2026 1 Tier Standard - SCHA

2026 Prior Authorization Criteria

CURRENT AS OF 11/01/2026

ACITRETIN

Products Affected

- *acitretin*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a dermatologist or an oncologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For prophylaxis of skin cancer in patients with previously treated skin cancers who have undergone an organ transplantation the request will be approved. For psoriasis: the patient has documented trial of, contraindication to, or medical reason for not using at least 2 of the treatment options listed: topical steroids, tazarotene, methotrexate, and cyclosporine.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ACL INHIBITORS

Products Affected

- NEXLETOL
- NEXLIZET

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	18 years or older
Prescriber Restrictions	Prescriber must be a cardiologist or specialist in the treatment of lipid disorders
Coverage Duration	New starts will be authorized for 4 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts ALL of the following must be provided: 1) Documentation of baseline low density lipoprotein cholesterol (LDL-C), 2) Member has tried and failed a high-intensity statin (i.e. atorvastatin 40-80 mg, rosuvastatin 20-40 mg) at maximum tolerated dose for 3 months via claim history or chart notes OR documentation has been provided that the member is not able to tolerate a statin. In addition to the initial criteria above if the new start is for the diagnosis of hypercholesterolemia, the following are required: 1) Member has a diagnosis of heterozygous familial hypercholesterolemia (FH) OR hypercholesterolemia, 2) Member has tried ezetimibe at a maximum tolerated dose and LDL-C is not at goal, or documentation has been provided that the patient is not able to tolerate ezetimibe. In addition to the initial criteria above if the new start is for cardiovascular risk reduction, the following are required: 1) Member has established cardiovascular disease (documented history of coronary artery disease, symptomatic peripheral artery disease, and/or cerebrovascular atherosclerotic disease, 2) Member does not have established cardiovascular disease but is considered high risk (one of the following): Diabetes Mellitus (Type 1 or Type 2) in females over 65 years of age or males over 60 years of age OR a Reynolds Risk score greater than 30% or a SCORE Risk score greater than 7.5% over 10 years OR a coronary artery calcium score greater than 400 Agaston units at any time in the past. For continuation of therapy or reauthorization requests for all indications:

PA Criteria	Criteria Details
	Documentation provided that the member has obtained clinical benefit from medication (e.g., LDL-C lowering from baseline)
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ACTEMRA

Products Affected

- ACTEMRA ACTPEN
- ACTEMRA SUBCUTANEOUS
- AVTOZMA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For pJIA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For sJIA, Giant Cell Arteritis and Systemic Sclerosis-Associated Interstitial Lung Disease: Approve
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ADALIMUMAB

Products Affected

- *adalimumab-fkjp (2 pen)*
- *adalimumab-fkjp (2 syringe) subcutaneous prefilled syringe kit 20 mg/0.4ml, 40 mg/0.8ml*
- SIMLANDI (1 PEN) SUBCUTANEOUS AUTO-INJECTOR KIT 40 MG/0.4ML, 80 MG/0.8ML
- SIMLANDI (1 SYRINGE)
- SIMLANDI (2 PEN)
- SIMLANDI (2 SYRINGE) SUBCUTANEOUS PREFILLED SYRINGE KIT 20 MG/0.2ML, 40 MG/0.4ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ankylosing spondylitis: Trial of, medical reason for not using, or contraindication to naproxen. For pJIA: Trial of, medical reason for not using, or contraindication to 1 of the following DMARDs: methotrexate or leflunomide. For RA: Trial of, medical reason for not using, or contraindication to 1 disease modifying antirheumatic drug (DMARD) (methotrexate, leflunomide, or sulfasalazine). For PsA, psoriasis, Hidradenitis Suppurativa, Crohn's Disease (CD), Ulcerative Colitis (UC) or Uveitis: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ADEMPAS

Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with Phosphodiesterase Inhibitors used for Pulmonary Hypertension or Other Soluble Guanylate Cyclase Stimulators.
Required Medical Information	Pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1)- Initial: [Note: documentation required] (1) PAH was confirmed by right heart catheterization AND (2) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg AND (3) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND (4) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. Continuation of therapy: patient has positive clinical response to treatment. Chronic thromboembolic pulmonary hypertension (CTEPH) (WHO Group 4) - Initial: [Note: documentation required] (1) Patient has persistent or recurrent CTEPH after pulmonary endarterectomy (PEA) OR (2) patient has inoperable CTEPH with the diagnosis confirmed by right heart catheterization AND by computed tomography (CT), magnetic resonance imaging (MRI), or pulmonary angiography. Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ALPHA-1 PROTEINASE INHIBITORS

Products Affected

- ARALAST NP INTRAVENOUS SOLUTION RECONSTITUTED 1000 MG, 500 MG
- GLASSIA
- PROLASTIN-C INTRAVENOUS SOLUTION
- ZEMAIRA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of hereditary alpha1-antitrypsin deficiency as evident by pretreatment serum AAT levels below 11 micromol/L and progressive FEV1 or FVC decline demonstrating symptomatic lung disease.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If the medication request is for Glassia or Aralast NP, the patient has a documented medical reason (such as trial, intolerance or contraindication) for not using Prolastin-C or Zemaira to treat their medical condition.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ALYFTREK

Products Affected

- ALYFTREK ORAL TABLET 10-50-125 MG, 4-20-50 MG

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with Trikafta, Kalydeco, Orkambi, or Symdeko. Patients with unknown CFTR gene mutations.
Required Medical Information	Documentation of CFTR gene that is responsive to vanzacaftor-tezacaftor-deutivacaftor treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist or a provider who specializes in treatment of CF.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Cystic Fibrosis (CF) - Initial: patient must have at least one F508del mutation in the CFTR gene or a mutation in the CFTR gene that is responsive to the requested medication. Continuation of therapy: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

AMBRISENTAN

Products Affected

- *ambrisentan*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1)- Initial: [Note: documentation required] (1) PAH was confirmed by right heart catheterization AND (2) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg AND (3) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND (4) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ANTINEOPLASTIC AGENTS

Products Affected

- *abiraterone acetate*
- ABIRTEGA
- AKEEGA
- ALECENSA
- ALUNBRIG
- AUGTYRO
- AVMAPKI FAKZYNJA CO-PACK
- AYVAKIT
- BALVERSA
- *bexarotene*
- BOSULIF ORAL CAPSULE
- BOSULIF ORAL TABLET 100 MG, 500 MG
- *bosutinib*
- BRAFTOVI ORAL CAPSULE 75 MG
- BRUKINSA
- CABOMETYX
- CALQUENCE ORAL TABLET
- CAPRELSA
- CAVHANZA
- 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)
- COPIKTRA
- COTELLIC
- DANZITEN
- *dasatinib*
- DAURISMO
- ENSACOVE
- ERIVEDGE
- ERLEADA
- *erlotinib hcl*
- EULEXIN
- *everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *everolimus oral tablet soluble*
- FOTIVDA
- FRUZAQLA
- GAVRETO
- *gefitinib*
- GILOTRIF
- GOMEKLI
- HERNEXEOS
- HYRNUO
- IBRANCE
- IBTROZI
- ICLUSIG
- IDHIFA
- *imatinib mesylate oral*
- IMKELDI
- INLURIYO
- INLYTA
- INQOVI
- INREBIC
- ITOVEBI
- IWILFIN
- JAYPIRCA
- JIDEYTRO
- KISQALI (200 MG DOSE)
- KISQALI (400 MG DOSE)
- KISQALI (600 MG DOSE)
- KOMZIFTI
- KOSELUGO
- KRAZATI
- *lapatinib ditosylate*
- LAZCLUZE
- *lenalidomide*
- 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)
- LEUKERAN
- LIFYORLI (125 MG DOSE) ORAL CAPSULE THERAPY PACK 1 X 25 MG & 1 X 100 MG
- LIFYORLI (150 MG DOSE) ORAL CAPSULE THERAPY PACK 2 X 25 MG & 1 X 100 MG

