or E.) Uncomplicated skin and skin structure infections
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	VRE: 4 weeks, Nosocomial or community acquired pneumonia: 3 weeks, All other indications: 2 weeks
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	Diagnosis of one of the following A.) Metastatic colorectal cancer, previously treated with fluoropyrimidine, oxaliplatin, and irinotecan-based regimens, an anti-VEGF therapy, and if RAS wild-type, an anti-EGFR therapy, B.) Metastatic esophagogastric cancer, Adenocarcinoma previously treated with at least 2 prior lines of chemotherapy that included a fluoropyrimidine, a platinum, either a taxane or irinotecan and if appropriate, HER2/neu-targeted therapy, or C.) Metastatic adenocarcinoma of the stomach previously treated with at least 2 prior lines of chemotherapy that included fluoropyrimidine, a platinum, either a taxane or irinotecan and if appropriate, HER2/neu-targeted therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

LORBRENA

Products Affected

- Lorbrena oral tablet 100 mg, 25 mg

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with strong CYP3A4 inducers
Required Medical Information	Diagnosis of metastatic, anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer with disease progression on either alectinib or ceritinib as the first ALK inhibitor for metastatic disease, or disease progression on crizotinib and at least one other ALK inhibitor for metastatic disease
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
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	Diagnosis of one of the following A.) HER2- negative, deleterious or suspected deleterious germline BRCA mutated metastatic breast cancer AND patient has been previously treated with chemotherapy in neoadjuvant, adjuvant, or metastatic setting, B.) Advanced ovarian cancer with known or suspected BRCA mutation as detected by an FDA-approved test AND patient has trial and failure, contraindication, or intolerance to 3 or more prior lines of chemotherapy, C.) Recurrent epithelial ovarian cancer, recurrent fallopian tube cancer, or recurrent primary peritoneal cancer AND used for maintenance treatment in patients who are in complete or partial response to platinum-based chemotherapy (e.g. cisplatin, carboplatin), or D.) Deleterious or suspected deleterious germline or somatic BRCA-mutated (gBRCAm or sBRCAm) epithelial ovarian, fallopian tube, or primary peritoneal cancer in patients with complete or partial response to first-line platinum-based chemotherapy, or E) Deleterious or suspected deleterious germline BRCA-mutated metastatic pancreatic adenocarcinoma and disease has not progressed on at least 16 weeks of a first-line platinum-based chemotherapy regimen F) Prostate cancer or G) ovarian cancer uses in combination with bevacizumab
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

MAYZENT

Products Affected

- Mayzent oral tablet 0.25 mg, 2 mg

PA Criteria	Criteria Details
Exclusion Criteria	(1) CYP2C9*3/*3 genotype, (2) In the last 6 months experienced myocardial infarction, unstable angina, stroke, TIA, decompensated heart failure requiring hospitalization, or Class III or IV heart failure, (3) Presence of Mobitz type II second-degree, third-degree AV block, or sick sinus syndrome, unless patient has a functioning pacemaker
Required Medical Information	Diagnosis of relapsing forms of multiple sclerosis (clinically isolated syndrome, relapsing-remitting disease, or secondary progressive disease)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

MEKINIST

Products Affected

- Mekinist oral tablet 0.5 mg, 2 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Locally advanced or metastatic anaplastic thyroid cancer (ATC) with BRAF V600E mutation and used in combination with dabrafenib and no satisfactory locoregional treatment options, B.) Malignant melanoma with lymph node involvement and following complete resection with BRAF V600E or V600K mutations and used in combination with dabrafenib, C.) Unresectable or metastatic malignant melanoma with BRAF V600E or V600K mutations and used in combination with dabrafenib or as monotherapy, or D.) Metastatic non-small cell lung cancer, with BRAF V600E mutation, in combination with dabrafenib.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

MEKTOVI

Products Affected

- Mektovi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of unresectable or metastatic malignant melonoma with documented BRAF V600E or V600K mutation as detected by an FDA approved test AND used in combination with encorafenib
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

MIGLUSTAT

Products Affected

- miglustat

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of mild to moderate type 1 Gaucher disease and patient is not a candidate for enzyme replacement therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

MYTESI

Products Affected

- Mytesi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-infectious diarrhea associated with HIV/AIDS in patients receiving anti-retroviral therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist or gastroenterologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NATPARA

Products Affected

- Natpara

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of hypoparathyroidism and used to control hypocalcemia
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NERLYNX

Products Affected

- Nerlynx

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Any of the following: A.) Diagnosis of early stage HER2- overexpressed breast cancer and used after completion of adjuvant trastuzumab based therapy OR B.) Advanced or metastatic HER2-positive breast cancer and patient has received 2 or more prior anti-HER2-based regimens in the metastatic setting, in combination with capecitabine
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NEUPOGEN

Products Affected

- Neupogen

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Chemotherapy induced febrile neutropenia (prophylaxis), B.) Severe chronic neutropenia, C.) Patient is undergoing autologous peripheral-blood progenitor cell transplant to mobilize progenitor cells for collection by leukapheresis, or D.) Hematopoietic subsyndrome of acute radiation syndrome (H-ARS)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NEXAVAR

