

ABIRATERONE

Products Affected

- *abiraterone acetate*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Castration-resistant metastatic prostate cancer (CRPC), or B.) High risk, castration-sensitive metastatic prostate cancer (CSPC). For treatment of CRPC and CSPC, abiraterone will be used in combination with prednisone AND one of the following applies 1.) Used in combination with a gonadotropin-releasing hormone (GnRH) analog (e.g. leuprolide, triptorelin), OR 2) Patient has received bilateral orchiectomy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ACITRETIN

Products Affected

- *acitretin*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Severely impaired liver or kidney function, B.) Chronic abnormally elevated blood lipid values, C.) Concomitant use of methotrexate or tetracyclines, D.) Pregnancy
Required Medical Information	Diagnosis of 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ACTIMMUNE

Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic granulomatous disease for use in reducing the frequency and severity of serious infections, or B.) Severe, malignant osteopetrosis (SMO)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

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 Doppler echocardiogram if patient is unable to undergo a right heart catheterization (e.g., patient is frail, elderly, etc.), 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 and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

AKEEGA

Products Affected

- AKEEGA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of deleterious or suspected deleterious BRCA-mutated (BRCAm) metastatic castration-resistant prostate cancer (mCRPC) AND used in combination with prednisone
Age Restrictions	18 years or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	end of plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ALECENSA

Products Affected

- ALECENSA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer as detected by an FDA-approved test
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ALPHA-1 PROTEINASE INHIBITOR

Products Affected

- PROLASTIN-C INTRAVENOUS SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	Immunoglobulin A (IgA) deficiency with antibodies against IgA
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	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ALUNBRIG

Products Affected

- ALUNBRIG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of anaplastic lymphoma kinase-positive (ALK) 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 or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ALVAIZ

Products Affected

- ALVAIZ

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic idiopathic thrombocytopenic purpura (ITP), B.) Chronic hepatitis C infection associated thrombocytopenia, or C.) Severe aplastic anemia with insufficient response to immunosuppressive therapy
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

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 classified as WHO Group I, 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.)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ARCALYST

Products Affected

- ARCALYST

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Cryopyrin-associated periodic syndromes (CAPS), including familial cold autoinflammatory syndrome (FCAS) and Muckle-Wells Syndrome (MWS), B.) Deficiency of interleukin-1 receptor antagonist (DIRA) and patient requires maintenance therapy for remission, or C.) Recurrent pericarditis (RP) and reduction in risk of recurrence
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ARIKAYCE

Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	Known sensitivity to any aminoglycoside
Required Medical Information	Diagnosis of pulmonary Mycobacterium avium complex (MAC) infection and used as part of a combination antibacterial regimen in treatment refractory patients (greater than 6 months of a multidrug background regimen)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist or pulmonologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

AUGTYRO

Products Affected

- AUGTYRO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Locally advanced or metastatic ROS1-positive non-small cell lung cancer, or B.) solid tumors that have an NTRK gene fusion are locally advanced, metastatic, or where surgical resection is likely to result in severe morbidity, and have progressed following treatment or have no satisfactory alternative therapy
Age Restrictions	12 years or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	end of plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

AURYXIA

Products Affected

- AURYXIA

PA Criteria	Criteria Details
Exclusion Criteria	Iron overload syndrome (e.g. hemochromatosis)
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	Plan Year
Other Criteria	Ferric Citrate is NOT approvable for iron deficiency anemia per Part D law
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

AUSTEDO

Products Affected

- AUSTEDO
- AUSTEDO XR ORAL TABLET EXTENDED RELEASE 24 HOUR 12 MG, 24 MG, 30 MG, 36 MG, 42 MG, 48 MG, 6 MG
- AUSTEDO XR PATIENT TITRATION ORAL TABLET EXTENDED RELEASE THERAPY PACK 6 & 12 & 24 MG

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Suicidal ideation and/or untreated or inadequately treated depression in a patient with Huntington's Disease, B.) Hepatic impairment, C.) Concomitant use of MAOIs, reserpine, tetrabenazine, or valbenazine
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 neurologist or psychiatrist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

AYVAKIT

Products Affected

- AYVAKIT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Unresectable or metastatic gastrointestinal stromal tumor, with a platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation, including PDGFRA D842V mutations, B.) Advanced Systemic Mastocytosis (AdvSM), including aggressive systemic mastocytosis (ASM), systemic mastocytosis with an associated hematological neoplasm (SMAHN), or mast cell leukemia (MCL), and platelet count of at least 50,000/mcL, or C.) Indolent systemic mastocytosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BALVERSA

Products Affected

- BALVERSA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of locally advanced or metastatic urothelial carcinoma and both of the following 1.) Susceptible fibroblast growth factor receptor (FGFR)3 or FGFR2 genetic alterations confirmed by an FDA-approved diagnostic test, and 2.) 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

BENLYSTA

Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Active, autoantibody-positive, system lupus erythematosus (SLE), or B.) Active lupus nephritis and patient is receiving standard therapy
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a nephrologist or rheumatologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BESREMI

Products Affected

- BESREMI

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Existence of, or history of severe psychiatric disorders (severe depression, suicidal ideation, or suicide attempt), B.) Hypersensitivity to interferons including interferon alfa-2b or excipients, C.) Hepatic impairment (Child-Pugh B or C), D.) History or presence of active serious or untreated autoimmune disease, or E.) Immunosuppressed transplant recipients
Required Medical Information	Diagnosis of polycythemia vera
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BEXAROTENE GEL

Products Affected

- *bexarotene*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of primary cutaneous T-cell lymphoma (CTCL Stage 1A/1B) and patient had an inadequate response, is intolerant to, or has a contraindication to at least one prior systemic therapy (e.g., corticosteroids) indicated for cutaneous manifestations of CTCL
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or dermatologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BEXAROTENE ORAL

Products Affected

- *bexarotene*

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) for cutaneous manifestations of CTCL
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an dermatologist, hematologist, or oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BOSENTAN

Products Affected

- *bosentan*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant cyclosporine A or glyburide therapy, or B.) Pregnancy
Required Medical Information	Diagnosis of pulmonary arterial hypertension (WHO Group I) and patient has New York Heart Association (NYHA) Functional Class II-IV, 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.)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

BOSULIF

Products Affected

- BOSULIF

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic, accelerated, or blast phase Philadelphia chromosome-positive (Ph+) chronic myelogenous leukemia (CML) with resistance or inadequate response to prior therapy, or B.) Newly diagnosed chronic phase Philadelphia chromosome-positive (Ph+) CML
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

BRAFTOVI

Products Affected

- BRAFTOVI ORAL CAPSULE 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) unresectable or metastatic melanoma with documented BRAF V600E or V600K mutation as detected by a FDA-approved test and used in combination with binimetinib, B.) metastatic colorectal cancer with documented BRAF V600E mutation as detected by a FDA-approved test, patient has received prior therapy, and braftovi used in combination with cetuximab, or C.) Metastatic non-small cell lung cancer with a BRAF V600E mutation as detected by an FDA-approved test and used in combination with binimetinib
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BRONCHITOL

Products Affected

- BRONCHITOL

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Cystic fibrosis of the lung
Age Restrictions	None
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	Plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

BRUKINSA

Products Affected

- BRUKINSA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following: A.) mantle cell lymphoma (MCL) and patient has received at least one prior therapy, B.) Treatment of adult patients with Waldenström macroglobulinemia, C.) Treatment of adult patients with relapsed or refractory marginal zone lymphoma who have received at least one anti-CD20-based regimen, D.) Chronic lymphocytic leukemia, E.) Small lymphocytic lymphoma, or F.) Relapsed or refractory follicular lymphoma, in combination with obinutuzumab, after 2 or more lines of systemic therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

BYLVAY

Products Affected

- BYLVAY
- BYLVAY (PELLETS)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Progressive familial intrahepatic cholestasis-associated pruritus, or B.) Cholestatic pruritus in patients with Alagille syndrome
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

CABLIVI

Products Affected

- CABLIVI

PA Criteria	Criteria Details
Exclusion Criteria	None
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CABOMETYX

Products Affected

- CABOMETYX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Advanced renal cell carcinoma, B.) Advanced hepatocellular carcinoma (HCC) and patient has been previously treated with sorafenib, C.) Advanced renal cell carcinoma and used as first line treatment in combination with nivolumab or D.) Treatment of adults and pediatric patients 12 years and older with locally advanced or metastatic differentiated thyroid cancer that has progressed following VEGFR-targeted therapy and who are radioactive iodine-refractory or ineligible
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CALQUENCE

Products Affected

- CALQUENCE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Mantle cell lymphoma (MCL) and patient has received at least 1 prior therapy, B.) Chronic lymphocytic leukemia (CLL), or C.) Small lymphocytic lymphoma (SLL)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CAMZYOS

Products Affected

- CAMZYOS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of symptomatic New York Heart Association (NYHA) class II-III obstructive hypertrophic cardiomyopathy (HCM) in adult patients
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CAPRELSA

Products Affected

- CAPRELSA

PA Criteria	Criteria Details
Exclusion Criteria	Congenital long QT syndrome
Required Medical Information	Diagnosis of metastatic or unresectable locally advanced medullary thyroid cancer (MTC) AND disease is symptomatic or progressive
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CARGLUMIC ACID

Products Affected

- *carglumic acid oral tablet soluble*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) N-acetyl glutamate synthase (NAGS) deficiency (confirmed by appropriate genetic testing) with acute or chronic hyperammonemia, or B.) Propionic or methylmalonic acidemia with acute hyperammonemia
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CAYSTON

Products Affected

- CAYSTON

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystic fibrosis (confirmed by appropriate diagnostic or genetic testing) and patient has Pseudomonas aeruginosa lung infection confirmed by positive culture
Age Restrictions	7 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

CHANTIX

Products Affected

- *varenicline tartrate (starter)*
- *varenicline tartrate oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of nicotine addiction and documentation of trial of previous smoking cessation therapies (nicotine replacement therapy or a therapeutic course of bupropion (7-9 weeks)). Otherwise, varenicline will require a prior authorization exception request indicating: (1) history of inadequate treatment response with a nicotine or bupropion smoking cessation therapy, OR (2) history of adverse event with a nicotine or bupropion smoking cessation therapy, OR (3) smoking cessation therapy with nicotine or bupropion is contraindicated.
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	12 weeks
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

CNS STIMULANTS

Products Affected

- *armodafinil*
- *modafinil oral*

PA Criteria	Criteria Details
Exclusion Criteria	None
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 disorder (SWD)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

COMETRIQ

Products Affected

- COMETRIQ (100 MG DAILY DOSE) ORAL KIT 80 & 20 MG
- COMETRIQ (140 MG DAILY DOSE) ORAL KIT 3 X 20 MG & 80 MG
- COMETRIQ (60 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of progressive, metastatic medullary thyroid cancer
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

COPIKTRA

Products Affected

- COPIKTRA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsed or refractory chronic lymphocytic leukemia (CLL), or B.) Relapsed or refractory small lymphocytic lymphoma (SLL). For CLL or SLL, the patient must have history of at least 2 prior therapies
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

COSENTYX

Products Affected

- COSENTYX (300 MG DOSE)
- COSENTYX SENSOREADY (300 MG)
- COSENTYX SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 75 MG/0.5ML
- COSENTYX UNOREADY

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Ankylosing spondylitis and patient has trial and failure, contraindication, or intolerance to two preferred products, (i.e. Enbrel, Humira, Rinvoq), B.) Moderate to severe plaque psoriasis in adults and patient has trial and failure, contraindication, or intolerance to two preferred products, (i.e. Enbrel, Humira, Skyrizi, Stelara), C.) Moderate to severe plaque psoriasis in patients 6 years to less than 18 years of age and patient has trial and failure, contraindication, or intolerance to two preferred products, (i.e. Enbrel, Stelara), D.) Active psoriatic arthritis in adult patient and has trial and failure, contraindication, or intolerance to two preferred products, (i.e. Enbrel, Humira, Rinvoq, Skyrizi, Stelara), E.) Active psoriatic arthritis in patients 2 years to less than 18 years of age, F.) Non-radiographic axial spondyloarthritis and patient has trial and failure, contraindication, or intolerance to one preferred product, (i.e. Rinvoq), G.) Active enthesitis-related arthritis, or H.) Moderate to severe hidradenitis suppurativa in adults and patient has trial and failure, contraindication, or intolerance to one preferred product, (i.e. Humira)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

COTELLIC

Products Affected

- COTELLIC

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.)unresectable or metastatic malignant melanoma with BRAF V600E OR V600K mutation, and documentation of combination therapy with vemurafenib (Zelboraf), or B.) Histiocytic neoplasms
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CYSTAGON

Products Affected

- CYSTAGON

PA Criteria	Criteria Details
Exclusion Criteria	Known serious hypersensitivity to penicillamine or cysteamine
Required Medical Information	Diagnosis of nephropathic cystinosis confirmed by the presence of increased cystine concentration in leukocytes or by genetic testing
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

CYSTEAMINE OPHTH

Products Affected

- CYSTADROPS
- CYSTARAN

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystinosis and patient has corneal cystine crystal accumulation
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DALFAMPRIDINE

Products Affected

- *dalfampridine er*

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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DAURISMO

Products Affected

- DAURISMO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of newly diagnosed acute myeloid leukemia (AML) and used in combination with cytarabine in patients 75 years of age or older OR in patients that 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 or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DAYBUE

Products Affected

- DAYBUE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Rett syndrome
Age Restrictions	2 years of age and older
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DEFERASIROX

Products Affected

- *deferasirox oral tablet*
- *deferasirox oral tablet soluble*

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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DEFERIPRONE

Products Affected

- *deferiprone*
- FERRIPROX ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Must meet all of the following 1.) Diagnosis of transfusional iron overload due to thalassemia syndromes, sickle cell disease, or other anemias, 2.) Patient has failed prior chelation therapy, and 3.) Patient has an absolute neutrophil count greater than $1.5 \times 10^9/L$
Age Restrictions	3 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DIACOMIT

Products Affected

- DIACOMIT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of severe myoclonic epilepsy in infancy (Dravet syndrome) in patients taking clobazam
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DICLOFENAC TOPICAL

Products Affected

- *diclofenac sodium external gel 3 %*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Actinic keratosis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DIFICID

Products Affected

- DIFICID

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of diarrhea associated with clostridioides difficile infection and patient has had an inadequate treatment response, intolerance, or contraindication to generic oral vancomycin
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	4 weeks
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DIMETHYL FUMARATE

Products Affected

- *dimethyl fumarate oral*
- *dimethyl fumarate starter pack oral capsule delayed release therapy pack*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DOJOLVI

Products Affected

- DOJOLVI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Long-chain fatty acid oxidation disorder (LC-FAOD)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DRONABINOL

Products Affected

- *dronabinol*

PA Criteria	Criteria Details
Exclusion Criteria	Sesame oil hypersensitivity
Required Medical Information	Diagnosis of one of the following A.) Anorexia associated to AIDS, or B.) Chemotherapy-induced nausea and vomiting
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