- LONSURF
- LORBRENA
- LUMAKRAS
- LYNPARZA ORAL TABLET
- LYTGOBI (12 MG DAILY DOSE)
- LYTGOBI (16 MG DAILY DOSE)
- LYTGOBI (20 MG DAILY DOSE)
- MEKINIST
- MEKTOVI
- *mercaptopurine oral suspension*
- MODEYSO
- NERLYNX
- *nilotinib d-tartrate*
- *nilotinib hcl*
- *nilutamide*
- NINLARO
- NUBEQA
- ODOMZO
- OGSIVEO ORAL TABLET 100 MG, 150 MG
- OJEMDA
- OJJAARA
- ONUREG
- ORGOVYX
- ORSERDU
- *pazopanib hcl*
- PEMAZYRE
- PIQRAY (200 MG DAILY DOSE)
- PIQRAY (250 MG DAILY DOSE)
- PIQRAY (300 MG DAILY DOSE)
- *pomalidomide*
- QINLOCK
- RASONQUE
- RETEVMO ORAL TABLET
- REVLIMID
- REVUFORJ
- REZLIDHIA
- ROMVIMZA
- ROZLYTREK
- RUBRACA
- RYDAPT
- SCEMBLIX
- SOLTAMOX
- *sorafenib tosylate*
- STIVARGA
- *sunitinib malate*
- TABLOID
- TABRECTA
- TAFINLAR
- TAGRISSO
- TALZENNA
- TAZVERIK
- TEPMETKO
- THALOMID ORAL CAPSULE 100 MG, 50 MG
- TIBSOVO
- *toremifene citrate*
- *tretinoin oral*
- TRUQAP ORAL TABLET 200 MG
- TRUQAP ORAL TABLET THERAPY PACK
- TUKYSA
- TURALIO ORAL CAPSULE 125 MG
- VANFLYTA
- VENCLEXTA
- VENCLEXTA STARTING PACK
- VEPPANU
- VERZENIO
- VITRAKVI
- VIZIMPRO
- VONJO
- VORANIGO
- WELIREG
- XALKORI
- XOSPATA
- XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 50 MG
- XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 10 MG, 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, 80 MG
- XPOVIO (80 MG TWICE WEEKLY)

- XTANDI
- YONSA
- YULITHIRA
- ZEJULA ORAL TABLET
- ZELBORAF
- ZENBEXUS
- ZOLINZA
- ZYDELIG
- ZYKADIA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an oncologist or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

APOMORPHINE

Products Affected

- *apomorphine hcl subcutaneous*

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with serotonin 5-HT3 receptor antagonists.
Required Medical Information	Reviewer will verify available patient claim history to confirm patient is not using 5-HT3 receptor antagonists.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If diagnosis is Parkinson's, the patient must have a documented trial of, contraindication to, or medical reason for not using two alternatives such as entacapone, tolcapone, rasagiline, selegiline, carbidopa/levodopa, bromocriptine, pramipexole or ropinirole.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

AQNEURSA

Products Affected

- AQNEURSA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts: 1) The member has a documented diagnosis of Niemann-Pick disease type C (NPC) AND 2) Documentation of genetic testing identifying disease-causing alleles in NPC1 or NPC2 AND 3) Documentation of disease-related neurological symptoms (e.g., developmental delay/regression, ataxia, cataplexy, seizures, motor-function decline, tremors, dysphagia) For reauthorization: Documentation that member has had positive response to therapy (e.g., stabilization in neurological status, decrease in functional Scale for Assessment and Rating of Ataxia [fSARA] score).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ARCALYST

Products Affected

- ARCALYST

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For deficiency of interleukein-1 receptor antagonist, documented trial of, contraindication to, or medical reason for not using Kineret. For continuation of therapy or reauthorization: Documentation has been provided that patient has clinically benefited from medication.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ARIKAYCE

Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Mycobacterium avium complex (MAC): (1) Documented diagnosis of MAC lung disease as verified by failure to acheive at least 2 negative sputum cultures following 6 consecutive months of a combination antibacterial drug regimen AND (2) Provider attestation that medication is being used as part of a combination antibacterial drug regimen.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist or an infectious disease specialist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ARISTADA