Products Affected

- Nexavar

PA Criteria	Criteria Details
Exclusion Criteria	Squamous cell lung cancer being treated with carboplatin and paclitaxel
Required Medical Information	Diagnosis of one of the following A.) Advanced renal cell carcinoma, B.) Locally recurrent or metastatic, progressive, differentiated thyroid carcinoma that is refractory to radioactive iodine treatment, or C.) Unresectable hepatocellular carcinoma
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NINLARO

Products Affected

- Ninlaro

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of multiple myeloma and documentation of combination therapy with lenalidomide and dexamethasone, used in patients with history of 1 prior therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NOXAFIL

Products Affected

- Noxafil oral suspension
- posaconazole oral tablet,delayed release (DR/EC)

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant treatment with sirolimus, B.) Concomitant use of CYP3A4 substrates that prolong QT interval (pimozide, quinidine), C.) Concomitant use of HMG-CoA Reductase inhibitors primarily metabolized through CYP3A4, or D.) Concomitant use of ergot alkaloids
Required Medical Information	Diagnosis of one of the following A.) Oropharyngeal candidiasis, or B.) Patient is severely immunocompromised and requires prophylaxis of invasive aspergillosis or candidiasis due to high risk of infection
Age Restrictions	13 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NUBEQA

Products Affected

- Nubeqa

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of non-metastatic, castration-resistant prostate cancer
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NUEDEXTA

Products Affected

- Nuedexta

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) History of prolonged QT interval, congenital long QT syndrome or Torsades de pointes, B.) Heart failure, C.) Complete AV block without an implanted pacemaker or high risk of complete AV block, D.) Concomitant use with quinidine, quinine, mefloquine, or drugs that prolong QT interval and are metabolized by CYP2D6 (e.g., thioridazine, pimozide), E.) Concomitant use with MAOIs or within 14 days of MAOI therapy
Required Medical Information	Diagnosis of pseudobulbar affect
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Diagnosis of hallucinations and delusions associated with Parkinson disease psychosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

NUVIGIL

Products Affected

- armodafinil
- modafinil oral tablet 100 mg, 200 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Obstructive sleep apnea (OSA) confirmed by sleep lab evaluation, B.) Narcolepsy confirmed by sleep lab evaluation, or C.) Shift work sleep disorder
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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

ODOMZO

Products Affected

- Odomzo

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of locally advanced basal cell carcinoma of the skin and one of the following A.) Cancer has recurred following surgery or radiation therapy, B.) Patient is not a candidate for surgery or radiation therapy.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ONUREG

Products Affected

- Onureg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	18 years and older
Prescriber Restrictions	N/A
Coverage Duration	3 years
Other Criteria	AML - Approve if the patient meets the following criteria (both A and B): A)Following intensive induction chemotherapy, the patient achieves one of the following according to the prescriber (i or ii): i. First complete remission OR ii. First complete remission with incomplete blood count recovery AND B) Patient is not able to complete intensive curative therapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

OPSUMIT

Products Affected

- Opsumit

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of pulmonary arterial hypertension WHO group I AND diagnosis was confirmed by right heart catheterization AND female patients are enrolled in the OPSUMIT REMS program
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ORENCIA

Products Affected

- Orencia

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: A.) Moderate to severe adult rheumatoid arthritis OR B.) Moderate to severe Juvenile idiopathic arthritis OR C.) Psoriatic arthritis. Must document 1.) trial and failure, contraindication or intolerance to Humira or Enbrel OR 2.) continuation of prior Orencia therapy for approval.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Diagnosis of hereditary tyrosinemia type 1
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ORGOVYX

Products Affected

- Orgovyx

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	18 years and older
Prescriber Restrictions	N/A
Coverage Duration	3 years
Other Criteria	Prostate Cancer-approve if the patient has advanced disease. Note: Advanced disease is defined as disease that has spread to other parts of the body, beyond the prostate. It can also include patients with persistent prostate specific antigen (PSA) levels or rising PSA levels after radiotherapy or surgery. Metastatic disease is also considered as advanced disease.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ORKAMBI

Products Affected

- Orkambi oral granules in packet
- Orkambi oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis (CF) with documented homozygous F508del mutation confirmed by FDA-approved CF mutation test
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist or prescribing practitioner is from a CF center accredited by the Cystic Fibrosis Foundation
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

OSPHENA

Products Affected

- Osphena

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) undiagnosed abnormal genital bleeding, B.) known or suspected estrogen-dependent neoplasia, C.) active or history of DVT, D.) active or history of pulmonary embolism, E.) active or history of arterial thromboembolic disease F.) pregnancy
Required Medical Information	Diagnosis of one of the following A.) moderate to severe dyspareunia due to vulvar and vaginal atrophy associated with menopause, or B.) moderate to severe vaginal dryness due to vulvar and vaginal atrophy associated with menopause. Patient has tried and failed, has a contraindication or intolerance to a low dose vaginal estrogen preparation (e.g. Premarin vaginal cream, Yuvafem).
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

OXANDRIN

Products Affected

- oxandrolone

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Breast or prostate cancer in men, B.) Breast cancer in women with hypercalcemia, C.) Pregnancy, D.) Nephrosis or nephrotic phase of nephritis, E.) Hypercalcemia
Required Medical Information	Diagnosis one of the following and receiving treatment as an adjunct therapy to promote weight gain A.) Extensive surgery, B.) Chronic infections, C.) Severe trauma, or D.) Failure to gain or maintain at least 90% of ideal body weight without definite pathophysiologic reasons, E.) Chronic corticosteroid administration, F.) Bone pain associated with osteoporosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PCSK9 INHIBITOR