DROXIDOPA

Products Affected

- *droxidopa*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of symptomatic neurogenic orthostatic hypotension (nOH) caused by primary autonomic failure (e.g., Parkinson disease, multiple system atrophy, pure autonomic failure), dopamine beta-hydroxylase deficiency, or non-diabetic autonomic neuropathy
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

DUPIXENT

Products Affected

- DUPIXENT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe atopic dermatitis and if patient is 2 years or older has trial/failure, contraindication, or intolerance to two of the following 1.) Topical corticosteroid and/or 2.) Topical calcineurin inhibitor, B.) Eosinophilic phenotype or oral corticosteroid-dependent moderate to severe asthma and used as an adjunct treatment, or C.) Chronic rhinosinusitis with nasal polyposis and used as an adjunct treatment, D.) Eosinophilic esophagitis, or E.) Prurigo nodularis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EMGALITY

Products Affected

- EMGALITY

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic or episodic migraine disorder and patient has documented trial, inadequate response, or contraindication to at least 2 generic formulary drugs used for migraine prevention (i.e., propranolol, topiramate, divalproex, timolol), or B.) Episodic cluster headache
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ENBREL

Products Affected

- ENBREL MINI
- ENBREL SUBCUTANEOUS SOLUTION 25 MG/0.5ML
- ENBREL SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- ENBREL SURECLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	None
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
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ENSPRYNG

Products Affected

- ENSPRYNG

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Active Hepatitis B infection, or B.) Active or untreated latent tuberculosis
Required Medical Information	Diagnosis of neuromyelitis optica spectrum disorder (NMOSD) in patients who are anti-aquaporin-4 (AQP4) antibody positive
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist, immunologist, or ophthalmologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EPIDIOLEX

Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Lennox-Gastaut syndrome, B.) Severe myoclonic epilepsy in infancy (Dravet syndrome), or C.) Seizures associated with tuberous sclerosis complex (TSC)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EPOETIN THERAPY

Products Affected

- RETACRIT INJECTION SOLUTION UNIT/ML, 4000 UNIT/ML, 40000
10000 UNIT/ML, 10000 UNIT/ML(1ML), UNIT/ML
2000 UNIT/ML, 20000 UNIT/ML, 3000

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Non-myeloid neoplastic disease and utilized for the treatment of chemotherapy induced anemia, B.) HIV infection and utilized for the treatment of zidovudine induced anemia, C.) Chronic kidney disease resulting in anemia, or D.) High risk surgical candidate status at risk for perioperative blood loss and undergoing elective, noncardiac, or nonvascular surgery
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ERIVEDGE

Products Affected

- ERIVEDGE

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Metastatic basal cell carcinoma, or 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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ERLEADA

Products Affected

- ERLEADA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Nonmetastatic, castration-resistant prostate cancer (nmCRPC), or B.) Metastatic, castration-sensitive prostate cancer (mCSPC). For treatment of nmCRPC and mCSPC, one of the following applies 1.) Used in combination with a gonadotropin-releasing hormone (GnRH) analog, OR 2) Patient has received bilateral orchiectomy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ERLOTINIB

Products Affected

- *erlotinib hcl*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Locally advanced, unresectable, or metastatic pancreatic cancer and erlotinib will be used in combination with gemcitabine, B.) Locally advanced or metastatic non-small cell lung cancer with 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 AND one of the following 1.) Erlotinib will be used as first-line treatment, 2.) Failure with at least one prior chemotherapy regimen, or 3.) No evidence of disease progression after four cycles of first-line platinum-based chemotherapy and erlotinib will be used as maintenance treatment
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EVEROLIMUS

Products Affected

- everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Hypersensitivity to everolimus or excipients, or B.) Hypersensitivity to rapamycin derivatives (e.g. sirolimus)
Required Medical Information	Diagnosis of one of the following A.) Renal angiomyolipoma and tuberous sclerosis complex (TSC) not requiring immediate surgery, B.) Advanced hormone receptor-positive, HER2 negative breast cancer in postmenopausal women and taken in combination with exemestane, after failure with letrozole or anastrozole, C.) Progressive, well-differentiated, nonfunctional neuroendocrine tumors of gastrointestinal or lung origin and disease is unresectable, locally advanced, or metastatic, D.) Pancreatic progressive neuroendocrine tumors and disease is unresectable, locally advanced, or metastatic, E.) Advanced renal cell carcinoma (RCC) after failure with sunitinib or sorafenib, F.) Subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex in patients who are not candidates for curative surgical resection
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EVEROLIMUS SUSPENSION

Products Affected

- *everolimus oral tablet soluble*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Hypersensitivity to everolimus , or B.) Hypersensitivity to rapamycin derivatives (e.g. sirolimus)
Required Medical Information	Diagnosis of one of the following A.) Tuberous sclerosis complex (TSC)-associated partial-onset seizures, or B.) Subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex in patients who are not candidates for curative surgical resection
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

EVRYSDI

Products Affected

- EVRYSDI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of spinal muscular atrophy (SMA)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

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, C.) Known or suspected gastrointestinal obstruction, including paralytic ileus, D.) Acute or severe bronchial asthma and used in an unmonitored setting (absence of resuscitative equipment)
Required Medical Information	Must meet all of the following 1.) Diagnosis of cancer-related breakthrough pain, 2.) Patient is currently receiving/tolerant to around-the-clock opioid therapy for persistent cancer pain, and 3.) Patient and prescriber are enrolled in the TIRF REMS Access Program
Age Restrictions	16 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FENTANYL PATCH

Products Affected

- *fentanyl*

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.) Mild or intermittent pain management, C.) Use in opioid non-tolerant patients, D.) Known or suspected gastrointestinal obstruction, including paralytic ileus, E.) Acute or severe bronchial asthma and used in an unmonitored setting (absence of resuscitative equipment)
Required Medical Information	Must meet all of the following 1.) Patient is opioid tolerant (taking for one week or longer at least 60mg of morphine or equivalent daily) and 2.) Patient has tried at least one extended release oral opioids or is unable to take extended release oral opioids secondary to allergy, adverse events, swallowing difficulty, or uncontrollable nausea/vomiting
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FILSPARI

Products Affected

- FILSPARI

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Pregnancy or B.) Concomitant use with angiotensin receptor blockers (ARBs), endothelin receptor antagonists (ERAs), or aliskiren
Required Medical Information	Diagnosis of treatment of primary immunoglobulin A (IgA) nephropathy at risk of rapid disease progression, generally a urine protein to creatinine ratio (UPCR) of 1.5 g/g or more, to reduce proteinuria
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FINGOLIMOD

Products Affected

- *fingolimod hcl*

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., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	10 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FINTEPLA

Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use of an MAOI, or B.) Use within 14 days of discontinuing an MAOI
Required Medical Information	Diagnosis of Severe myoclonic epilepsy in infancy (Dravet syndrome) or seizures associated with Lennox-Gastaut syndrome
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FIRMAGON

Products Affected

- FIRMAGON (240 MG DOSE)
- FIRMAGON SUBCUTANEOUS SOLUTION RECONSTITUTED 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced prostate cancer
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

FOTIVDA

Products Affected

- FOTIVDA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of relapsed or refractory advanced renal cell cancer (RCC) following 2 or more prior systemic therapies
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

FRUZAQLA

Products Affected

- FRUZAQLA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic colorectal cancer (mCRC) and all of the following: A.) patient has been previously treated with fluoropyrimidine, oxaliplatin, irinotecan-based chemotherapy, B.) an anti-VEGF therapy, and C.) if RAS wild-type and medically appropriate, patient has also been previously treated with anti-EGFR therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	3 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GALAFOLD

Products Affected

- GALAFOLD

PA Criteria	Criteria Details
Exclusion Criteria	None
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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GATTEX

Products Affected

- GATTEX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of short bowel syndrome and patient is dependent on parenteral support
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GAVRETO

Products Affected

- GAVRETO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Metastatic RET fusion-positive non-small cell lung cancer (NSCLC) as detected by an FDA approved test, B.) Advanced or metastatic RET-mutant medullary thyroid cancer and patient requires systemic therapy, or C.) Advanced or metastatic RET fusion-positive thyroid and patient requires systemic therapy and is radioactive iodine-refractory, when radioactive iodine is appropriate
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

GEFITINIB

Products Affected

- *gefitinib*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic non-small cell lung cancer (NSCLC) and must meet all of the following 1.) Tumor has 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 and 2.) Used as first-line treatment
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GILOTRIF

Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Metastatic non-small cell lung cancer (NSCLC) in patients whose tumors have non-resistant epidermal growth factor receptor (EGFR) mutations as detected by an FDA-approved test, or B.) Metastatic squamous NSCLC with progression after platinum-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GLATIRAMER

Products Affected

- *glatiramer acetate*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

GLEOSTINE

Products Affected

- GLEOSTINE ORAL CAPSULE 10 MG, 100 MG, 40 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Must meet one of the following: A.) Hodgkin's disease in patient who has relapsed during or failed to respond to primary therapy and is being used in combination with other agents OR B.) Intracranial tumor
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

GLP1

Products Affected

- MOUNJARO
- OZEMPIC (0.25 OR 0.5 MG/DOSE) SUBCUTANEOUS SOLUTION PEN-INJECTOR 2 MG/3ML
- OZEMPIC (1 MG/DOSE) SUBCUTANEOUS SOLUTION PEN-INJECTOR 4 MG/3ML
- OZEMPIC (2 MG/DOSE)
- RYBELSUS
- TRULICITY
- VICTOZA SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Must meet all of the following 1.) The drug is prescribed for an FDA-approved indication, 2.) For a diagnosis of Type 2 Diabetes Mellitus the patient has a trial and failure, contraindication or intolerance to metformin or any metformin combination product
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

GROWTH HORMONE

Products Affected

- OMNITROPE SUBCUTANEOUS SOLUTION CARTRIDGE
- OMNITROPE SUBCUTANEOUS SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Use for growth promotion in pediatric patients with closed epiphyses, B.) Acute critical illness caused by complications following open-heart or abdominal surgery, multiple accidental trauma, or acute respiratory failure, C.) Active malignancy, D.) Active proliferative or severe nonproliferative diabetic retinopathy, E.) Prader-Willi Syndrome in patients who are severely obese, have a history of upper airway obstruction or sleep apnea, or have severe respiratory impairment
Required Medical Information	Diagnosis of pediatric indication: A.) GHD and bone age at least 1 year or 2 standard deviations (SD) delayed compared with chronological age and 2 stim tests with peak GH secretion below 10 ng/mL or IGF-1/IGFBP3 level more than 2 SDS below mean if CNS pathology, h/o irradiation, or proven genetic cause, B.) SGA and birth weight or length 2 or more SDS below mean for gestational age and fails to manifest catch up growth by age 2 (height 2 or more SDS below mean for age and gender), C.) CRI and metabolic abnormalities have been corrected, and patient has not had renal transplant D.) SHOX deficiency or Noonan syndrome E.) PWS confirmed by genetic testing, F.) Turner Syndrome confirmed by chromosome analysis. For GHD, CRI, SHOX deficiency, Noonan syndrome, and PWS one of the following height more than 3 SDS below mean for age and gender, or height more than 2 SDS below mean with GV more than 1 SDS below mean, or GV over 1 year 2 SDS below mean. OR Diagnosis of an adult indication: A.) childhood- or adult-onset GHD confirmed by 2 standard GH stim tests (provide assay): 1 test must be insulin tolerance test (ITT) with blood glucose nadir less than 40 mg/dL (2.2 mmol/L). If contraindicated, use a standardized stim test (i.e. arginine plus GH releasing hormone [preferred], glucagon, arginine), B.) GHD with at least 1 other pituitary hormone deficiency and failed at least 1 GH stim test (ITT preferred), C.) GHD with panhypopituitarism (3 or more pituitary hormone deficiencies), D.) GHD with irreversible hypothalamic-pituitary structural lesions due to tumors, surgery or radiation of pituitary or hypothalamus

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
	region AND a subnormal IGF-1 (after at least 1 month off GH therapy) AND Objective evidence of GHD complications, such as: low bone density, increased visceral fat mass, or cardiovascular complications AND Completed linear growth (GV less than 2 cm/year) AND GH has been discontinued for at least 1 month (if previously receiving GH)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an Endocrinologist or Nephrologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

GUANFACINE ER

Products Affected

- *guanfacine hcl er*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of attention deficit hyperactivity disorder (ADHD)
Age Restrictions	6 years of age to 17 years of age
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

HEPATITIS C

Products Affected

- MAVYRET
- *sofosbuvir-velpatasvir*
- VOSEVI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of HCV genotype, subtype and quantitative HCV RNA (viral load) testing any time prior to therapy. Must document cirrhosis status, prior treatment history (if any), and planned duration of treatment. All genotypes will require trial/failure, contraindication to, or intolerance to Mavyret or Sofosbuvir-Velpatasvir prior to the approval of Vosevi. Genotype and subtype are not required for: (1) initial treatment of patients without cirrhosis if using Sofosbuvir-Velpatasvir or Mavyret OR (2) treatment of patients with compensated cirrhosis if using Mavyret
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, hepatologist, or infectious disease specialist
Coverage Duration	Duration of approval per AASLD Guidelines
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

HUMIRA

Products Affected

- HUMIRA (2 PEN)
- HUMIRA (2 SYRINGE) SUBCUTANEOUS PREFILLED SYRINGE KIT 10 MG/0.1ML, 20 MG/0.2ML, 40 MG/0.4ML, 40 MG/0.8ML
- HUMIRA-CD/UC/HS STARTER SUBCUTANEOUS PEN-INJECTOR KIT 80 MG/0.8ML
- HUMIRA-PED \geq 40KG UC STARTER
- HUMIRA-PSORIASIS/UVEIT STARTER

PA Criteria	Criteria Details
Exclusion Criteria	None
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, G.) Moderate to severe ulcerative colitis in patients who have had an inadequate response to immunosuppressants (e.g. corticosteroids, azathioprine), H.) Non-infectious uveitis (including intermediate, posterior, and panuveitis), or I.) Moderate to severe hidradenitis suppurativa
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin
Prior Authorization 2024
Last Updated 9/24/2024

PA Criteria	Criteria Details
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18
Last Updated: 09/24/2024
Effective: 10/01/2024

HYFTOR

Products Affected

- HYFTOR

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Facial angiofibroma associated with tuberous sclerosis
Age Restrictions	6 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IBRANCE

Products Affected

- IBRANCE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Advanced or metastatic, hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer used in combination with fulvestrant and disease has progressed following endocrine therapy, or B.) Advanced or metastatic, hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer and used in combination with an aromatase inhibitor in a male or female patient as initial endocrine-based therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ICATIBANT

Products Affected

- *icatibant acetate subcutaneous solution
prefilled syringe*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following and used as treatment for acute attacks A.) Hereditary angioedema (HAE) with C1 inhibitor deficiency (Type 1) confirmed by laboratory testing, or B.) HAE with C1 inhibitor dysfunction (Type 2) confirmed by laboratory testing, or C.) HAE with normal C1 inhibitor (Type 3) confirmed by laboratory testing and one of the following 1.) Positive test for an F12, angiotensin-1, or plasminogen gene mutation, or 2.) Family history of angioedema and the angioedema was refractory to a trial of an antihistamine for at least one month
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an allergist, hematologist, or immunologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ICLUSIG

Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	None
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, B.) Chronic phase, chronic myeloid leukemia (CML) in adult patients with resistance or intolerance to at least two prior kinase inhibitors, or C.) 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IDHIFA

Products Affected

- IDHIFA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of relapsed or refractory acute myeloid leukemia (AML) with an isocitrate dehydrogenase 2 (IDH2) 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IMATINIB

Products Affected

- *imatinib mesylate*

PA Criteria	Criteria Details
Exclusion Criteria	None
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 stromal 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
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IMBRUVICA

Products Affected

- IMBRUVICA ORAL CAPSULE
- IMBRUVICA ORAL SUSPENSION
- IMBRUVICA ORAL TABLET 140 MG, 280 MG, 420 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL), B.) Chronic lymphocytic leukemia (CLL)/Small lymphocytic lymphoma (SLL) with 17p deletion, C.) Waldenstrom's macroglobulinemia (WM), or D.) Chronic graft vs host disease (cGVHD) after failure of at least one first-line corticosteroid therapy
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

INBRIJA

Products Affected

- INBRIJA

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concurrent use with nonselective monoamine oxidase inhibitors (MAOIs) (e.g. phenelzine and tranylcypromine), B.) Recent use (within 2 weeks) with a nonselective MAOI
Required Medical Information	Must meet all of the following A.) Diagnosis of Parkinson's disease and used for intermittent treatment of off episodes, and B.) Concurrent therapy with carbidopa/levodopa
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

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, or C.) Intravenous administration
Required Medical Information	Diagnosis of one of the following A.) Severe primary insulin-like growth factor-1 (IGF-1) deficiency and utilized for pediatric treatment of growth failure, or B.) Growth hormone (GH) gene deletion and patient has developed neutralizing antibodies to GH and utilized for pediatric treatment of growth failure
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

INLYTA

Products Affected

- INLYTA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following 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), or B.) Advanced renal cell carcinoma and used as first-line therapy in combination with avelumab or pembrolizumab
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

INQOVI

Products Affected

- INQOVI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of myelodysplastic syndromes (MDS), including previously treated and untreated, de novo and secondary MDS with the following French-American-British subtypes (refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, and chronic myelomonocytic leukemia [CMML]) and intermediate-1, intermediate-2, and high-risk International Prognostic Scoring System groups
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

INREBIC

Products Affected

- INREBIC

PA Criteria	Criteria Details
Exclusion Criteria	None
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 and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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, or B.) Known or suspected estrogen-dependent neoplasia
Required Medical Information	Diagnosis of one of the following A.) moderate to severe dyspareunia due to menopause, or B.) atrophic vaginitis due to menopause
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Initial: 3 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ISTURISA

Products Affected

- ISTURISA ORAL TABLET 1 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Cushing's disease in patients for whom pituitary surgery is not an option or has not been curative
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ITRACONAZOLE

Products Affected

- *itraconazole oral*

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.), C.) Concurrent use of CYP2D6 inhibitors (e.g., bupropion, fluoxetine, paroxetine, quinidine, terbinafine), D.) Renal or hepatic impairment and concomitant use of colchicine, fesoterodine, solifenacin, or telithromycin, E.) Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Systemic fungal infection (e.g., aspergillosis, histoplasmosis, blastomycosis), or B.) Onychomycosis confirmed by one of the following: positive potassium hydroxide (KOH) preparation, fungal culture, or nail biopsy
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ITRACONAZOLE SOLN

Products Affected

- *itraconazole oral*

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.), C.) Concurrent use of CYP2D6 inhibitors (e.g., bupropion, fluoxetine, paroxetine, quinidine, terbinafine), D.) Renal or hepatic impairment and concomitant use of colchicine, fesoterodine, solifenacin, or telithromycin, E.) Pregnancy
Required Medical Information	Diagnosis of candidiasis (esophageal or oropharyngeal) that is refractory to treatment with fluconazole
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IVABRADINE

Products Affected

- *ivabradine hcl*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Decompensated acute heart failure, B.) hypotension (i.e. blood pressure less than 90/50 mmHg), C.) sick sinus syndrome or sinoatrial block or 3rd degree AV block (unless a functioning demand pacemaker is present), D.) bradycardia (i.e., resting heart rate less than 60 bpm prior to treatment), E.) Severe hepatic impairment (Child-Pugh C), F.) Pacemaker dependent (heart rate maintained exclusively by the pacemaker), G.) Concomitant use of strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of one of the following A.) Adult patients with stable, symptomatic chronic heart failure with left ventricular ejection fraction 35% or less, who are in sinus rhythm with resting heart rate 70 beats per minute or more and either are on maximally tolerated doses of beta-blockers or have a contraindication to beta-blocker use, or B.) Pediatric patients with stable, symptomatic heart failure due to dilated cardiomyopathy and are in sinus rhythm with an elevated heart rate
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

IVERMECTIN

Products Affected

- *ivermectin oral*

PA Criteria	Criteria Details
Exclusion Criteria	Prevention or treatment of COVID-19
Required Medical Information	Diagnosis of one of the following: A.) Strongyloidiasis of the intestinal tract or B.) Onchocerciasis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	1 month
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

IWILFIN

Products Affected

- IWILFIN

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of high-risk neuroblastoma to be used to reduce the risk of relapse in adult and pediatric patients who have demonstrated at least a partial response to prior multiagent, multimodality therapy including anti-GD2 immunotherapy
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

JAKAFI

Products Affected

- JAKAFI

PA Criteria	Criteria Details
Exclusion Criteria	None
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 AND patient has had an inadequate response to or is intolerant of hydroxyurea, C.) Acute graft versus host disease AND disease is refractory to steroid therapy, or D.) Chronic graft-versus-host disease (cGVHD) after failure of corticosteroid therapy (alone or in combination with a calcineurin inhibitor) and up to one additional line of systemic therapy
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

JAYPIRCA

Products Affected

- JAYPIRCA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) relapsed or refractory mantle cell lymphoma (MCL) and is being used after at least two lines of systemic therapy, including a BTK inhibitor or B.) chronic lymphocytic leukemia or small lymphocytic lymphoma who have received at least 2 prior lines of therapy, including a Bruton tyrosine kinase inhibitor and a B-cell lymphoma 2 inhibitor.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

JOENJA

Products Affected

- JOENJA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of activated phosphoinositide 3-kinase (PI3K) delta syndrome
Age Restrictions	12 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KALYDECO

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystic fibrosis (CF) 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	None
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KESIMPTA

Products Affected

- KESIMPTA

PA Criteria	Criteria Details
Exclusion Criteria	Active Hepatitis B infection
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KINERET

Products Affected

- KINERET SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe active rheumatoid arthritis and patient has trial and failure, contraindication, or intolerance to two preferred products, (i.e. Humira, Enbrel, Rinvoq), B.) Cryopyrin-associated periodic syndromes (i.e., neonatal-onset multisystem inflammatory disease), or C.) Deficiency of interleukin-1 receptor antagonist
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KISQALI

Products Affected

- KISQALI (200 MG DOSE)
- KISQALI (400 MG DOSE)
- KISQALI (600 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced or metastatic hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer and one of the following A.) The patient is a pre-or perimenopausal woman or man and the requested drug will be used in combination with an aromatase inhibitor as initial endocrine-based therapy, B.) The patient is a postmenopausal woman or man, the requested drug will be used in combination with an aromatase inhibitor as initial endocrine-based therapy, and the patient has experienced disease progression, an intolerable adverse event, or contraindication to Ibrance (palbociclib) or Verzenio (abemaciclib), C.) The patient is a pre-or perimenopausal woman or man and the requested drug is being used with fulvestrant as initial endocrine-based therapy, or D.) The patient is a postmenopausal woman or man, the requested drug is being used following disease progression on endocrine therapy, and the patient has experienced disease progression, an intolerable adverse event, or contraindication to Ibrance (palbociclib) or Verzenio (abemaciclib)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Part B Prerequisite	No

Formulary ID: 24459 version 18
Last Updated: 09/24/2024
Effective: 10/01/2024

KISQALI FEMARA

Products Affected

- KISQALI FEMARA (200 MG DOSE)
- KISQALI FEMARA (400 MG DOSE)
- KISQALI FEMARA (600 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced or metastatic hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer and one of the following A.) The patient is pre-or perimenopausal woman or male and the requested drug will be used as initial endocrine-based therapy, B.) The patient is postmenopausal, the requested drug will be used as initial endocrine-based therapy, and the patient has experienced disease progression, an intolerable adverse event, or contraindication to Ibrance (palbociclib) or Verzenio (abemaciclib)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KOSELUGO

Products Affected

- KOSELUGO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of neurofibromatosis type 1 (NF1) in a patient who has symptomatic, inoperable plexiform neurofibromas (PN)
Age Restrictions	2 years of age to 17 years of age
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

KRAZATI

Products Affected

- KRAZATI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following: A.) KRAS G12C-mutated locally advanced or metastatic non-small cell lung cancer (NSCLC) as determined by an FDA-approved test and patient has received at least one prior systemic therapy, or B.) KRAS G12C-mutated locally advanced or metastatic colorectal cancer in combination with cetuximab, as determined by an FDA-approved test, who have received prior treatment with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LAPATINIB

Products Affected

- *lapatinib ditosylate*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced or metastatic breast cancer with tumors that overexpress human epidermal growth factor receptor 2 (HER2) AND meets one of the following A.) Used in combination with capecitabine in a patient who has received prior therapy including an anthracycline, a taxane, and trastuzumab, OR B.) Used in combination with letrozole in a postmenopausal female for whom hormonal therapy is indicated
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LENALIDOMIDE

Products Affected

- *lenalidomide*

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 or without additional cytogenetic abnormalities, 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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LENVIMA

Products Affected

- LENVIMA (10 MG DAILY DOSE)
- LENVIMA (12 MG DAILY DOSE)
- LENVIMA (14 MG DAILY DOSE)
- LENVIMA (18 MG DAILY DOSE)
- LENVIMA (20 MG DAILY DOSE)
- LENVIMA (24 MG DAILY DOSE)
- LENVIMA (4 MG DAILY DOSE)
- LENVIMA (8 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
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, in combination with everolimus, following one prior anti-angiogenic therapy, C.) Unresectable hepatocellular carcinoma, first-line therapy, D.) Advanced endometrial carcinoma that is not microsatellite instability-high or mismatch repair deficient, in combination with pembrolizumab, when disease has progressed following prior systemic therapy and patient is not a candidate for curative surgery or radiation, or E.) Advanced renal cell carcinoma, in combination with pembrolizumab and used as first-line therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LEUKINE

Products Affected

- LEUKINE INJECTION SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Exclusion Criteria	None
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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LEUPROLIDE

Products Affected

- ELIGARD
- *leuprolide acetate (3 month)*
- *leuprolide acetate injection*
- LUPRON DEPOT (1-MONTH)
- LUPRON DEPOT (3-MONTH)
- LUPRON DEPOT (4-MONTH)
- LUPRON DEPOT (6-MONTH)
- LUPRON DEPOT-PED (1-MONTH) INTRAMUSCULAR KIT 7.5 MG
- LUPRON DEPOT-PED (3-MONTH) INTRAMUSCULAR KIT 11.25 MG
- LUPRON DEPOT-PED (6-MONTH)

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Pregnancy, B.) Undiagnosed abnormal uterine bleeding
Required Medical Information	Must meet one of the following: 1.) Eligard only: Advanced or metastatic prostate cancer, 2.) For Lupron depot and leuprolide products only: A.) Advanced or metastatic prostate cancer and patient has failed or is intolerant to Eligard (7.5 mg 1-month, 22.5 mg 3-month, 30 mg 4-month, & 45 mg 6-month depots only), B.) Endometriosis (3.75 mg 1-month & 11.25 mg 3-month depots only), C.) Anemia due to uterine leiomyomata (Fibroids) (3.75 mg 1-month & 11.25 mg 3-month depots only) and patient is preoperative, or D.) Central precocious puberty (idiopathic or neurogenic) in children
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

L-GLUTAMINE

Products Affected

- *l-glutamine oral packet*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of sickle cell disease AND one of the following 1.) Patient has acute complications and is being treated with Hydroxyurea, or 2.) Patient has acute complications and is unable to tolerate Hydroxyurea
Age Restrictions	5 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LIDOCAINE PATCH

Products Affected

- lidocaine external patch 5 %

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Pain associated with diabetic neuropathy, B.) Pain associated with cancer-related neuropathy, C.) Post-herpetic neuralgia, D.) Chronic back pain, or E.) Osteoarthritis of the knee or hip
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LINEZOLID

Products Affected

- *linezolid intravenous solution 600 mg/300ml*
- *linezolid oral*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use of an MAOI, or B.) Use within 14 days of discontinuing an MAOI
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	None
Prescriber Restrictions	None
Coverage Duration	1 month
Other Criteria	IV formulation: B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

LONSURF

Products Affected

- LONSURF

PA Criteria	Criteria Details
Exclusion Criteria	None
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, or B.) Metastatic gastric or gastroesophageal junction 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
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LORBRENA

Products Affected

- LORBRENA

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with strong CYP3A4 inducers
Required Medical Information	Diagnosis of anaplastic lymphoma kinase (ALK)-positive metastatic non-small cell lung cancer (NSCLC) 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LUMAKRAS

Products Affected

- LUMAKRAS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of KRAS G12C-mutated locally advanced or metastatic non-small cell lung cancer (NSCLC) as determined by an FDA-approved test, and patient has received at least one prior systemic therapy (e.g., immune checkpoint inhibitor, platinum-based chemotherapy)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LUPKYNIS

Products Affected

- LUPKYNIS

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of strong CYP3A4 inhibitors (e.g., ketoconazole, itraconazole, clarithromycin)
Required Medical Information	Diagnosis of systemic lupus erythematosus (SLE) with active lupus nephritis (LN) Classes III, IV, V (alone or in combination), and all of the following: 1.) Baseline renal function of 45 mL/min/1.73 m ² or greater, 2.) Will be used in combination with a background immunosuppressive therapy regimen (e.g. mycophenolate, oral steroids, etc). Renewal: Documentation of positive clinical response to therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a rheumatologist or nephrologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LYNPARZA

Products Affected

- LYNPARZA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) HER2-negative, deleterious or suspected deleterious germline BRCA mutated high-risk early or metastatic breast cancer AND patient has been previously treated with chemotherapy in neoadjuvant, adjuvant, or metastatic setting, B.) 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), C.) 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, D.) 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, E.) Advanced epithelial ovarian, fallopian tube, or primary peritoneal cancer in patients who are in complete or partial response to first-line platinum-based chemotherapy and whose cancer is associated with homologous recombination deficiency positive status defined by either a deleterious or suspected deleterious BRCA-mutation and/or genomic instability AND are using in combination with bevacizumab for maintenance treatment, F.) Deleterious or suspected deleterious germline or somatic homologous recombination repair gene mutated metastatic castration-resistant prostate cancer in patients who have progressed following prior treatment with enzalutamide or abiraterone, or G.) Deleterious or suspected deleterious BRCA-mutated metastatic castration-resistant prostate cancer in combination with abiraterone and prednisone or prednisolone
Age Restrictions	18 years of age and older