Products Affected

- ARISTADA INITIO 441 MG/1.6ML, 662 MG/2.4ML, 882 MG/3.2ML
- ARISTADA INTRAMUSCULAR PREFILLED SYRINGE 1064 MG/3.9ML,

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented history of receiving oral aripiprazole without any clinically significant side effects.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least two of the following: Abilify Maintena, Abilify Asimtufii, Invega Sustenna, Invega Trinza, Invega Hafyera, or Risperidone Microspheres ER.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

AUVELITY

Products Affected

- AUVELITY ORAL TABLET EXTENDED RELEASE 45-105 MG
- AUVELITY TITRATION PACK

PA Criteria	Criteria Details
Exclusion Criteria	Seizure disorder
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using to two generic antidepressants.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

AZTREONAM LYSINE

Products Affected

- CAYSTON

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist, infectious disease specialist, or an expert in the treatment of cystic fibrosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

BENLYSTA

Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a rheumatologist, nephrologist, or specialist in the treatment of autoimmune disorders.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts for systemic lupus erythematosus (SLE): concurrent use of two of the following or medical reason for not using glucocorticoids, azathioprine, methotrexate, mycophenolate, or hydroxychloroquine, chloroquine, and cyclophosphamide. For continuation of therapy or reauthorization for SLE: documentation of clinical response to therapy (i.e. fewer flares that required steroid treatment, lower average daily oral prednisone dose, improved daily function either as measured through a validated functional scale or through improved daily performance documented at clinic visits, etc.) For new starts for lupus nephritis (LN): concurrent use of or medical reason for not using background immunosuppressive therapy regimen. For continuation of therapy or reauthorization for LN: Documentation of improvement in renal function (i.e. reduction in UPCR).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

BESREMI

Products Affected

- BESREMI
- BESREMI PEN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a hematologist, oncologist, or specialist for submitted diagnosis.
Coverage Duration	The request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

BOSENTAN

Products Affected

- *bosentan oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of pulmonary arterial hypertension (PAH) WHO Group I classification and PAH Functional Class.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or cardiologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

BRINSUPRI

Products Affected

- BRINSUPRI

PA Criteria	Criteria Details
Exclusion Criteria	Patients with bronchiectasis due to cystic fibrosis.
Required Medical Information	N/A
Age Restrictions	12 years of age and older.
Prescriber Restrictions	Prescribed by or in consultation with a pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For non-cystic fibrosis bronchiectasis - Initial: (1) patient has documented history of bronchiectasis diagnosed by chest computed tomography scan AND (2) prescriber confirmation that bronchiectasis is not primarily driven by chronic obstructive pulmonary disease or asthma. Continuation of therapy: patient has positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CAMZYOS

Products Affected

- CAMZYOS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For all new starts, documentation of ALL of the following must be provided: 1) Diagnosis of symptomatic New York Heart Association (NYHA) class II or III obstructive hypertrophic cardiomyopathy (oHCM) AND 2) Patient has a left ventricular ejection fraction (LVEF) greater than or equal to 55% AND 3) Assessment of Valsalva left ventricular outflow tract (LVOT) gradient AND 4) Trial of, medical reason for not using or contraindication to BOTH of the following: Beta blockers (i.e. metoprolol, propranolol, atenolol) AND Non-dihydropyridine calcium channel blockers (i.e. verapamil, diltiazem) AND 5) Prescriber attests that patient is not using moderate to strong CYP2C19 or CYP3A4 inhibitors or inducers. For continuation of therapy or reauthorization, all of the following must be provided: 1) Documentation of clinical benefit as evidenced by an improvement from baseline in oHCM symptoms (i.e., improvement in fatigue, chest pain, shortness of breath, LVOT, peak oxygen consumption, etc.) OR improvement or no worsening of NYHA functional class AND 2) Member must also have a left ventricular ejection fraction (LVEF) greater than or equal to 50%.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PA Criteria	Criteria Details
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

CARDAMYST

Products Affected

- CARDAMYST

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documented history of acute symptomatic episodes of paroxysmal supraventricular tachycardia.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CARGLUMIC ACID