Products Affected

- Praluent Pen
- Repatha
- Repatha Pushtronex
- Repatha SureClick

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	PRALUENT: Must meet criteria #1, #2 or #3. REPATHA: Must meet criteria #1, #2, #3 or #4. 1.) Diagnosis of primary hyperlipidemia including heterozygous familial hypercholesterolemia (HeFH). 2.) Myocardial infarction prophylaxis, stroke prophylaxis, and to reduce risk of coronary revascularization in pts with established CVD. 3.) Diagnosis of clinical atherosclerotic cardiovascular disease (CVD) as defined as one of the following: a. acute coronary syndrome, b. history of myocardial infarction, c. stable/unstable angina, d. coronary or other arterial revascularization, e. stroke, f. transient ischemic stroke (TIA), g. peripheral arterial disease presumed to be atherosclerotic region. 4.) Primary hyperlipidemia homozygous familial hypercholesterolemia (HoFH) confirmed by genotyping OR diagnosis based on the following: a. History of untreated LDL-C greater than 500 mg/dL AND xanthoma before 10 years of age OR b. Documentation of HeFH in both parents. REQUIRED DOCUMENTATION FOR INITIAL THERAPY: A.) Baseline and current LDL-C, LDL-C greater than or equal to 70 mg/dL, AND used in combination with maximally tolerated high-intensity statin OR patient is statin intolerant and LDL-C greater than or equal to 70 mg/dL. FOR CONTINUING THERAPY: Will continue to be used in combination with maximally tolerated statin (unless statin intolerant).
Age Restrictions	Repatha: 13 years of age and older for diagnosis HoFM. Diagnosis CVD and HeFH: 18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist, endocrinologist, or lipid specialist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.

PA Criteria	Criteria Details
Off-Label Uses	N/A

PEGASYS

Products Affected

- Pegasys subcutaneous solution
- Pegasys subcutaneous syringe

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Autoimmune hepatitis or other autoimmune condition known to be exacerbated by interferon or B.) Uncontrolled depression
Required Medical Information	Diagnosis of one of the following A.) chronic hepatitis C and criteria applied consistent with current AASLD-IDSA guidance with compensated liver disease, or B.) chronic hepatitis B infection
Age Restrictions	Hepatitis B: 3 years of age and older. Hepatitis C: 5 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a hepatologist, gastroenterologist, or infectious disease specialist
Coverage Duration	HBV: 12 months, HCV: based on current AASLD guidelines
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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

PENNSAID

Products Affected

- Pennsaid topical solution in metered-dose pump

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of osteoarthritis
Age Restrictions	Nnoe
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PHENYLBUTYRATE

Products Affected

- sodium phenylbutyrate

PA Criteria	Criteria Details
Exclusion Criteria	Management of acute hyperammonemia
Required Medical Information	Diagnosis of urea cycle disorders involving deficiencies of carbamoylphosphate synthetase, ornithine transcarbamoylase, or argininosuccinic acid
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PIQRAY

Products Affected

- Piqray

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of hormone receptor (HR) positive, HER2-negative, PIK3CA-mutated, advanced or metastatic breast cancer and used in combination with fulvestrant for postmenopausal women, and men following progression on or after endocrine- based regimen.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

POMALYST

Products Affected

- Pomalyst

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Must meet all of the following A.) Disease multiple myeloma has progressed on or within 60 days of completion of the last therapy, B.) If female of reproductive potential ALL of the below: Two negative pregnancy tests obtained prior to initiating therapy with Pomalyst, monthly negative pregnancy tests during therapy, C.) Patient has been counseled about the use of 2 forms of reliable contraception before, during, and 1 month after discontinuing therapy with Pomalyst, D.) Patient assessment to determine if prophylactic aspirin or antithrombic treatment (warfarin, clopidogrel) will need to be taken to reduce the risk of VTE (embolism, stroke), and OR AIDS-related Kaposi sarcoma (KS) after failure of highly active antiretroviral therapy (HAART) or in patients with KS who are HIV-negative and E.) Registered and certified to be compliant with Pomalyst REMS program
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PULMONARY FIBROSIS

Products Affected

- Esbriet oral capsule
- Esbriet oral tablet 801 mg
- Ofev

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Esbriet: Diagnosis of idiopathic pulmonary fibrosis. Ofev: A.) Diagnosis of idiopathic pulmonary fibrosis OR B.) Systemic sclerosis-associated interstitial lung disease OR C.) Chronic fibrosing interstitial lung diseases (ILDs) with a progressive phenotype.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PULMOZYME

Products Affected

- Pulmozyme

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted 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

QUININE SULFATE

Products Affected

- quinine sulfate

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following: A.) Prolongation of QT interval, B.) Glucose-6-phosphate dehydrogenase deficiency, C.) Myasthenia gravis, D.) Known hypersensitivity to mefloquine or quinidine, E.) Optic neuritis, F.) Diagnosis of Blackwater fever
Required Medical Information	Diagnosis of one of the following A.) uncomplicated Plasmodium falciparum malaria, B.) uncomplicated Plasmodium vivax malaria, or C.) babesiosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	1 month
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

RAVICTI

Products Affected

- Ravicti

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of urea cycle disorders
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