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

LYTGOBI

Products Affected

- LYTGOBI (12 MG DAILY DOSE)
- LYTGOBI (16 MG DAILY DOSE)
- LYTGOBI (20 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of unresectable, locally advanced or metastatic intrahepatic cholangiocarcinoma harboring fibroblast growth factor receptor 2 (FGFR2) gene fusions or other rearrangements and previously treated
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

MATULANE

Products Affected

- MATULANE

PA Criteria	Criteria Details
Exclusion Criteria	Inadequate marrow reserve
Required Medical Information	Diagnosis of Hodgkin's Disease, Stages III and IV and used in combination with other anticancer drugs
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

MAYZENT

Products Affected

- MAYZENT
- MAYZENT STARTER PACK

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) CYP2C9*3/*3 genotype, B.) In the last 6 months experienced myocardial infarction, unstable angina, stroke, TIA, decompensated heart failure requiring hospitalization, Class III-IV heart failure, or C.) 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, including clinically isolated syndrome, relapsing-remitting disease, or active secondary progressive disease and the following A.) Patients with relapsing forms of multiple sclerosis have history of/or contraindication to Avonex, Betaseron, Glatiramer, Fingolimod, or Dimethyl Fumarate
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

MEKINIST

Products Affected

- MEKINIST

PA Criteria	Criteria Details
Exclusion Criteria	None
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 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, D.) Metastatic non-small cell lung cancer, with BRAF V600E mutation, in combination with dabrafenib, E.) Unresectable or metastatic solid tumors with BRAF V600E mutation, in combination with dabrafenib, and have progressed following prior treatment and have no satisfactory alternative treatment options, F.) Low-grade glioma with a BRAF V600E mutation and require systemic therapy, in combination with dabrafenib
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

MEKTOVI

Products Affected

- MEKTOVI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Unresectable or metastatic malignant melanoma with documented BRAF V600E or V600K mutation as detected by an FDA approved test AND used in combination with encorafenib or B.) Metastatic non-small cell lung cancer with a BRAF V600E 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

METHOXSALEN

Products Affected

- *methoxsalen rapid*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Aphakia, B.) Melanoma or a history of melanoma, C.) Invasive squamous cell carcinomas, or D.) History of a light sensitive disease/skin photosensitivity disorder such systemic lupus erythematosus (SLE), porphyria cutanea tarda, erythropoietic protoporphyria, variegate porphyria, xeroderma pigmentosum or albinism
Required Medical Information	Diagnosis of one of the following A.) Psoriasis, or B.) Vitiligo
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist, immunologist, or dermatologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

MIFEPRISTONE

Products Affected

- *mifepristone oral tablet 300 mg*

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 endogenous Cushing syndrome in patients with type 2 diabetes mellitus or glucose intolerance and must meet all of the following 1.) Used to control hyperglycemia secondary to hypercortisolism, and 2.) Patient has failed 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

MIGLUSTAT

Products Affected

- *miglustat*
- YARGESA

PA Criteria	Criteria Details
Exclusion Criteria	None
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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

MS INTERFERONS

Products Affected

- AVONEX PEN INTRAMUSCULAR AUTO-INJECTOR KIT
- AVONEX PREFILLED INTRAMUSCULAR PREFILLED SYRINGE KIT
- BETASERON SUBCUTANEOUS KIT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

NERLYNX

Products Affected

- NERLYNX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Early stage HER2-positive breast cancer and used following adjuvant trastuzumab therapy, or B.) Advanced or metastatic HER2-positive breast cancer, used in combination with capecitabine, AND patient has received 2 or more prior anti-HER2-based regimens in the metastatic setting
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

NICOTINE

Products Affected

- NICOTROL

PA Criteria	Criteria Details
Exclusion Criteria	Beneficiary continues to smoke
Required Medical Information	Beneficiary must have successful cessation at 12 weeks for one additional authorization period of 12 weeks.
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Initial: 12 weeks, Renewal: 12 weeks (Maximum 24 weeks treatment)
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

NINLARO

Products Affected

- NINLARO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of multiple myeloma, used in combination with lenalidomide and dexamethasone, AND patient has history of at least 1 prior therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

NITISINONE

Products Affected

- *nitisinone oral capsule 10 mg, 2 mg, 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of hereditary tyrosinemia type 1 confirmed by one of the following A.) Biochemical testing (e.g., detection of succinylacetone in urine), or B.) DNA testing (mutation analysis)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

NUBEQA

Products Affected

- NUBEQA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Non-metastatic, castration-resistant prostate cancer (nmCRPC) or B.) Metastatic hormone-sensitive prostate cancer in combination with docetaxel. For treatment of nmCRPC, one of the following applies 1.) Used in combination with a gonadotropin-releasing hormone (GnRH) analog or 2) Patient has received bilateral orchiectomy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

NUCALA

Products Affected

- NUCALA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Severe asthma with eosinophilic phenotype and used as an adjunct treatment, B.) Eosinophilic granulomatosis with polyangiitis (EGPA), C.) Hypereosinophilic syndrome lasting at least 6 months without an identifiable non-hematologic secondary cause, or D.) Chronic rhinosinusitis with nasal polyps and used as an adjunct treatment
Age Restrictions	6 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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, F.) History of quinine-, mefloquine-, or quinidine-induced thrombocytopenia, bone marrow depression, or lupus-like syndrome
Required Medical Information	Diagnosis of pseudobulbar affect
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

NUPLAZID

Products Affected

- NUPLAZID ORAL CAPSULE
- NUPLAZID ORAL TABLET 10 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Parkinson's disease and both of the following apply A.) Used for treatment of hallucinations and/or delusions associated with Parkinson's disease psychosis, and B.) Diagnosis of Parkinson's disease was made prior to the onset of psychotic symptoms
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

OCTREOTIDE

Products Affected

- *octreotide acetate injection solution 100 mcg/ml, 1000 mcg/ml, 200 mcg/ml, 50 mcg/ml, 500 mcg/ml*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Acromegaly confirmed by high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range and patient has had inadequate response to or is ineligible for surgery, radiation, or bromocriptine mesylate, or B.) Metastatic carcinoid syndrome with associated diarrhea or flushing, or C.) Vasoactive intestinal peptide-secreting tumors (VIPomas) with associated diarrhea
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

OFEV

Products Affected

- OFEV

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Idiopathic pulmonary fibrosis (IPF), B.) Systemic sclerosis-associated interstitial lung disease (ILD), or C.) Chronic fibrosing interstitial lung disease with a progressive phenotype
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

OGSIVEO

Products Affected

- OGSIVEO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of progressing desmoid tumors who require systemic treatment
Age Restrictions	18 years or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	end of plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

OJEMDA

Products Affected

- OJEMDA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of relapsed or refractory pediatric low-grade glioma harboring a BRAF fusion or rearrangement, or BRAF V600 mutation
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

OJJAARA

Products Affected

- OJJAARA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of intermediate or high-risk myelofibrosis (MF), including primary MF or secondary MF [postpolycythemia vera (PV) and post-essential thrombocythemia (ET)], in adults with anemia.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ONUREG

Products Affected

- ONUREG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of acute myeloid leukemia (AML) used in maintenance treatment for adult patients who achieved first complete remission (CR) or complete remission with incomplete blood count recovery (CRi) following intensive induction chemotherapy and are not able to complete intensive curative therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

OPSUMIT

Products Affected

- OPSUMIT

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of pulmonary arterial hypertension (WHO Group I), 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.)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ORGOVYX

Products Affected

- ORGOVYX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced prostate cancer
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ORKAMBI

Products Affected

- ORKAMBI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystic fibrosis (CF) with documented homozygous F508del mutation confirmed by FDA-approved CF mutation test
Age Restrictions	None
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ORSERDU

Products Affected

- ORSERDU

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced or metastatic, ER-positive, HER2-negative, ESR1-mutated, breast cancer in postmenopausal women or adult man after at least 1 line of endocrine therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

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 deep vein thrombosis (DVT), pulmonary embolism (PE), or a history of these conditions, D.) Active arterial thromboembolic disease (e.g. stroke, myocardial infarction) or a history of these conditions, or E.) 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
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

OTEZLA

Products Affected

- OTEZLA ORAL TABLET 30 MG
- OTEZLA ORAL TABLET THERAPY PACK 10 & 20 & 30 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following: A.) Active psoriatic arthritis and patient has trial and failure or intolerance or contraindication to two preferred products, (i.e. Cosentyx, Humira, Enbrel, Skyrizi, Stelara, Rinvoq), B.) Moderate to severe plaque psoriasis, patient is a candidate for phototherapy or systemic therapy, and patient has trial and failure or intolerance or contraindication to two preferred products, (i.e. Cosentyx, Humira, Enbrel, Skyrizi, Stelara), C.) Mild plaque psoriasis, patient is a candidate for phototherapy or systemic therapy, and patient has trial and failure or intolerance or contraindication to at least one topical psoriasis product (e.g., medium to high potency corticosteroid and/or vitamin D analog), or D.) Behcet's Disease and patient has active oral ulcers
Age Restrictions	6 years of age and older
Prescriber Restrictions	PsA: Prescribed by or in consultation with a dermatologist or rheumatologist. Plaque psoriasis: Prescribed by or in consultation with a dermatologist.
Coverage Duration	plan year
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PANRETIN

Products Affected

- PANRETIN

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of AIDS-related Kaposi's sarcoma and both of the following 1.) Used to treat cutaneous lesions, and 2.) Systemic anti-Kaposi's Sarcoma therapy is not indicated (e.g., patient does not have more than 10 new KS lesions in the prior month, symptomatic lymphedema, symptomatic pulmonary KS, or symptomatic visceral involvement)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or HIV specialist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PAZOPANIB

Products Affected

- *pazopanib hcl*

PA Criteria	Criteria Details
Exclusion Criteria	None
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
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PEGYLATED INTERFERON

Products Affected

- PEGASYS SUBCUTANEOUS SOLUTION 180 MCG/ML
- PEGASYS SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Autoimmune hepatitis, B.) Hepatic decompensation (Child-Pugh score greater than 6 (Class B and C) in cirrhotic patients before treatment, OR hepatic decompensation (Child-Pugh score greater than or equal to 6) in cirrhotic patients co-infected with hepatitis C and HIV before treatment, C.) Hypersensitivity reactions, including urticaria, bronchoconstriction, anaphylaxis, or Stevens-Johnson syndrome to alfa interferons or any component of the product, or D.) Pregnancy with concomitant ribavirin use
Required Medical Information	Diagnosis of one of the following A.) Chronic hepatitis B infection, or B.) Chronic hepatitis C and required criteria will be applied consistent with current AASLD-IDSa guidance with compensated liver disease
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, hepatologist, or infectious disease specialist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PEMAZYRE

Products Affected

- PEMAZYRE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Previously treated, unresectable locally advanced or metastatic cholangiocarcinoma with confirmed fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement as detected by an FDA-approved test or B.) Relapsed or refractory myeloid/lymphoid neoplasms with fibroblast growth factor receptor 1 rearrangement
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist, gastroenterologist, or hepatologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PHENYL BUTYRATE

Products Affected

- sodium phenylbutyrate oral powder 3 gm/tsp
- sodium phenylbutyrate oral tablet

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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PIQRAY

Products Affected

- PIQRAY (200 MG DAILY DOSE)
- PIQRAY (250 MG DAILY DOSE)
- PIQRAY (300 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of hormone receptor (HR) positive, HER2-negative, PIK3CA-mutated, advanced or metastatic breast cancer and must meet all of the following 1.) Used in combination with fulvestrant, 2.) Disease has progressed on or after an endocrine-based regimen, and 3.) Patient is a male or postmenopausal female
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PIRFENIDONE

Products Affected

- *pirfenidone*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of idiopathic pulmonary fibrosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

POMALYST

Products Affected

- POMALYST

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) AIDS-related Kaposi sarcoma and patient has failure on highly active antiretroviral therapy (HAART), B.) Kaposi sarcoma in HIV-negative adults, or C.) Multiple myeloma and in combination with dexamethasone in adults who have received at least 2 prior therapies (including lenalidomide and a proteasome inhibitor) and have demonstrated disease progression on or within 60 days of completion of the last therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

POSACONAZOLE

Products Affected

- NOXAFIL ORAL PACKET
- *posaconazole oral*

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, B.) Patient is severely immunocompromised and requires prophylaxis of invasive aspergillosis or candidiasis due to high risk of infection, or C.) Invasive aspergillosis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	12 weeks
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

POSACONAZOLE SUSPENSION

Products Affected

- *posaconazole oral*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Oropharyngeal candidiasis, B.) Patient is severely immunocompromised and requires prophylaxis of invasive aspergillosis or candidiasis due to high risk of infection, or C.) Invasive aspergillosis
Age Restrictions	13 years of age and older
Prescriber Restrictions	None
Coverage Duration	12 weeks
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PREVYMIS

Products Affected

- PREVYMIS ORAL

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use with pimozide or ergot alkaloids (ergotamine, dihydroergotamine), B.) Concomitant use with pitavastatin or simvastatin when coadministered with cyclosporine
Required Medical Information	Diagnosis of one of the following A.) Prophylaxis of cytomegalovirus (CMV) infection and disease in adult CMV-seropositive recipients [R+] of an allogeneic hematopoietic stem cell transplant, or B.) Prophylaxis of CMV disease in adult kidney transplant recipients at high risk (Donor CMV seropositive/Recipient CMV seronegative [D+/R-])
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PROMACTA

Products Affected

- PROMACTA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic idiopathic thrombocytopenic purpura (ITP), B.) Chronic hepatitis C infection associated thrombocytopenia, or C.) Severe aplastic anemia with insufficient response to immunosuppressive therapy or in combination with standard immunosuppressive therapy
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

QINLOCK

Products Affected

- QINLOCK

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced gastrointestinal stromal tumor (GIST) and patient has received prior treatment with 3 or more kinase inhibitors, including imatinib
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

QUININE SULFATE

Products Affected

- *quinine sulfate oral*

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, G.) Use solely for treatment or prevention of nocturnal leg cramps
Required Medical Information	Diagnosis of one of the following A.) uncomplicated Plasmodium falciparum malaria, B.) uncomplicated Plasmodium vivax malaria, or C.) babesiosis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	1 month
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

RAVICTI

Products Affected

- RAVICTI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of urea cycle disorders
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

REPATHA

Products Affected

- REPATHA
- REPATHA SURECLICK
- REPATHA PUSHTRONEX SYSTEM

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) primary hyperlipidemia including heterozygous familial hypercholesterolemia (HeFH), B.) homozygous familial hypercholesterolemia, C.) established cardiovascular disease and myocardial infarction prophylaxis, stroke prophylaxis, or coronary revascularization prophylaxis is required, or D.) clinical atherosclerotic cardiovascular disease (CVD) as defined as one of the following 1.) acute coronary syndrome, 2.) history of myocardial infarction, 3.) stable/unstable angina, 4.) coronary or other arterial revascularization, 5.) stroke, 6.) transient ischemic stroke (TIA), or 7.) peripheral arterial disease presumed to be atherosclerotic region
Age Restrictions	10 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

RETEVMO

Products Affected

- RETEVMO ORAL CAPSULE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Advanced or metastatic RET-mutant medullary thyroid cancer (MTC) in patients who require systemic therapy, B.) Metastatic RET fusion-positive non-small cell lung cancer (NSCLC), C.) Advanced or metastatic RET fusion-positive thyroid cancer in patients who require systemic therapy and are refractory to radioactive iodine, if appropriate, or D.) Locally advanced or metastatic solid tumors with a RET gene fusion that have progressed on or following prior systemic treatment or who have no satisfactory alternative treatment options
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

REZLIDHIA

Products Affected

- REZLIDHIA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of relapsed or refractory acute myeloid leukemia (AML) with a susceptible IDH1 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
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

REZUROCK

Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of chronic graft-vs-host disease and patient has failed at least 2 prior lines of systemic therapy.
Age Restrictions	12 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

RILUZOLE

Products Affected

- *riluzole*
- TEGLUTIK

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of amyotrophic lateral sclerosis (ALS)
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

RINVOQ

Products Affected

- RINVOQ

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe rheumatoid arthritis, B.) Active psoriatic arthritis, C.) Moderate to severe atopic dermatitis and patient has trial/failure, contraindication, or intolerance to two of the following 1.) Topical corticosteroid and/or 2.) Topical calcineurin inhibitor, D.) Moderately to severely active ulcerative colitis who have had an inadequate response or intolerance to one or more tumor necrosis factor blockers, E.) Active ankylosing spondylitis who have had an inadequate response or intolerance to one or more tumor necrosis factor blockers, F.) Active nonradiographic axial spondyloarthritis with objective signs of inflammation who have had an inadequate response or intolerance to tumor necrosis factor blocker therapy, G.) Moderate to severe active Crohn's disease who have had an inadequate response or intolerance to tumor necrosis factor blocker therapy, or H.) Active polyarticular juvenile idiopathic arthritis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Part B Prerequisite	No

Formulary ID: 24459 version 18
Last Updated: 09/24/2024
Effective: 10/01/2024

RINVOQ LQ

Products Affected

- RINVOQ LQ

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Active psoriatic arthritis, or B.) Active polyarticular juvenile idiopathic arthritis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

RIVFLOZA

Products Affected

- RIVFLOZA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Primary hyperoxaluria type 1 and the patient has relatively preserved kidney function (eGFR is greater than or equal to 30mL/min/1.73m ²)
Age Restrictions	9 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ROZLYTREK

Products Affected

- ROZLYTREK

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) ROS1-positive metastatic non-small cell lung cancer (NSCLC), or B.) Solid tumors that have a neurotrophic tyrosine receptor kinase (NTRK) gene fusion without a known acquired resistance mutation, are metastatic or where surgical resection is likely to result in severe morbidity, and have either progressed following treatment or have no satisfactory alternative therapy
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

RUBRACA

Products Affected

- RUBRACA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer, used as maintenance treatment, and patient is in complete or partial response to platinum-based chemotherapy, or B.) Deleterious BRCA mutation (germline and/or somatic)-associated metastatic castration-resistant prostate cancer and patient has been treated with androgen receptor-directed therapy and a taxane-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

RYDAPT

Products Affected

- RYDAPT

PA Criteria	Criteria Details
Exclusion Criteria	None
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SAPROPTERIN

Products Affected

- *sapropterin dihydrochloride oral packet*
- *sapropterin dihydrochloride oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of hyperphenylalaninemia (HPA) caused by tetrahydrobiopterin (BH4)-responsive phenylketonuria (PKU)
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Initial: 2 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

SCEMBLIX

Products Affected

- SCEMBLIX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP), previously treated with two or more tyrosine kinase inhibitors (TKIs), or B.) Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP) with the T315I mutation
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SIGNIFOR

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Cushing disease and patient has had inadequate response to or is not a candidate for surgery. For renewal: Documentation of a clinically meaningful reduction in 24-hour urinary free cortisol (UFC) levels or improvement in signs or symptoms of the disease
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SILDENAFIL

Products Affected

- *sildenafil citrate oral tablet 20 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Nitrate therapy, including intermittent use, B.) Concomitant use with riociguat or other guanylate cyclase stimulators, C.) Concomitant use with HIV protease inhibitors or elvitegravir/cobicistat/tenofovir/emtricitabine
Required Medical Information	Diagnosis of pulmonary arterial hypertension (WHO Group I), 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.)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SKYRIZI

Products Affected

- SKYRIZI PEN
- SKYRIZI SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Moderate to severe plaque psoriasis and patient is a candidate for systemic therapy or phototherapy, B.) Active psoriatic arthritis, C.) Moderately to severely active Crohn's disease, or D.) Moderately to severely active ulcerative colitis
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

SODIUM OXYBATE

Products Affected

- *sodium oxybate*

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 one of the following A.) Narcolepsy with excessive daytime drowsiness and has trial of/or contraindication to a central nervous system (CNS) stimulant drug (e.g., amphetamine, dextroamphetamine, methylphenidate) or a CNS wakefulness promoting drug (e.g., armodafinil, modafinil), or B.) Cataplexy and narcolepsy
Age Restrictions	7 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

SOHONOS

Products Affected

- SOHONOS

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of fibrodysplasia ossificans progressiva (FOP)
Age Restrictions	8 years and older for females and 10 years and older for males
Prescriber Restrictions	None
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SOMAVERT

Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of acromegaly confirmed by high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range and patient has had an 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SORAFENIB

Products Affected

- *sorafenib tosylate*

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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SPRYCEL

Products Affected

- SPRYCEL

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Newly diagnosed Philadelphia chromosome-positive (Ph+) chronic myeloid leukemia (CML) in chronic phase, B.) Chronic, accelerated, or myeloid or lymphoid blast phase Ph+ CML with resistance or intolerance to prior therapy, C.) Ph+ acute lymphoblastic leukemia (ALL) with resistance or intolerance to prior therapy, or D.) Newly diagnosed Ph+ ALL in combination with chemotherapy
Age Restrictions	1 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

STELARA

Products Affected

- STELARA SUBCUTANEOUS SOLUTION 45 MG/0.5ML
- STELARA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Moderate to severely active Crohn disease, B.) Moderate to severe plaque psoriasis, C.) Active psoriatic arthritis, or D.) Moderate to severe active ulcerative colitis
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	Screening for latent tuberculosis infection is required prior to initiation of treatment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

STIVARGA

Products Affected

- STIVARGA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Metastatic colorectal cancer in patients previously treated with fluoropyrimidine, oxaliplatin, and irinotecan containing chemotherapy, anti-VEGF therapy, and if RAS wild type, anti-EGFR therapy, B.) Liver carcinoma in patients previously treated with sorafenib, or C.) Locally advanced, unresectable or metastatic gastrointestinal stromal tumor (GIST) after treatment with imatinib and sunitinib
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SUNITINIB

Products Affected

- *sunitinib malate*

PA Criteria	Criteria Details
Exclusion Criteria	None
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 carcinoma 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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

SYMDEKO

Products Affected

- SYMDEKO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystic fibrosis (CF) and must meet one of the following 1.) Patient is homozygous for the F508del mutation, or 2.) 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 or prescribing practitioner is from a CF center accredited by the Cystic Fibrosis Foundation
Coverage Duration	Initial: 6 months, Renewal: 12 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

SYMLIN

Products Affected

- SYMLINPEN 120 SUBCUTANEOUS SOLUTION PEN-INJECTOR
- SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Confirmed diagnosis of gastroparesis, B.) Hypoglycemia unawareness
Required Medical Information	Diagnosis of type 1 or type 2 diabetes mellitus and patient uses mealtime insulin therapy and has failed to achieve desired glucose control
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TABRECTA

Products Affected

- TABRECTA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic non-small cell lung cancer (NSCLC) in patients whose tumors have a mutation that leads to mesenchymal-epithelial transition (MET) exon 14 skipping 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TAFINLAR

Products Affected

- TAFINLAR

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Locally advanced or metastatic anaplastic thyroid carcinoma with BRAF V600E mutation, in combination with trametinib and no satisfactory locoregional treatment options, B.) Metastatic non-small cell lung cancer with BRAF V600E mutation, in combination with trametinib OR in patients previously treated as monotherapy, C.) Unresectable or metastatic malignant melanoma with BRAF V600E or V600K mutation, D.) Unresectable or metastatic solid tumors with BRAF V600E mutation, in combination with trametinib, and have progressed following prior treatment and have no satisfactory alternative treatment options, or E.) Low-grade glioma with a BRAF V600E mutation and require systemic therapy, in combination with trametinib
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TAGRISO

Products Affected

- TAGRISO

PA Criteria	Criteria Details
Exclusion Criteria	None
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, B.) Metastatic non-small cell lung cancer with T790M EGFR mutation (as confirmed by an FDA-approved test) AND whose disease has progressed on or after EGFR tyrosine kinase inhibitor therapy, C.) Non-small cell lung cancer (NSCLC) with tumor epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations (as confirmed by an FDA-approved test) AND patient requires adjuvant therapy after tumor resection, or D.) First-line treatment of adult patients with locally advanced or metastatic non-small cell lung cancer whose tumors have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R mutations, as detected by an FDA-approved test, in combination with pemetrexed and platinum-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TAKHZYRO

Products Affected

- TAKHZYRO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following and used as routine prophylaxis A.) Hereditary angioedema (HAE) with C1 inhibitor deficiency (Type 1) confirmed by laboratory testing, or B.) HAE with C1 inhibitor dysfunction (Type 2) confirmed by laboratory testing, or C.) HAE with normal C1 inhibitor (Type 3) confirmed by laboratory testing and one of the following 1.) Positive test for an F12, angiopoietin-1, or plasminogen gene mutation, or 2.) Family history of angioedema and the angioedema was refractory to a trial of an antihistamine for at least one month
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a hematologist, immunologist, or allergist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TALZENNA

Products Affected

- TALZENNA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Deleterious or suspected deleterious germline breast cancer susceptibility gene (BRCA)-mutated (gBRCAm), human epidermal growth factor receptor 2 (HER2)-negative locally advanced or metastatic breast cancer, or B.) Homologous recombination repair gene-mutated metastatic castration-resistant prostate cancer in combination with enzalutamide
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TASIGNA

Products Affected

- TASIGNA

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Long QT syndrome, B.) Uncorrected hypokalemia, C.) Uncorrected hypomagnesemia
Required Medical Information	Diagnosis of one of the following A.) Newly diagnosed chronic phase Philadelphia chromosome-positive chronic myelogenous leukemia (CML), B.) Chronic phase or accelerated phase Philadelphia chromosome-positive CML in a patient with resistance or intolerance to prior therapy that included imatinib, or C.) Chronic phase Philadelphia chromosome-positive CML in a patient with resistance or intolerance to prior tyrosine-kinase inhibitor therapy
Age Restrictions	1 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TAVNEOS

Products Affected

- TAVNEOS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of severe active anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis (granulomatosis with polyangiitis [GPA] and microscopic polyangiitis [MPA]) and both of the following apply 1.) Used as adjunctive treatment, and 2.) Used in combination with standard therapy including glucocorticoids
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TAZAROTENE

Products Affected

- *tazarotene external cream*
- *tazarotene external gel*
- TAZORAC EXTERNAL CREAM 0.05 %

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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TAZVERIK

Products Affected

- TAZVERIK

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Metastatic or locally advanced epithelioid sarcoma in patients not eligible for complete resection, B.) Relapsed or refractory follicular lymphoma in patients whose tumors are positive for an EZH2 mutation as detected by an FDA-approved test and who have received at least 2 prior systemic therapies, or C.) Relapsed or refractory follicular lymphoma in patients who have no satisfactory alternative treatment options
Age Restrictions	16 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TEPMETKO

Products Affected

- TEPMETKO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic non-small cell lung cancer (NSCLC) with mesenchymal-epithelial transition (MET) exon 14 skipping alterations
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TERIPARATIDE

Products Affected

- *teriparatide (recombinant) subcutaneous solution pen-injector 620 mcg/2.48ml*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Osteoporosis in postmenopausal female patient with high risk for fracture and patient has contraindication or has tried/had inadequate response to a bisphosphonate or Tymlos, B.) Primary or hypogonadal osteoporosis in male patient with high risk for fracture and patient has contraindication or has tried/had inadequate response to a bisphosphonate, or C.) Osteoporosis due to associated sustained systemic glucocorticoid therapy in patient with high risk for fracture and patient has contraindication or has tried/had inadequate response to a bisphosphonate
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Initial: 12 months, Renewal: 12 months (Maximum 24 month treatment per patient lifetime)
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TETRABENAZINE

Products Affected

- *tetrabenazine*

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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

THALOMID

Products Affected

- THALOMID

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Multiple myeloma that is newly diagnosed, or B.) Erythema nodosum leprosum (ENL)
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist, infectious disease specialist, or dermatologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TIBSOVO

Products Affected

- TIBSOVO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsed or refractory acute myeloid leukemia with a susceptible isocitrate dehydrogenase-1 mutation (as detected by an FDA-approved test), B.) Previously treated, locally advanced or metastatic cholangiocarcinoma with an isocitrate dehydrogenase-1 mutation (as detected by an FDA-approved test.), C.) Acute myeloid leukemia (newly-diagnosed) with susceptible isocitrate dehydrogenase-1 mutation and meets one of the following: 1.) Patient is 75 years of age or older, or 2.) Patient has comorbidities that preclude intensive induction chemotherapy, or D.) Relapsed or refractory myelodysplastic syndromes with a susceptible isocitrate dehydrogenase-1 mutation (as detected by an FDA-approved test)
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist, hematologist, hepatologist, or oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TOBI

Products Affected

- TOBI PODHALER

PA Criteria	Criteria Details
Exclusion Criteria	Known sensitivity to any aminoglycoside
Required Medical Information	Diagnosis of cystic fibrosis (confirmed by appropriate diagnostic or genetic testing) and patient has suspected or confirmed <i>Pseudomonas aeruginosa</i> infection in the lungs
Age Restrictions	6 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TOLVAPTAN

Products Affected

- *tolvaptan*

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Diagnosis of Autosomal Dominant Polycystic Kidney Disease (ADPKD), B.) Urgent need to raise serum sodium acutely, C.) Inability to sense or appropriately respond to thirst, D.) Hypovolemic hyponatremia, E.) Concomitant use of strong CYP 3A Inhibitors (e.g. clarithromycin, ketoconazole, ritonavir), or F.) Anuria
Required Medical Information	Diagnosis of clinically significant hypervolemic or euvolemic hyponatremia (serum sodium less than 125 mEq/L or less marks hyponatremia that is symptomatic and has resisted correction with fluid restriction), including in patients with heart failure and syndrome of inappropriate antidiuretic hormone (SIADH)
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	1 month
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TOPICAL RETINOIDS

Products Affected

- *adapalene external cream*
- *adapalene external gel 0.3 %*
- *tretinoin external*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of mild to moderate acne vulgaris
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TOPICAL TESTOSTERONE