REGRANEX

Products Affected

- Regranex

PA Criteria	Criteria Details
Exclusion Criteria	Known neoplasm at the site of application
Required Medical Information	Diagnosis of lower extremity diabetic neuropathic ulcers that extend into the subcutaneous tissue or beyond and have an adequate blood supply
Age Restrictions	16 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	6 months
Other Criteria	N/A
Indications	All Medically-accepted 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

REVATIO

Products Affected

- sildenafil (Pulmonary Arterial Hypertension) oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	Receiving nitrate therapy (includes intermittent use)
Required Medical Information	Diagnosis of pulmonary arterial hypertension (WHO Group I) that was confirmed by right heart catheterization.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

REVLIMID

Products Affected

- Revlimid

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Multiple myeloma and medication will be used in combination with dexamethasone, B.) Autologous hematopoietic stem-cell transplantation (HSCT) in multiple myeloma patients, C.) Transfusion-dependent anemia due to low- or intermediate-1-risk myelodysplastic syndrome (MDS) associated with a deletion 5q cytogenetic abnormality with or without additional cytogenetic abnormalities, or D.) Mantle cell lymphoma whose disease has relapsed or progressed after two prior therapies, one of which included bortezomib, E.) Follicular lymphoma and used in combination with rituximab, or F.) Marginal zone lymphoma and used in combination with rituximab
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

RINVOQ

Products Affected

- Rinvoq

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of moderate to severe rheumatoid arthritis and patient has had an inadequate response or intolerance to methotrexate.
Age Restrictions	18 years of age or older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ROZLYTREK

Products Affected

- Rozlytrek oral capsule 100 mg, 200 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: A) ROS1-positive metastatic non-small cell lung cancer (NSCLC), OR B) Solid tumors that 1) have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion without a known acquired resistance mutation, AND 2) are metastatic or where surgical resection is likely to result in severe morbidity, AND 3) have either progressed following treatment or have no satisfactory alternative therapy
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

RUBRACA

Products Affected

- Rubraca

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis for which Rubraca is being used. BRCA-mutation (germline or somatic) status. Other medications tried for the diagnosis provided
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Authorization will be for 3years
Other Criteria	Ovarian, Fallopian Tube or Primary Peritoneal Cancer-treatment-Approve if the patient meets the following criteria (i and ii): i.The patient has a BRCA-mutation (germline or somatic) as confirmed by an approved test, AND ii.The patient has progressed on two or more prior lines of chemotherapy. Maintenance Therapy of Ovarian, Fallopian tube or Primary peritoneal cancer-Approve if the patient is in complete or partial response after at least two platinum-based chemotherapy regimens. Castration-Resistant Prostate Cancer - Approve if the patient meets the following criteria (A, B, C, and D): A) The patient has metastatic disease that is BRCA-mutation positive (germline and/or somatic) AND B) The patient meets one of the following criteria (i or ii): i. The medication is used concurrently with a gonadotropin-releasing hormone (GnRH) analog OR ii. The patient has had a bilateral orchiectomy AND C) The patient has been previously treated with at least one androgen receptor-directed therapy AND D) The patient meets one of the following criteria (i or ii): i. The patient has been previously treated with at least one taxane-based chemotherapy OR ii. The patient is not a candidate or is intolerant to taxane-based chemotherapy.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

RUCONEST

Products Affected

- Ruconest

PA Criteria	Criteria Details
Exclusion Criteria	Known allergy to rabbits or rabbit-derived products (leporine protein hypersensitivity)
Required Medical Information	Diagnosis of Hereditary angioedema (HAE)
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hematologist, immunologist, or allergist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

RYDAPT

Products Affected

- Rydapt

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) treatment naive FLT3 mutation-positive acute myelogenous leukemia (AML) and must be used in combination with standard cytarabine and daunorubicin induction and cytarabine consolidation therapy, or B.) systemic mastocytosis or mast cell leukemia
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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

SANDOSTATIN

Products Affected

- octreotide acetate injection solution

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) acromegaly and patient has inadequate response to or is ineligible for surgery, radiation, or bromocriptine mesylate, B.) metastatic carcinoid syndrome, or C.) vasoactive intestinal peptide-secreting tumors (VIPomas) with associated diarrhea
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SIGNIFOR

Products Affected

- Signifor

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Cushing disease and patient has inadequate response to or is not a candidate for surgery
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Must meet all of the following A.) Diagnosis of pulmonary multidrug resistant tuberculosis (MDR-TB) and B.) used in combination with at least 3 other agents.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist
Coverage Duration	24 weeks
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SKYRIZI

Products Affected

- Skyrizi subcutaneous syringe kit

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of moderate to severe plaque psoriasis in adults who are candidates for systemic therapy or phototherapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SOLTAMOX

Products Affected

- Soltamox

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant coumarin-type anticoagulant therapy, B.) history of thromboembolic disease such as DVT or PE
Required Medical Information	Diagnosis of breast cancer and documentation of inability to swallow tablet formulation
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SOMATULINE

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	Diagnosis of one of the following A.) acromegaly in patient with inadequate response to or is ineligible for surgery or radiotherapy, B.) carcinoid syndrome, or C.) gastroenteropancreatic neuroendocrine tumors (GEP-NETs)
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SOMAVERT