Products Affected

- testosterone transdermal gel 10 mg/act (2%), 12.5 mg/act (1%), 20.25 mg/1.25gm (1.62%), 20.25 mg/act (1.62%), 25 mg/2.5gm (1%), 40.5 mg/2.5gm (1.62%), 50 mg/5gm (1%)

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Carcinoma of the breast (males only), B.) Known or suspected carcinoma of the prostate, C.) Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Hypogonadotropic hypogonadism, or B.) Primary hypogonadism. Diagnosis of hypogonadism must be confirmed by a low-for-age serum testosterone (total or free) level defined by the normal laboratory reference value
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TOREMIFENE

Products Affected

- *toremifene citrate*

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 and patient must have previous inadequate response or intolerance to tamoxifen
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	6 months
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TORPENZ

Products Affected

- TORPENZ

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Hypersensitivity to everolimus or excipients, or B.) Hypersensitivity to rapamycin derivatives (e.g. sirolimus)
Required Medical Information	Diagnosis of one of the following A.) Renal angiomyolipoma and tuberous sclerosis complex (TSC) not requiring immediate surgery, B.) Advanced hormone receptor-positive, HER2 negative breast cancer in postmenopausal women and taken in combination with exemestane, after failure with letrozole or anastrozole, C.) Subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex in patients who are not candidates for curative surgical resection
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TRELSTAR

Products Affected

- TRELSTAR MIXJECT

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of advanced prostate cancer
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TRIENTINE

Products Affected

- *trientine hcl*

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Wilson's disease in patients that are intolerant to penicillamine
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TRIKAFTA

Products Affected

- TRIKAFTA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of cystic fibrosis (CF) and patient has at least 1 F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene or a mutation in the CFTR gene that is responsive to elexacaftor/tezacaftor/ivacaftor verified by an FDA-cleared CF mutation test
Age Restrictions	2 years of age and older
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TRUQAP

Products Affected

- TRUQAP

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, locally advanced or metastatic breast cancer with 1 or more PIK3CA/AKT1/PTEN-alterations as detected by an FDA-approved test and, A.) patient has had disease progression following 1 or more endocrine-based regimen(s) in the metastatic setting or recurrence on or within 12 months of completing adjuvant therapy, and B.) will be used in combination with fulvestrant injection.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TUKYSA

Products Affected

- TUKYSA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following: A.) advanced unresectable or metastatic HER2-positive breast cancer (including brain metastases) in patients who have received one or more prior anti-HER2-based regimens in the metastatic setting and drug is being used in combination with trastuzumab and capecitabine, or B.) unresectable or metastatic RAS wild-type, HER2-positive colorectal cancer that has progressed following treatment with fluoropyrimidine, oxaliplatin, and irinotecan-based chemotherapy and drug is being used in combination with trastuzumab
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

TURALIO

Products Affected

- TURALIO ORAL CAPSULE 125 MG

PA Criteria	Criteria Details
Exclusion Criteria	None
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 and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

TYMLOS

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of osteoporosis in men or postmenopausal women and one of the following A.) osteoporotic fracture or multiple risk factors for fracture, or B.) previous trial of/or contraindication to bisphosphonate
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Initial: 12 months, Renewal: 12 months (Maximum 24 month treatment per patient lifetime)
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

UBRELVY

Products Affected

- UBRELVY

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of strong CYP3A4 inhibitors (e.g., ketoconazole, itraconazole, clarithromycin)
Required Medical Information	Diagnosis of migraine disorder with or without aura and patient has documented trial, inadequate response, or contraindication to at least 1 generic formulary triptan
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VALCHLOR

Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Exclusion Criteria	None
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	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VANFLYTA

Products Affected

- VANFLYTA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Patient must have all of the following A.) Newly diagnosed acute myeloid leukemia with FLT3-ITD mutation, B.) Used in combination with standard cytarabine and anthracycline induction and cytarabine consolidation, and as maintenance monotherapy following consolidation chemotherapy, and C.) Must be enrolled in the VANFLYTA REMS program
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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 or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

VERQUVO

Products Affected

- VERQUVO

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use of other soluble guanylate cyclase (sGC) stimulators, or B.) Pregnancy
Required Medical Information	Diagnosis of chronic heart failure (HF), NYHA Class II to IV and all of the following 1.) Left ventricular ejection fraction less than 45%, 2.) Previous hospitalization for HF within 6 months or outpatient IV diuretic treatment for HF within 3 months
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VERZENIO

Products Affected

- VERZENIO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, node-positive, early breast cancer and ALL of the following: 1.) Patient is at high risk of recurrence, and 2.) Requested drug will be used in combination with endocrine therapy (tamoxifen or an aromatase inhibitor) for adjuvant treatment, OR B.) Advanced or metastatic, hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer and one of the following 1.) Used in combination with fulvestrant in a patient with disease progression following endocrine therapy, 2.) Used as monotherapy in a patient with disease progression following endocrine therapy and prior chemotherapy in the metastatic setting, or 3.) For postmenopausal women, and men, used as initial endocrine-based treatment in combination with an aromatase inhibitor
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VIGABATRIN

Products Affected

- *vigabatrin*
- VIGADRONE
- VIGPODER

PA Criteria	Criteria Details
Exclusion Criteria	None
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
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

VIJOICE

Products Affected

- VIJOICE

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of severe manifestations of PIK3CA-Related Overgrowth Spectrum (PROS) in patients who require systemic therapy
Age Restrictions	2 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

VITRAKVI

Products Affected

- VITRAKVI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of metastatic or surgically unresectable neurotrophic receptor tyrosine kinase (NTRK) gene fusion positive solid tumors without a known acquired resistance mutation and used in patients with unsatisfactory alternative treatments or who have progressed following treatment
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VIZIMPRO

Products Affected

- VIZIMPRO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis 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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VONJO

Products Affected

- VONJO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of intermediate or high-risk primary or secondary myelofibrosis in adults AND a platelet count less than 50 X 10(9)/L
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

VORICONAZOLE

Products Affected

- voriconazole intravenous
- voriconazole oral

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Concomitant use of carbamazepine, CYP3A4 substrates (e.g., terfenadine, astemizole, cisapride, pimozone, or quinidine), B.) Concomitant use with high-dose ritonavir (400mg every 12 hours), C.) Concomitant use with ergot alkaloids, D.) Concomitant use with long-acting barbiturates, E.) Concomitant use with rifabutin or rifampin, F.) Concomitant use with sirolimus, or G.) Concomitant use with efavirenz at standard doses of 400mg/day or higher
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 <i>Scedosporium apiospermum</i> or <i>Fusarium</i> species
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist
Coverage Duration	6 months
Other Criteria	IV formulation: B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VUMERITY

Products Affected

- VUMERITY

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsing forms of multiple sclerosis (e.g., clinically isolated syndrome, relapsing-remitting MS, active secondary progressive disease, or progressive-relapsing MS), or B.) Patient has experienced a first clinical episode and has MRI features consistent with multiple sclerosis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a neurologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

VYNDAMAX

Products Affected

- VYNDAMAX

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of wild type or hereditary transthyretin related familial amyloid cardiomyopathy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

WELIREG

Products Affected

- WELIREG

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of one of the following A.) Von Hippel-Lindau (VHL) disease and therapy is required for any of the following disease associated tumors that do not require immediate surgery 1.) Renal cell carcinoma (RCC), 2.) Central nervous system (CNS) hemangioblastoma, or 3.) Pancreatic neuroendocrine tumor (pNET), or B.) Advanced renal cell carcinoma following a programmed death receptor-1 or programmed death-ligand 1 inhibitor and a vascular endothelial growth factor tyrosine kinase inhibitor
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XALKORI

Products Affected

- XALKORI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK)-positive or ROS1-positive as detected by an FDA-approved test, B.) Relapsed or refractory systemic anaplastic large cell lymphoma that is anaplastic lymphoma kinase (ALK) positive as detected by an FDA-approved test, or C.) Unresectable, recurrent, or refractory inflammatory myofibroblastic tumors that are anaplastic lymphoma kinase (ALK)-positive
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XDEMVY

Products Affected

- XDEMVY

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Demodex blepharitis
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XERMELO

Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of carcinoid syndrome diarrhea and both of the following 1.) Diarrhea is inadequately controlled by a stable dose of somatostatin analog (SSA) therapy (e.g., octreotide, lanreotide) for at least 3 months, and 2.) Used in combination with SSA therapy
Age Restrictions	None
Prescriber Restrictions	Prescribed by or in consultation with an oncologist, endocrinologist, or gastroenterologist
Coverage Duration	Plan year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

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 and used for the prevention of skeletal related events, B.) Multiple myeloma and used for the prevention of skeletal related events, C.) Hypercalcemia of malignancy refractory to bisphosphonate therapy, or D.) Giant cell tumor of bone that is unresectable or where surgical resection is likely to result in severe morbidity
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XOLAIR

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Chronic idiopathic urticaria in patients who remain symptomatic despite H1 antihistamine therapy and patient will continue to receive concurrent H1 antihistamine therapy unless contraindicated or not tolerated, 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 and an additional controller medication (i.e. long acting beta2-agonist, leukotriene modifier, or sustained-release theophylline) and patient has trial and failure, contraindication, or intolerance to Dupixent or Nucala, C.) Nasal polyps in patients with inadequate response to nasal corticosteroids, requested drug will be used as adjunctive treatment, and patient has trial and failure, contraindication, or intolerance to Dupixent, or D.) Reduction of allergic reactions (type I), including anaphylaxis, that may occur with accidental exposure to 1 or more foods in with IgE-mediated food allergy and is being used in conjunction with food allergen avoidance
Age Restrictions	1 year of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	B vs D determination required per CMS guidance
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Part B Prerequisite	No

Formulary ID: 24459 version 18
Last Updated: 09/24/2024
Effective: 10/01/2024

XOSPATA

Products Affected

- XOSPATA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of relapsed or refractory acute myeloid leukemia (AML) with a FMS-like tyrosine kinase 3 (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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XPOVIO

Products Affected

- XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 50 MG
- XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 40 MG
- XPOVIO (40 MG TWICE WEEKLY) ORAL TABLET THERAPY PACK 40 MG
- XPOVIO (60 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 60 MG
- XPOVIO (60 MG TWICE WEEKLY)
- XPOVIO (80 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 40 MG
- XPOVIO (80 MG TWICE WEEKLY)

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Relapsed or refractory multiple myeloma being used in combination with dexamethasone in a 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, B.) Multiple myeloma being used in combination with bortezomib and dexamethasone in a patient who has received at least 1 prior therapy, C.) Relapsed or refractory diffuse large B-cell lymphoma not otherwise specified, or D.) Relapsed or refractory DLBCL arising from follicular lymphoma and patient has received at least 2 lines of systemic therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

PA Criteria	Criteria Details
Off-Label Uses	N/A
Part B Prerequisite	No

XTANDI

Products Affected

- XTANDI

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Castration-resistant prostate cancer (CRPC), B.) Metastatic, castration-sensitive prostate cancer (mCSPC). For treatment of CRPC and mCSPC, one of the following applies 1.) Used in combination with a gonadotropin-releasing hormone (GnRH) analog or 2) Patient has received bilateral orchiectomy, or C.) Nonmetastatic castration-sensitive prostate cancer with biochemical recurrence at high risk for metastasis
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

XURIDEN

Products Affected

- XURIDEN

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of hereditary orotic aciduria
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

YONSA

Products Affected

- YONSA

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of metastatic, castration-resistant prostate cancer (mCRPC) and used in combination with methylprednisolone. For treatment of mCRPC, one of the following applies: 1.) Used in combination with a gonadotropin-releasing hormone (GnRH) analog or 2) Patient has received bilateral orchiectomy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or urologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

ZARXIO

Products Affected

- ZARXIO

PA Criteria	Criteria Details
Exclusion Criteria	None
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	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZEJULA

Products Affected

- ZEJULA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of Advanced or recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer and used as maintenance therapy in a patient who is in a complete or partial response to platinum-based chemotherapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZELBORAF

Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	None
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 or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZIEXTENZO

Products Affected

- ZIEXTENZO

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of a non-myeloid malignancy and drug is being used as prophylaxis for chemotherapy-induced neutropenia
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZILBRYSQ

Products Affected

- ZILBRYSQ

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of generalized myasthenia gravis in adults who are anti-acetylcholine receptor (AChR) antibody positive
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZOKINVY

Products Affected

- ZOKINVY

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of one of the following A.) Hutchinson-Gilford Progeria Syndrome, or B.) Treatment of processing deficient progeroid laminopathies with either heterozygous LMNA mutation with progerin-like protein accumulation or homozygous or compound heterozygous ZMPSTE24 mutations
Age Restrictions	None
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZOLINZA

Products Affected

- ZOLINZA

PA Criteria	Criteria Details
Exclusion Criteria	None
Required Medical Information	Diagnosis of primary cutaneous T-cell lymphoma (CTCL) in patients who have progressive, persistent or recurrent disease on or following two systemic therapies (e.g., bexarotene, romidepsin, etc)
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZURZUVAE

Products Affected

- ZURZUVAE

PA Criteria	Criteria Details
Exclusion Criteria	Any of the following A.) Actively suicidal, B.) Currently pregnant, C.) History of bipolar disorder, schizophrenia, or schizoaffective disorder
Required Medical Information	Diagnosis of postpartum depression
Age Restrictions	18 years of age and older
Prescriber Restrictions	None
Coverage Duration	14 days
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZYDELIG

Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Exclusion Criteria	History of toxic epidermal necrosis with any drug
Required Medical Information	Diagnosis of Chronic lymphocytic leukemia, used in combination with rituximab and patient has relapsed on at least one prior therapy
Age Restrictions	18 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist
Coverage Duration	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

ZYKADIA

Products Affected

- ZYKADIA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	None
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	Plan Year
Other Criteria	None
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

PART B VERSUS PART D

Products Affected

- ABELCET INTRAVENOUS SUSPENSION 5 MG/ML
- *acetylcysteine inhalation solution 10 %, 20 %*
- *acyclovir sodium intravenous solution 50 mg/ml*
- *albuterol sulfate inhalation nebulization solution (2.5 mg/3ml) 0.083%, 0.63 mg/3ml, 1.25 mg/3ml, 2.5 mg/0.5ml*
- *amphotericin b intravenous solution reconstituted 50 mg*
- *amphotericin b liposome intravenous suspension reconstituted 50 mg*
- *aprepitant oral capsule 125 mg, 40 mg, 80 & 125 mg, 80 mg*
- *azathioprine oral tablet 100 mg, 50 mg, 75 mg*
- *budesonide inhalation suspension 0.25 mg/2ml, 0.5 mg/2ml, 1 mg/2ml*
- *calcitonin (salmon) nasal solution 200 unit/act*
- *calcitriol oral capsule 0.25 mcg, 0.5 mcg*
- *calcitriol oral solution 1 mcg/ml*
- *caspofungin acetate intravenous solution reconstituted 50 mg, 70 mg*
- *cefoxitin sodium intravenous solution reconstituted 1 gm, 10 gm, 2 gm*
- *cinacalcet hcl oral tablet 30 mg, 60 mg, 90 mg*
- CLINIMIX E/DEXTROSE (2.75/5) INTRAVENOUS SOLUTION 2.75 %
- CLINIMIX E/DEXTROSE (4.25/10) INTRAVENOUS SOLUTION 4.25 %
- CLINIMIX E/DEXTROSE (4.25/5) INTRAVENOUS SOLUTION 4.25 %
- CLINIMIX E/DEXTROSE (5/15) INTRAVENOUS SOLUTION 5 %
- CLINIMIX E/DEXTROSE (5/20) INTRAVENOUS SOLUTION 5 %
- CLINIMIX/DEXTROSE (4.25/10) INTRAVENOUS SOLUTION 4.25 %
- CLINIMIX/DEXTROSE (4.25/5) INTRAVENOUS SOLUTION 4.25 %
- CLINIMIX/DEXTROSE (5/15) INTRAVENOUS SOLUTION 5 %
- CLINIMIX/DEXTROSE (5/20) INTRAVENOUS SOLUTION 5 %
- CLINISOL SF INTRAVENOUS SOLUTION 15 %
- *colistimethate sodium (cba) injection solution reconstituted 150 mg*
- *cromolyn sodium inhalation nebulization solution 20 mg/2ml*
- *cyclophosphamide oral capsule 25 mg, 50 mg*
- *cyclophosphamide oral tablet 25 mg, 50 mg*
- *cyclosporine modified oral capsule 100 mg, 25 mg, 50 mg*
- *cyclosporine modified oral solution 100 mg/ml*
- *cyclosporine oral capsule 100 mg, 25 mg*
- *dextrose intravenous solution 10 %, 5 %*
- *diphtheria-tetanus toxoids dt intramuscular suspension 25-5 lfu/0.5ml*
- ENGERIX-B INJECTION SUSPENSION 20 MCG/ML
- ENGERIX-B INJECTION SUSPENSION PREFILLED SYRINGE 10 MCG/0.5ML, 20 MCG/ML
- ENVARUS XR ORAL TABLET EXTENDED RELEASE 24 HOUR 0.75 MG, 1 MG, 4 MG
- *everolimus oral tablet 0.25 mg, 0.5 mg, 0.75 mg, 1 mg*

Formulary ID: 24459 version 16

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

- GAMMAGARD INJECTION SOLUTION 2.5 GM/25ML
- GAMMAGARD S/D LESS IGA INTRAVENOUS SOLUTION RECONSTITUTED 10 GM, 5 GM
- GAMUNEX-C INJECTION SOLUTION 1 GM/10ML
- GENGRAF ORAL CAPSULE 100 MG, 25 MG
- GENGRAF ORAL SOLUTION 100 MG/ML
- *granisetron hcl oral tablet 1 mg*
- HEPLISAV-B INTRAMUSCULAR SOLUTION PREFILLED SYRINGE 20 MCG/0.5ML
- IMOVAX RABIES INTRAMUSCULAR SUSPENSION RECONSTITUTED 2.5 UNIT/ML
- INTRALIPID INTRAVENOUS EMULSION 20 %, 30 %
- *ipratropium bromide inhalation solution 0.02 %*
- *ipratropium-albuterol inhalation solution 0.5-2.5 (3) mg/3ml*
- ISOLYTE-P IN D5W INTRAVENOUS SOLUTION
- ISOLYTE-S PH 7.4 INTRAVENOUS SOLUTION
- *methotrexate sodium (pf) injection solution 50 mg/2ml*
- *methotrexate sodium injection solution 50 mg/2ml*
- *methotrexate sodium oral tablet 2.5 mg*
- *methylprednisolone oral tablet 16 mg, 32 mg, 4 mg, 8 mg*
- *multiple electro type 1 ph 5.5 intravenous solution*
- *mycophenolate mofetil oral capsule 250 mg*
- *mycophenolate mofetil oral suspension reconstituted 200 mg/ml*
- *mycophenolate mofetil oral tablet 500 mg*
- *mycophenolate sodium oral tablet delayed release 180 mg, 360 mg*
- NUTRILIPID INTRAVENOUS EMULSION 20 %
- *ondansetron hcl oral solution 4 mg/5ml*
- *ondansetron hcl oral tablet 4 mg, 8 mg*
- *ondansetron oral tablet dispersible 4 mg, 8 mg*
- *paricalcitol oral capsule 1 mcg, 2 mcg, 4 mcg*
- *pentamidine isethionate inhalation solution reconstituted 300 mg*
- *pentamidine isethionate injection solution reconstituted 300 mg*
- PLASMA-LYTE A INTRAVENOUS SOLUTION
- PLENAMINE INTRAVENOUS SOLUTION 15 %
- *prednisolone oral solution 15 mg/5ml*
- *prednisolone sodium phosphate oral solution 10 mg/5ml, 20 mg/5ml, 25 mg/5ml, 6.7 (5 base) mg/5ml*
- *prednisolone sodium phosphate oral tablet dispersible 10 mg, 15 mg, 30 mg*
- PREDNISON INTENSOL ORAL CONCENTRATE 5 MG/ML
- *prednisone oral solution 5 mg/5ml*
- *prednisone oral tablet 1 mg, 10 mg, 2.5 mg, 20 mg, 5 mg, 50 mg*
- PREHEVBRIO INTRAMUSCULAR SUSPENSION 10 MCG/ML
- PREMASOL INTRAVENOUS SOLUTION 10 %
- PROGRAF ORAL PACKET 0.2 MG, 1 MG
- PROSOL INTRAVENOUS SOLUTION 20 %
- PULMOZYME INHALATION SOLUTION 2.5 MG/2.5ML
- RABAVERT INTRAMUSCULAR SUSPENSION RECONSTITUTED

Formulary ID: 24459 version 16

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

- RECOMBIVAX HB INJECTION SUSPENSION 10 MCG/ML, 40 MCG/ML, 5 MCG/0.5ML
- RECOMBIVAX HB INJECTION SUSPENSION PREFILLED SYRINGE 10 MCG/ML, 5 MCG/0.5ML
- *sirolimus oral solution 1 mg/ml*
- *sirolimus oral tablet 0.5 mg, 1 mg, 2 mg*
- *tacrolimus oral capsule 0.5 mg, 1 mg, 5 mg*
- TDVAX INTRAMUSCULAR SUSPENSION 2-2 LF/0.5ML
- TENIVAC INTRAMUSCULAR INJECTABLE 5-2 LFU, 5-2 LFU (INJECTION)
- *tigecycline intravenous solution reconstituted 50 mg*
- *tobramycin inhalation nebulization solution 300 mg/5ml*
- TPN ELECTROLYTES INTRAVENOUS CONCENTRATE
- TRAVASOL INTRAVENOUS SOLUTION 10 %
- TREXALL ORAL TABLET 10 MG, 15 MG, 5 MG, 7.5 MG
- TROPHAMINE INTRAVENOUS SOLUTION 10 %
- XATMEP ORAL SOLUTION 2.5 MG/ML

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.

Formulary ID: 24459 version 16

Last Updated: 09/24/2024

Effective: 10/01/2024

Index

A

ABELCET INTRAVENOUS
SUSPENSION 5 MG/ML..... 270
abiraterone acetate 1
acetylcysteine inhalation solution 10 %, 20
% 270
acitretin 2
ACTIMMUNE..... 3
acyclovir sodium intravenous solution 50
mg/ml 270
adapalene external cream..... 223
adapalene external gel 0.3 % 223
ADEMPAS 4
AKEEGA 5
albuterol sulfate inhalation nebulization
solution (2.5 mg/3ml) 0.083%, 0.63
mg/3ml, 1.25 mg/3ml, 2.5 mg/0.5ml .. 270
ALECENSA..... 6
ALUNBRIG..... 8
ALVAIZ..... 9
ambrisentan..... 10
amphotericin b intravenous solution
reconstituted 50 mg..... 270
amphotericin b liposome intravenous
suspension reconstituted 50 mg 270
aprepitant oral capsule 125 mg, 40 mg, 80 &
125 mg, 80 mg 270
ARCALYST 11
ARIKAYCE..... 12
armodafinil..... 36
AUGTYRO..... 13
AURYXIA 14
AUSTEDO..... 15
AUSTEDO XR ORAL TABLET
EXTENDED RELEASE 24 HOUR 12
MG, 24 MG, 30 MG, 36 MG, 42 MG, 48
MG, 6 MG..... 15
AUSTEDO XR PATIENT TITRATION
ORAL TABLET EXTENDED

RELEASE THERAPY PACK 6 & 12 &
24 MG 15
AVONEX PEN INTRAMUSCULAR
AUTO-INJECTOR KIT..... 141
AVONEX PREFILLED
INTRAMUSCULAR PREFILLED
SYRINGE KIT..... 141
AYVAKIT 16
azathioprine oral tablet 100 mg, 50 mg, 75
mg 270

B

BALVERSA 17
BENLYSTA SUBCUTANEOUS..... 18
BESREMI 19
BETASERON SUBCUTANEOUS KIT 141
bexarotene 20, 21
bosentan 22
BOSULIF..... 23
BRAFTOVI ORAL CAPSULE 75 MG ... 24
BRONCHITOL..... 25
BRUKINSA 26
budesonide inhalation suspension 0.25
mg/2ml, 0.5 mg/2ml, 1 mg/2ml 270
BYLVAY 27
BYLVAY (PELLETS)..... 27

C

CABLIVI 28
CABOMETYX 29
calcitonin (salmon) nasal solution 200
unit/act..... 270
calcitriol oral capsule 0.25 mcg, 0.5 mcg 270
calcitriol oral solution 1 mcg/ml 270
CALQUENCE 30
CAMZYOS..... 31
CAPRELSA 32
carglumic acid oral tablet soluble 33
caspofungin acetate intravenous solution
reconstituted 50 mg, 70 mg..... 270
CAYSTON..... 34

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

cefoxitin sodium intravenous solution reconstituted 1 gm, 10 gm, 2 gm.....	270
cinacalcet hcl oral tablet 30 mg, 60 mg, 90 mg	270
CLINIMIX E/DEXTROSE (2.75/5) INTRAVENOUS SOLUTION 2.75 %	270
CLINIMIX E/DEXTROSE (4.25/10) INTRAVENOUS SOLUTION 4.25 %	270
CLINIMIX E/DEXTROSE (4.25/5) INTRAVENOUS SOLUTION 4.25 %	270
CLINIMIX E/DEXTROSE (5/15) INTRAVENOUS SOLUTION 5 %	270
CLINIMIX E/DEXTROSE (5/20) INTRAVENOUS SOLUTION 5 %	270
CLINIMIX/DEXTROSE (4.25/10) INTRAVENOUS SOLUTION 4.25 %	270
CLINIMIX/DEXTROSE (4.25/5) INTRAVENOUS SOLUTION 4.25 %	270
CLINIMIX/DEXTROSE (5/15) INTRAVENOUS SOLUTION 5 %	270
CLINIMIX/DEXTROSE (5/20) INTRAVENOUS SOLUTION 5 %	270
CLINISOL SF INTRAVENOUS SOLUTION 15 %	270
colistimethate sodium (cba) injection solution reconstituted 150 mg.....	270
COMETRIQ (100 MG DAILY DOSE) ORAL KIT 80 & 20 MG	37
COMETRIQ (140 MG DAILY DOSE) ORAL KIT 3 X 20 MG & 80 MG	37
COMETRIQ (60 MG DAILY DOSE).....	37
COPIKTRA.....	38
COSENTYX (300 MG DOSE).....	39, 40
COSENTYX SENSOREADY (300 MG) 39, 40	
COSENTYX SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 75 MG/0.5ML	39, 40
COSENTYX UNOREADY	39, 40
COTELLIC	41
cromolyn sodium inhalation nebulization solution 20 mg/2ml	270

cyclophosphamide oral capsule 25 mg, 50 mg	270
cyclophosphamide oral tablet 25 mg, 50 mg	270
cyclosporine modified oral capsule 100 mg, 25 mg, 50 mg	270
cyclosporine modified oral solution 100 mg/ml	270
cyclosporine oral capsule 100 mg, 25 mg	270
CYSTADROPS.....	43
CYSTAGON.....	42
CYSTARAN.....	43
D	
dalfampridine er	44
DAURISMO	45
DAYBUE.....	46
deferasirox oral tablet	47
deferasirox oral tablet soluble.....	47
deferiprone	48
dextrose intravenous solution 10 %, 5 %	270
DIACOMIT.....	49
diclofenac sodium external gel 3 %	50
DIFICID.....	51
dimethyl fumarate oral	52
dimethyl fumarate starter pack oral capsule delayed release therapy pack	52
diphtheria-tetanus toxoids dt intramuscular suspension 25-5 lfu/0.5ml	270
DOJOLVI.....	53
dronabinol	54
droxidopa	55
DUPIXENT.....	56
E	
ELIGARD	123
EMGALITY.....	57
ENBREL MINI.....	58
ENBREL SUBCUTANEOUS SOLUTION 25 MG/0.5ML	58
ENBREL SUBCUTANEOUS SOLUTION PREFILLED SYRINGE	58
ENBREL SURECLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR	58

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

ENGERIX-B INJECTION SUSPENSION 20 MCG/ML	270	GATTEX.....	77
ENGERIX-B INJECTION SUSPENSION PREFILLED SYRINGE 10 MCG/0.5ML, 20 MCG/ML	270	GAVRETO	78
ENSPRYNG	59	gefitinib.....	79
ENVARBUS XR ORAL TABLET EXTENDED RELEASE 24 HOUR 0.75 MG, 1 MG, 4 MG	270	GENGRAF ORAL CAPSULE 100 MG, 25 MG	271
EPIDIOLEX.....	60	GENGRAF ORAL SOLUTION 100 MG/ML.....	271
ERIVEDGE.....	62	GILOTRIF	80
ERLEADA	63	glatiramer acetate	81
erlotinib hcl	64	GLEOSTINE ORAL CAPSULE 10 MG, 100 MG, 40 MG.....	82
everolimus oral tablet 0.25 mg, 0.5 mg, 0.75 mg, 1 mg	270	granisetron hcl oral tablet 1 mg	271
everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg	65	guanfacine hcl er	86
everolimus oral tablet soluble	66	H	
EVRYSDI	67	HEPLISAV-B INTRAMUSCULAR SOLUTION PREFILLED SYRINGE 20 MCG/0.5ML	271
F		HUMIRA (2 PEN)	88, 89
fentanyl	69	HUMIRA (2 SYRINGE) SUBCUTANEOUS PREFILLED SYRINGE KIT 10 MG/0.1ML, 20 MG/0.2ML, 40 MG/0.4ML, 40 MG/0.8ML	88, 89
fentanyl citrate buccal lozenge on a handle	68	HUMIRA-CD/UC/HS STARTER SUBCUTANEOUS PEN-INJECTOR KIT 80 MG/0.8ML	88, 89
FERRIPROX ORAL SOLUTION.....	48	HUMIRA-PED>/=40KG UC STARTER	88, 89
FILSPARI	70	HUMIRA-PSORIASIS/UVEIT STARTER	88, 89
fingolimod hcl.....	71	HYFTOR.....	90
FINTEPLA.....	72	I	
FIRMAGON (240 MG DOSE).....	73	IBRANCE.....	91
FIRMAGON SUBCUTANEOUS SOLUTION RECONSTITUTED 80 MG	73	icatibant acetate subcutaneous solution prefilled syringe	92
FOTIVDA.....	74	ICLUSIG.....	93
FRUZAQLA	75	IDHIFA	94
G		imatinib mesylate	95
GALAFOLD	76	IMBRUVICA ORAL CAPSULE.....	96
GAMMAGARD INJECTION SOLUTION 2.5 GM/25ML.....	270	IMBRUVICA ORAL SUSPENSION.....	96
GAMMAGARD S/D LESS IGA INTRAVENOUS SOLUTION RECONSTITUTED 10 GM, 5 GM....	271	IMBRUVICA ORAL TABLET 140 MG, 280 MG, 420 MG.....	96
GAMUNEX-C INJECTION SOLUTION 1 GM/10ML.....	271		