Products Affected

- Somavert

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of acromegaly and patient has inadequate response to or is ineligible for surgery or radiation therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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 oral tablet 100 mg, 140 mg, 20 mg, 50 mg, 70 mg, 80 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Philadelphia chromosome-positive chronic myelogenous leukemia (Ph+ CML) that is newly diagnosed in the chronic phase, B.) Ph+ CML in chronic, accelerated, or lymphoid blast phase with resistance or intolerance to prior therapy, C.) Diagnosis of Ph+ acute lymphoblastic leukemia with resistance or intolerance to prior therapy, or D.) Newly diagnosed Ph+ acute lymphoblastic leukemia in combination with chemotherapy
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	Diagnosis of one of the following A.) moderate to severely active Crohn's disease and patient has trial and failure or intolerance or contraindication to Humira, B.) moderate to severe plaque psoriasis and patient has trial and failure or intolerance or contraindication to Humira or Enbrel, or C.) active psoriatic arthritis and patient has trial and failure or intolerance or contraindication to Humira or Enbrel, or D.) moderate to severely active ulcerative colitis and patient has trial and failure or intolerance or contraindication to Humira
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a rheumatologist or gastroenterologist or dermatologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

STIVARGA

Products Affected

- Stivarga

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) metastatic colorectal cancer in patients previously treated with ALL of the following per the indication: 1. (fluoropyrimidine-, oxaliplatin-, and irinotecan)-based chemotherapy 2. anti-VEGF bevacizumab 3. anti-EGFR panitumumab OR cetuximab (for KRAS mutation-negative patients only), B.) liver carcinoma in patients previously treated with sorafenib, or C.) locally advanced, unresectable or metastatic gastrointestinal stromal tumor (GIST) who have been previously treated with imatinib and sunitinib
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SUTENT

Products Affected

- Sutent

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) gastrointestinal stromal tumor after disease progression on or intolerance to imatinib, B.) pancreatic neuroendocrine tumors in a patient with unresectable, locally advanced, or metastatic disease, C.) advanced renal cell carcinoma, or D.) renal cell carinoma and used as adjuvant therapy following nephrectomy in patients who are at high risk for recurrence.
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SYMDEKO

Products Affected

- Symdeko

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis and One of the following A.) Patient is homozygous for the F508del mutation, or B.) Patient has at least one mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene that is responsive to tezacaftor/ivacaftor verified by an FDA-cleared CF mutation test
Age Restrictions	6 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SYNAREL

Products Affected

- Synarel

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) pregnancy, B.) breastfeeding, C.) undiagnosed abnormal vaginal bleeding
Required Medical Information	Diagnosis of one of the following A.) central precocious puberty, or B.) endometriosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

SYNRIBO

Products Affected

- Synribo

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of chronic or accelerated phase chronic myeloid leukemia (CML) and patient has tried and failed or has a contraindication or intolerance to 2 tyrosine kinase inhibitors
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	A.) Diagnosis of locally advanced or metastatic anaplastic thyroid carcinoma with BRAF V600E mutation, in combination with trametinib and no satisfactory locoregional treatment options OR B.) Diagnosis of metastatic non-small cell lung cancer with BRAF V600E mutation, in combination with trametinib OR in patients previously treated as monotherapy OR C.) Diagnosis of unresectable or metastatic malignant melanoma with BRAF V600E or V600K mutation AND 1) used as monotherapy OR 2) in combination with trametinib OR 3) used as adjuvant therapy following complete resection in patients with lymph node involvement AND used in combination with trametinib.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TAGRISSO

Products Affected

- Tagrisso

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) metastatic, non-small cell lung cancer (NSCLC) with EGFR exon 19 deletion or exon 21 L858R mutation and used as first line therapy, or B.) Metastatic, non-small cell lung cancer with confirmed presence of T790M EGFR mutation AND whose disease has progressed on or after EGFR tyrosine kinase inhibitor based therapy (Diagnosis should be confirmed by an FDA-approved test)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TAKHZYRO

Products Affected

- Takhzyro

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of hereditary angioedema and used in prevention of attacks
Age Restrictions	12 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TALZENNA

Products Affected

- Talzenna

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of deleterious or suspected deleterious germline BRCA-mutated (gBRCAm), HER2-negative locally advanced or metastatic breast cancer
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TARGRETIN

Products Affected

- bexarotene
- Targretin topical

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of cutaneous T-cell lymphoma (CTCL) and patient is not a candidate for or had an inadequate response, is intolerant to, or has a contraindication to at least one prior systemic therapy (e.g., corticosteroids). Female patients of child-bearing potential have a documented negative pregnancy test one week prior to the initiation of therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TASIGNA

Products Affected

- Tasigna

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Long QT syndrome, B.) Uncorrected hypokalemia, C.) Uncorrected hypomagnesemia, or D.) Concomitant use with a drug known to prolong the QT interval or strong cytochrome P450 3A4 inhibitors
Required Medical Information	Diagnosis of one of the following A.) Newly diagnosed, chronic phase Philadelphia chromosome-positive chronic myelogenous leukemia (CML), B.) Chronic-phase and accelerated-phase Philadelphia chromosome-positive CML in patients resistant or intolerant to prior therapy that include imatinib, or C.) Chonic phase Phildelphia chromosome-positive CML in patients with resistance or intolerance to prior tyrosine-kinase inhibitor therapy
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TAZORAC

Products Affected

- tazarotene
- Tazorac topical cream 0.05 %
- Tazorac topical gel

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) acne vulgaris and patient has trial with at least one generic topical acne product, or B.) stable moderate to severe plaque psoriasis with 20% or less body surface area involvement and patient has trial with at least one other topical psoriasis product (e.g., medium to high potency corticosteroid and/or vitamin D analogs)
Age Restrictions	12 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TAZVERIK

Products Affected

- Tazverik

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of (A) metastatic or locally advanced epithelioid sarcoma in patients not eligible for complete resection (B) Follicular lymphoma, Relapsed or refractory, EZH2 mutation-positive, in patients who have received at least 2 prior systemic therapies
Age Restrictions	16 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TEGSEDI

Products Affected

- Tegsedi

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Platelet count less than 100,000 per microliter, B.) urinary protein to creatinine ratio (UPCR) of 1000 mg/g or higher
Required Medical Information	Diagnosis of Polyneuropathy of hereditary transthyretin-mediated amyloidosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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

THALOMID

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	One of the following A.) Newly diagnosed multiple myeloma, in combination with dexamethasone, B.) severe erythema nodosum leprosum with cutaneous manifestations and the medication will not be used as monotherapy if the member has moderate to severe neuritis or C.) acute treatment of the cutaneous manifestations of moderate to severe erythema nodosum leprosum.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or infectious disease specialist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TIBSOVO

Products Affected

- Tibsovo

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Acute myeloid leukemia in relapsed or refractory patients, with susceptible isocitrate dehydrogenase-1 mutation, or B.) Newly-diagnosed acute myeloid leukemia with susceptible isocitrate dehydrogenase-1 mutation in patients 75 years or older or with comorbidities that preclude intensive induction chemotherapy.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TOPICAL AGENTS FOR ATOPIC DERMATITIS

Products Affected

- tacrolimus topical ointment 0.03 %

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	Authorize use in patients who have tried a prescription strength topical corticosteroid (brand or generic) for the current condition. Dermatologic condition on or around the eyes, eyelids, axilla, or genitalia, authorize use without a trial of a prescription strength topical corticosteroid.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TOREMIFENE

Products Affected

- toremifene

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following: A.) Acquired or congenital long QT syndrome, B.) Uncorrected hypokalemia, C.) Uncorrected hypomagnesemia
Required Medical Information	Diagnosis of metastatic breast cancer, in postmenopausal women with estrogen-receptor positive or unknown tumors, and patient must have previous inadequate response or intolerance to tamoxifen.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	6 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TRACLEER

Products Affected

- bosentan
- Tracleer oral tablet for suspension

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Receiving concomitant cyclosporine A or glyburide therapy, B.) Aminotransferase elevations are accompanied by signs or symptoms of liver dysfunction or injury or increases in bilirubin at least 2 times the upper limit of normal, or C.) Pregnancy
Required Medical Information	Diagnosis of pulmonary arterial hypertension that was confirmed by right heart catheterization or Doppler echocardiogram if patient is unable to undergo a right heart catheterization (e.g., patient is frail, elderly, etc.) AND all of the following: A.) Patient has WHO Group I PAH, B.) Patient has New York Heart Association (NYHA) Functional Class II-IV, and C.) Female patients of reproductive potential must use two forms of reliable contraception
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Diagnosis of advanced prostate cancer and used in palliative treatment
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TRETINOIN

Products Affected

- tretinoin topical cream

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of mild to moderate acne vulgaris
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TRIENTINE

Products Affected

- trientine

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of Wilson's disease and intolerance to penicillamine
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

TRIKAFTA

Products Affected

- Trikafta

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cystic fibrosis and patient has at least 1 F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene verified by an FDA-cleared CF mutation test
Age Restrictions	12 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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	Diagnosis of symptomatic tenosynovial giant cell tumor (TGCT) associated with severe morbidity or functional limitations and not amenable to improvement with surgery.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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/Lapatinib 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/Lapatinib will be used in combination with Xeloda or 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/Lapatinib 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 patient has not been previously treated with a HER2-inhibitor Tykerb/Lapatinib.
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Bone cancer-chordoma, colon or rectal cancer

UPTRAVI

Products Affected

- Uptravi oral tablet 1,000 mcg, 1,200 mcg, 1,400 mcg, 1,600 mcg, 200 mcg, 400 mcg, 600 mcg, 800 mcg
- Uptravi oral tablets,dose pack

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of pulmonary arterial hypertension (WHO Group 1) confirmed by right heart catheterization and patient has tried and had an insufficient response to at least one other PAH agent (e.g., sildenafil) therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VALCHLOR

Products Affected

- Valchlor

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of cutaneous T-cell lymphoma (stage IA and IB mycosis fungoides-type) and patient has received prior skin-directed therapy (e.g. Topical corticosteroids, phototherapy, or topical nitrogen mustard)
Age Restrictions	18 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VENCLEXTA

Products Affected

- Venclexta
- Venclexta Starting Pack

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with strong CYP3A inhibitor during the initial and titration phase in patients with CLL or SLL
Required Medical Information	Diagnosis of one of the following A.) chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) or B.) Newly-diagnosed acute myeloid leukemia (AML) and used in combination with azacitidine, decitabine or low-dose cytarabine in patients 75 years or older or who have comorbidities that preclude use of intensive induction chemotherapy.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VERZENIO