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

IMOVAX RABIES INTRAMUSCULAR SUSPENSION RECONSTITUTED 2.5 UNIT/ML	271
INBRIJA	97
INCRELEX.....	98
INLYTA.....	99
INQOVI	100
INREBIC.....	101
INTRALIPID INTRAVENOUS EMULSION 20 %, 30 %	271
INTRAROSA.....	102
ipratropium bromide inhalation solution 0.02 %	271
ipratropium-albuterol inhalation solution 0.5-2.5 (3) mg/3ml	271
ISOLYTE-P IN D5W INTRAVENOUS SOLUTION.....	271
ISOLYTE-S PH 7.4 INTRAVENOUS SOLUTION.....	271
ISTURISA ORAL TABLET 1 MG, 5 MG	103
itraconazole oral.....	104, 105
ivabradine hcl.....	106
ivermectin oral	107
IWILFIN	108
J	
JAKAFI.....	109
JAYPIRCA	110
JOENJA	111
K	
KALYDECO.....	112
KESIMPTA.....	113
KINERET SUBCUTANEOUS SOLUTION PREFILLED SYRINGE	114
KISQALI (200 MG DOSE)	115
KISQALI (400 MG DOSE)	115
KISQALI (600 MG DOSE)	115
KISQALI FEMARA (200 MG DOSE) ..	116
KISQALI FEMARA (400 MG DOSE) ..	116
KISQALI FEMARA (600 MG DOSE) ..	116
KOSELUGO	117
KRAZATI.....	118

L	
lapatinib ditosylate	119
lenalidomide.....	120
LENVIMA (10 MG DAILY DOSE)	121
LENVIMA (12 MG DAILY DOSE)	121
LENVIMA (14 MG DAILY DOSE)	121
LENVIMA (18 MG DAILY DOSE)	121
LENVIMA (20 MG DAILY DOSE)	121
LENVIMA (24 MG DAILY DOSE)	121
LENVIMA (4 MG DAILY DOSE)	121
LENVIMA (8 MG DAILY DOSE)	121
LEUKINE INJECTION SOLUTION RECONSTITUTED.....	122
leuprolide acetate (3 month)	123
leuprolide acetate injection	123
l-glutamine oral packet	124
lidocaine external patch 5 %	125
linezolid intravenous solution 600 mg/300ml	126
linezolid oral	126
LONSURF	127
LORBRENA.....	128
LUMAKRAS	129
LUPKYNIS.....	130
LUPRON DEPOT (1-MONTH)	123
LUPRON DEPOT (3-MONTH)	123
LUPRON DEPOT (4-MONTH)	123
LUPRON DEPOT (6-MONTH)	123
LUPRON DEPOT-PED (1-MONTH) INTRAMUSCULAR KIT 7.5 MG	123
LUPRON DEPOT-PED (3-MONTH) INTRAMUSCULAR KIT 11.25 MG.	123
LUPRON DEPOT-PED (6-MONTH)	123
LYNPARZA ORAL TABLET	131, 132
LYTGOBI (12 MG DAILY DOSE)	133
LYTGOBI (16 MG DAILY DOSE)	133
LYTGOBI (20 MG DAILY DOSE)	133
M	
MATULANE	134
MAVYRET.....	87
MAYZENT	135
MAYZENT STARTER PACK.....	135

Formulary ID: 24459 version 18
Last Updated: 09/24/2024
Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

MEKINIST	136	ODOMZO	151
MEKTOVI	137	OFEV	152
methotrexate sodium (pf) injection solution		OGSIVEO	153
50 mg/2ml	271	OJEMDA	154
methotrexate sodium injection solution 50		OJJAARA	155
mg/2ml	271	OMNITROPE SUBCUTANEOUS	
methotrexate sodium oral tablet 2.5 mg..	271	SOLUTION CARTRIDGE.....	84, 85
methoxsalen rapid	138	OMNITROPE SUBCUTANEOUS	
methylprednisolone oral tablet 16 mg, 32		SOLUTION RECONSTITUTED... ..	84, 85
mg, 4 mg, 8 mg	271	ondansetron hcl oral solution 4 mg/5ml..	271
mifepristone oral tablet 300 mg	139	ondansetron hcl oral tablet 4 mg, 8 mg...	271
miglustat.....	140	ondansetron oral tablet dispersible 4 mg, 8	
modafinil oral.....	36	mg	271
MOUNJARO	83	ONUREG	156
multiple electro type 1 ph 5.5 intravenous		OPSUMIT	157
solution.....	271	ORGOVYX.....	158
mycophenolate mofetil oral capsule 250 mg		ORKAMBI.....	159
.....	271	ORSERDU	160
mycophenolate mofetil oral suspension		OSPHENA	161
reconstituted 200 mg/ml	271	OTEZLA ORAL TABLET 30 MG	162
mycophenolate mofetil oral tablet 500 mg		OTEZLA ORAL TABLET THERAPY	
.....	271	PACK 10 & 20 & 30 MG	162
mycophenolate sodium oral tablet delayed		OZEMPIC (0.25 OR 0.5 MG/DOSE)	
release 180 mg, 360 mg	271	SUBCUTANEOUS SOLUTION PEN-	
N		INJECTOR 2 MG/3ML	83
NERLYNX	142	OZEMPIC (1 MG/DOSE)	
NICOTROL	143	SUBCUTANEOUS SOLUTION PEN-	
NINLARO.....	144	INJECTOR 4 MG/3ML	83
nitisinone oral capsule 10 mg, 2 mg, 5 mg		OZEMPIC (2 MG/DOSE)	83
.....	145	P	
NOXAFIL ORAL PACKET.....	171	PANRETIN.....	163
NUBEQA.....	146	paricalcitol oral capsule 1 mcg, 2 mcg, 4	
NUCALA.....	147	mcg.....	271
NUEDEXTA.....	148	pazopanib hcl	164
NUPLAZID ORAL CAPSULE.....	149	PEGASYS SUBCUTANEOUS SOLUTION	
NUPLAZID ORAL TABLET 10 MG....	149	180 MCG/ML	165
NUTRILIPID INTRAVENOUS		PEGASYS SUBCUTANEOUS SOLUTION	
EMULSION 20 %.....	271	PREFILLED SYRINGE	165
O		PEMAZYRE.....	166
octreotide acetate injection solution 100		pentamidine isethionate inhalation solution	
mcg/ml, 1000 mcg/ml, 200 mcg/ml, 50		reconstituted 300 mg.....	271
mcg/ml, 500 mcg/ml	150		

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

pentamidine isethionate injection solution reconstituted 300 mg.....	271
PIQRAY (200 MG DAILY DOSE).....	168
PIQRAY (250 MG DAILY DOSE).....	168
PIQRAY (300 MG DAILY DOSE).....	168
pirfenidone	169
PLASMA-LYTE A INTRAVENOUS SOLUTION.....	271
PLENAMINE INTRAVENOUS SOLUTION 15 %	271
POMALYST	170
posaconazole oral.....	171, 172
prednisolone oral solution 15 mg/5ml	271
prednisolone sodium phosphate oral solution 10 mg/5ml, 20 mg/5ml, 25 mg/5ml, 6.7 (5 base) mg/5ml.....	271
prednisolone sodium phosphate oral tablet dispersible 10 mg, 15 mg, 30 mg.....	271
PREDNISONE INTENSOL ORAL CONCENTRATE 5 MG/ML.....	271
prednisone oral solution 5 mg/5ml	271
prednisone oral tablet 1 mg, 10 mg, 2.5 mg, 20 mg, 5 mg, 50 mg	271
PREHEVBRIO INTRAMUSCULAR SUSPENSION 10 MCG/ML	271
PREMASOL INTRAVENOUS SOLUTION 10 %	271
PREVYMIS ORAL	173
PROGRAF ORAL PACKET 0.2 MG, 1 MG	271
PROLASTIN-C INTRAVENOUS SOLUTION.....	7
PROMACTA	174
PROSOL INTRAVENOUS SOLUTION 20 %	271
PULMOZYME INHALATION SOLUTION 2.5 MG/2.5ML.....	271
Q	
QINLOCK.....	175
quinine sulfate oral.....	176

R	
RABAVERT INTRAMUSCULAR SUSPENSION RECONSTITUTED...	271
RAVICTI	177
RECOMBIVAX HB INJECTION SUSPENSION 10 MCG/ML, 40 MCG/ML, 5 MCG/0.5ML	271
RECOMBIVAX HB INJECTION SUSPENSION PREFILLED SYRINGE 10 MCG/ML, 5 MCG/0.5ML	271
REGSPANEX.....	178
REPATHA	179
REPATHA PUSHTRONEX SYSTEM..	179
REPATHA SURECLICK.....	179
RETACRIT INJECTION SOLUTION 10000 UNIT/ML, 10000 UNIT/ML(1ML), 2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML, 4000 UNIT/ML, 40000 UNIT/ML	61
RETEVMO ORAL CAPSULE.....	180
REZLIDHIA	181
REZUROCK.....	182
riluzole	183
RINVOQ.....	184
RINVOQ LQ.....	185
RIVFLOZA.....	186
ROZLYTREK.....	187
RUBRACA	188
RYBELSUS	83
RYDAPT.....	189
S	
sapropterin dihydrochloride oral packet .	190
sapropterin dihydrochloride oral tablet...	190
SCEMBLIX	191
SIGNIFOR	192
sildenafil citrate oral tablet 20 mg	193
sirolimus oral solution 1 mg/ml	271
sirolimus oral tablet 0.5 mg, 1 mg, 2 mg	272
SKYRIZI PEN	194
SKYRIZI SUBCUTANEOUS.....	194
sodium oxybate	195

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

sodium phenylbutyrate oral powder 3 gm/tsp	167	TEPMETKO	216
sodium phenylbutyrate oral tablet.....	167	teriparatide (recombinant) subcutaneous solution pen-injector 620 mcg/2.48ml	217
sofosbuvir-velpatasvir.....	87	testosterone transdermal gel 10 mg/act (2%), 12.5 mg/act (1%), 20.25 mg/1.25gm (1.62%), 20.25 mg/act (1.62%), 25 mg/2.5gm (1%), 40.5 mg/2.5gm (1.62%), 50 mg/5gm (1%)	224
SOHONOS.....	196	tetrabenazine	218
SOMAVERT.....	197	THALOMID	219
sorafenib tosylate	198	TIBSOVO	220
SPRYCEL	199	tigecycline intravenous solution reconstituted 50 mg.....	272
STELARA SUBCUTANEOUS SOLUTION 45 MG/0.5ML	200	TOBI PODHALER.....	221
STELARA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE	200	tobramycin inhalation nebulization solution 300 mg/5ml	272
STIVARGA	201	tolvaptan.....	222
sunitinib malate.....	202	toremifene citrate	225
SYMDEKO.....	203	TORPENZ.....	226
SYMLINPEN 120 SUBCUTANEOUS SOLUTION PEN-INJECTOR.....	204	TPN ELECTROLYTES INTRAVENOUS CONCENTRATE	272
SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR	204	TRAVASOL INTRAVENOUS SOLUTION 10 %	272
SYNAREL	205	TRELSTAR MIXJECT	227
T		tretinoin external	223
TABRECTA	206	TREXALL ORAL TABLET 10 MG, 15 MG, 5 MG, 7.5 MG	272
tacrolimus oral capsule 0.5 mg, 1 mg, 5 mg	272	trientine hcl	228
TAFINLAR.....	207	TRIKAFTA.....	229
TAGRISO	208	TROPHAMINE INTRAVENOUS SOLUTION 10 %	272
TAKHZYRO.....	209	TRULICITY	83
TALZENNA	210	TRUQAP.....	230
TASIGNA	211	TUKYSA	231
TAVNEOS	212	TURALIO ORAL CAPSULE 125 MG..	232
tazarotene external cream	213	TYMLOS	233
tazarotene external gel	213	U	
TAZORAC EXTERNAL CREAM 0.05 %	213	UBRELVY	234
TAZVERIK.....	214	V	
TDVAX INTRAMUSCULAR SUSPENSION 2-2 LF/0.5ML	272	VALCHLOR.....	235
TEGLUTIK.....	183	VANFLYTA	236
TEGSEDI.....	215	varenicline tartrate (starter).....	35
TENIVAC INTRAMUSCULAR INJECTABLE 5-2 LFU, 5-2 LFU (INJECTION)	272		

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024

My Choice Wisconsin

Prior Authorization 2024

Last Updated 9/24/2024

varenicline tartrate oral tablet	35	XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 40 MG. 255, 256
VENCLEXTA.....	237	XPOVIO (40 MG TWICE WEEKLY) ORAL TABLET THERAPY PACK 40 MG
VENCLEXTA STARTING PACK	237	255, 256
VERQUVO	238	XPOVIO (60 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 60 MG. 255, 256
VERZENIO.....	239	XPOVIO (60 MG TWICE WEEKLY).. 255, 256
VICTOZA SUBCUTANEOUS SOLUTION PEN-INJECTOR.....	83	XPOVIO (80 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 40 MG. 255, 256
vigabatrin	240	XPOVIO (80 MG TWICE WEEKLY).. 255, 256
VIGADRONE.....	240	XTANDI
VIGPODER	240	257
VIJOICE	241	XURIDEN.....
VITRAKVI.....	242	258
VIZIMPRO	243	Y
VONJO	244	YARGESA.....
voriconazole intravenous	245	140
voriconazole oral.....	245	YONSA.....
VOSEVI.....	87	259
VUMERITY	246	Z
VYNDAMAX.....	247	ZARXIO
W		260
WELIREG.....	248	ZEJULA ORAL TABLET.....
X		261
XALKORI.....	249	ZELBORAF.....
XATMEP ORAL SOLUTION 2.5 MG/ML	272	262
XDEMVY	250	ZIEXTENZO
XERMELO	251	263
XGEVA.....	252	ZILBRYSQ.....
XOLAIR	253	264
XOSPATA	254	ZOKINVY
XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 50 MG	255, 256	265
		ZOLINZA
		266
		ZURZUVAE.....
		267
		ZYDELIG
		268
		ZYKADIA ORAL TABLET
		269

Formulary ID: 24459 version 18

Last Updated: 09/24/2024

Effective: 10/01/2024