Products Affected

- Verzenio

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	HR status, HER2 status, previous medications/therapies tried, concomitant therapy, menopausal status
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 Verzenio 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 Verzenio will be used in combination with anastrozole, exemestane, or letrozole 3. Patient is postmenopausal and meets the following conditions: Verzenio will be used in combination with Faslodex. 4. patient is premenopausal or perimenopausal and meets the following conditions: The patient is receiving ovarian suppression/ablation with a gonadotropin-releasing hormone (GnRH) agonist, or has had surgical bilateral oophorectomy or ovarian irradiation AND Verzenio will be used in combination with Faslodex 5. patient is postmenopausal, premenopausal/perimenopausal (patient is receiving ovarian suppression/ablation with GnRH agonist or has had surgical bilateral oophorectomy or ovarian irradiation) or 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) and meets the following conditions: Verzenio will be used as monotherapy

PA Criteria	Criteria Details
	AND patient's breast cancer has progressed on at least one prior endocrine therapy (e.g., anastrozole, exemestane, letrozole, tamoxifen, Fareston, exemestane plus Afinitor, Faslodex, Afinitor plus Faslodex or tamoxifen, megestrol acetate, fluoxymesterone, ethinyl estradiol) AND patient has tried chemotherapy for metastatic breast cancer. 6. pt is a man who is receiving GnRH agonist AND Verzenio with be used in combination with anastrozole, exemestane or letrozole 7. Patient is a man and Verzenio will be used in combination with Faslodex
Indications	All FDA-approved Indications, Some Medically-accepted Indications.
Off-Label Uses	Men with breast cancer

VIGABATRIN

Products Affected

- vigabatrin
- Vigadrone

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Infantile spasms, or B.) Refractory complex partial seizures and the drug is being used as adjunctive therapy in patients who have responded inadequately to two alternative treatments (e.g. carbamazepine, lamotrigine, levetiracetam, oxcarbazepine, valproic acid, divalproex sodium). Patient and prescriber must be enrolled in the Vigabatrin REMS program.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VITRAKVI

Products Affected

- Vitrakvi oral capsule 100 mg, 25 mg
- Vitrakvi oral solution

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of metastatic or surgically unresectable neurotrophic receptor tyrosine kinase (NTRK) gene fusion positive solid tumors and used in patients with unsatisfactory alternative treatments or who have progressed following treatment
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VIZIMPRO

Products Affected

- Vizimpro

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	First line treatment of metastatic non-small cell lung cancer with confirmed epidermal growth factor receptor (EGFR) exon 19 deletion or exon 21 L858R substitution mutations as detected by an FDA-approved test
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VORICONAZOLE

Products Affected

- voriconazole intravenous
- voriconazole oral suspension for reconstitution
- voriconazole oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Invasive aspergillosis, B.) Candidemia, C.) Esophageal Candidiasis, D.) Invasive candidiasis of the skin and abdomen, kidney, bladder wall, and wounds, or E.) Serious fungal infection due to Scedosporium apiospermum or Fusarium species
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist
Coverage Duration	6 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

VOTRIENT

Products Affected

- Votrient

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) advanced renal cell carcinoma, or B.) advanced soft tissue sarcoma and patient received at least one prior chemotherapy (e.g., doxorubicin, dacarbazine, ifosfamide, epirubicin, docetaxel, or vinorelbine)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XALKORI

Products Affected

- Xalkori

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK)-positive or ROS1-positive as detected by an FDA-approved test
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XELJANZ

Products Affected

- Xeljanz oral tablet
- Xeljanz XR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following: A.) Moderate to severe rheumatoid arthritis (RA), B.) Active psoriatic arthritis, or C.) Moderate to severe ulcerative colitis (UC).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XENAZINE

Products Affected

- tetrabenazine oral tablet 12.5 mg, 25 mg

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Actively suicidal, B.) Untreated or inadequately treated depression, C.) Impaired hepatic function, D.) Concomitant use of monoamine oxidase inhibitors, E.) Concomitant use of reserpine or within 20 days of discontinuing reserpine
Required Medical Information	Diagnosis of chorea associated with Huntington's disease
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XGEVA

Products Affected

- Xgeva

PA Criteria	Criteria Details
Exclusion Criteria	Hypocalcemia (calcium less than 8.0 mg/dL)
Required Medical Information	Diagnosis of one of the following A.) bone metastases from a solid tumor, B.) giant cell tumor of the bone that is unresectable or where surgical resection is likely to result in severe morbidity, C.) hypercalcemia of malignancy refractory to bisphosphonate therapy, or D.) multiple myeloma used for the prevention of skeletal related events
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XOLAIR

Products Affected

- Xolair

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Chronic idiopathic urticaria in patients who remain symptomatic despite H1 antihistamine therapy, or B.) Moderate to severe persistent asthma in patients with a positive skin test or in vitro reactivity to a perennial aeroallergen and symptoms are inadequately controlled with inhaled corticosteroids
Age Restrictions	6 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an allergist, immunologist, pulmonologist, or dermatologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XOSPATA

Products Affected

- Xospata

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of relapsed or refractory acute myeloid leukemia, with presence of FLT3 mutation as detected by an FDA-approved test
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XPOVIO

Products Affected

- Xpovio

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of relapsed or refractory multiple myeloma being used in combination with dexamethasone in patient who has received at least 4 prior therapies and is refractory to at least 2 proteasome inhibitors, at least 2 immunomodulatory agents, and an anti-CD38 monoclonal antibody. For use as monotherapy for the treatment of adult patients with relapsed or refractory diffuse large B-cell lymphoma
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	12 months
Other Criteria	Monotherapy for the treatment of adult patients with relapsed or refractory diffuse large B-cell lymphoma
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XTANDI

Products Affected

- Xtandi

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) castration-resistant prostate cancer, or B.) metastatic, castration-sensitive prostate cancer
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XURIDEN

Products Affected

- Xuriden

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of hereditary orotic aciduria
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

XYREM

Products Affected

- Xyrem

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant treatment with sedative hypnotic agents, B.) Succinic semialdehyde dehydrogenase deficiency
Required Medical Information	Diagnosis of cataplexy and excessive daytime sleepiness in patients with narcolepsy
Age Restrictions	7 years of age and older
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

YONSA

Products Affected

- Yonsa

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Pregnancy, B.) Patients with severe baseline hepatic impairment (Child-Pugh Class C)
Required Medical Information	Diagnosis of metastatic castration-resistant prostate cancer and All of the following A.) used in combination with methylprednisolone
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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	3 years
Other Criteria	Ovarian, fallopian tube, or primary peritoneal cancer, maintenance therapy - approve if the patient is in complete or partial response after platinum-based chemotherapy regimen. Ovarian, fallopian tube, or primary peritoneal cancer, treatment-approve per label if the patient has tried at least three prior chemotherapy regimens and has homologous recombination deficiency (HRD)-positive disease as confirmed by an approved test.
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	Diagnosis of one of the following A.) Unresectable or metastatic melanoma and patient has positive BRAF-V600E mutation documented by an FDA-approved test, or B.) Erdheim-Chester disease and patient has documented BRAF V600 mutation
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

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. PBPC-prescribed by or in consultation with an oncologist, hematologist, or physician that specializes in transplantation.
Coverage Duration	Cancer pts receiving chemo-6 mo. PBPC-1 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, Some Medically-accepted Indications.
Off-Label Uses	Patients undergoing PBPC collection and therapy

ZOLINZA

Products Affected

- Zolinza

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	3 years
Other Criteria	Cutaneous T-Cell Lymphoma including Mycosis Fungoides/Sezary Syndrome-approve.
Indications	All FDA-approved Indications.
Off-Label Uses	N/A

ZYDELIG

Products Affected

- Zydelig

PA Criteria	Criteria Details
Exclusion Criteria	History of severe allergic reactions including anaphylaxis and toxic epidermal necrolysis
Required Medical Information	Diagnosis of one of the following A.) Chronic lymphocytic leukemia and all of the following: Used in combination with rituximab, patient has relapsed on at least one prior therapy (e.g., purine analogues [fludarabine, pentostatin, cladribine], alkylating agents [chlorambucil, cyclophosphamide], or monoclonal antibodies [rituximab]), and patient does not have any co-morbidities that prevents the use of cytotoxic chemotherapy (i.e. severe neutropenia or thrombocytopenia, creatinine clearance less than 60 mL/minute), B.) Non-Hodgkins lymphoma (Follicular, B-Cell) and the patient has relapsed on at least two prior systemic therapies (e.g., rituximab, alkylating agents [cyclophosphamide, chlorambucil], anthracyclines [doxorubicin, daunorubicin], purine analogs [fludarabine]), or C.) Small lymphocytic lymphoma and the patient has relapsed on at least two prior systemic therapies (e.g., rituximab, alkylating agents [cyclophosphamide, chlorambucil], anthracyclines [doxorubicin, daunorubicin], purine analogs [fludarabine])
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ZYKADIA

Products Affected

- Zykadia oral tablet

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

ZYTIGA

Products Affected

- abiraterone oral tablet 250 mg, 500 mg

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Diagnosis of one of the following A.) Castration-resistant metastatic prostate cancer and used in combination with prednisone, or B.) High risk, castration-sensitive metastatic prostate cancer and used in combination with prednisone
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	12 months
Other Criteria	N/A
Indications	All Medically-accepted 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 oral capsule
- aprepitant oral capsule,dose pack
- azathioprine
- Brovana
- budesonide inhalation
- calcitriol oral
- caspofungin
- Clinimix 5%/D15W Sulfite Free
- Clinimix 4.25%/D10W Sulf Free
- Clinimix 4.25%/D5W Sulfit Free
- Clinimix 5%-D20W(sulfite-free)
- 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
- Envarsus XR
- everolimus (immunosuppressive)
- Gengraf oral capsule 100 mg, 25 mg
- Gengraf oral solution
- granisetron HCl oral
- Hepatamine 8%
- Intralipid intravenous emulsion 20 %
- Intralipid intravenous emulsion 30 %
- ipratropium bromide inhalation
- ipratropium-albuterol
- methotrexate sodium
- methotrexate sodium (PF) injection solution
- mycophenolate mofetil
- mycophenolate sodium
- Nephramine 5.4 %
- Nutrilipid
- Octagam
- ondansetron
- ondansetron HCl oral
- Panzyga
- paricalcitol oral
- pentamidine inhalation
- Premasol 10 %
- Privigen
- 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
- Syndros
- tacrolimus oral
- tobramycin in 0.225 % NaCl
- Travasol 10 %
- TrophAmine 10 %
- Xatmep
- Zortress oral tablet 0.25 mg, 0.5 mg, 0.75 mg, 1 mg

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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