Products Affected

- *carglumic acid oral tablet soluble*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CASPOFUNGIN

Products Affected

- *caspofungin acetate*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Documentation of a consultation with an infectious disease specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CERDELGA

Products Affected

- CERDELGA

PA Criteria	Criteria Details
Exclusion Criteria	Patients with undetermined CYP2D6 metabolizer status.
Required Medical Information	Patient's CYP2D6 metabolizer status, as determined by an FDA approved test. For reauthorization, documentation has been provided that patient has obtained clinical benefit from medication (e.g. increased platelet count, improvement in anemia, PFTs, improvement in radiographic scans, improved quality of life).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a specialist in treatment of Gaucher's disease.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CGRP ANTAGONISTS

Products Affected

- AIMOVIG
- EMGALITY
- EMGALITY (300 MG DOSE)
- NURTEC
- QULIPTA
- UBRELVY
- ZAVZPRET

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For acute migraine new starts - for Ubrelvy and Nurtec requests, must have trial of, contraindication to or medical reason for not using a triptan. For Zavzpret requests, must have trial of, contraindication to or medical reason for not using Ubrelvy or Nurtec. For migraine prophylaxis new starts - at least 4 migraine days per month or one or more severe migraine attacks lasting for greater than 12 hours despite use of abortive therapy (e.g. triptans or NSAIDs). For Emgality requests for episodic cluster headache new starts - approve. For continuation of therapy or reauthorization - For acute migraine (Nurtec, Ubrelvy, Zavzpret), must show documentation of improvement in migraine symptoms (pain, photophobia, phonophobia). For migraine prevention (Nurtec, Emgality, Qulipta, Aimovig), must show documentation of improvement in migraine symptoms. For episodic cluster headache treatment, must show documentation of reduction in frequency of headaches.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

CHOLBAM

Products Affected

- CHOLBAM

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: Patient has documented diagnosis of either: 1) bile acid synthesis disorder due to a single enzyme defect or 2) peroxisomal disorders. For continuation of therapy or reauthorization: prescriber attests: 1) the patient has clinical improvement with therapy (i.e. liver function tests) AND 2) there is no evidence of biliary obstruction or cholestasis
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a hepatologist, gastroenterologist, or metabolic specialist
Coverage Duration	New starts will be authorized for 3 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CIBINQO

Products Affected

- CIBINQO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For atopic dermatitis: Trial of, contraindication to, or medical reason for not using Rinvoq or Dupixent.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

CIMZIA

Products Affected

- CIMZIA (1 SYRINGE)
- CIMZIA (2 SYRINGE)
- CIMZIA SUBCUTANEOUS KIT 2 X 200 MG
- CIMZIA-STARTER

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ankylosing spondylitis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: Enbrel, an adalimumab product, Cosentyx, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For Crohns Disease: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an adalimumab product or an ustekinumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For non-radiographic axial spondyloarthritis: approve. For psoriasis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: Tremfya, an ustekinumab product, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For PsA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: Cosentyx, an ustekinumab product, Tremfya, Xeljanz, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. Polyarticular juvenile idiopathic arthritis (pJIA) - Initial: trial of, medical reason for not using, or contraindication to 2 of the following:

PA Criteria	Criteria Details
	Enbrel, an adalimumab product or Xeljanz or or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

CORLANOR

Products Affected

- CORLANOR ORAL SOLUTION
- *ivabradine hcl*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	New starts for chronic heart failure must have all of the following: 1) LVEF of 35% or less 2) Sinus rhythm and have resting heart rate greater than or equal to 70 bpm. For pediatric patients with heart failure due to dilated cardiomyopathy: approve.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a cardiologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not receiving a beta blocker.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

CORTROPHIN

Products Affected

- CORTROPHIN
- CORTROPHIN GEL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	New starts for MS exacerbation, rheumatic disorders, collagen diseases, dermatologic diseases, serum sickness, edematous state (e.g. nephrotic syndrome without uremia), and respiratory diseases: trial of, contraindication to, or medical reason for not using oral corticosteroids. New starts for ophthalmic disease: trial of, contraindication to, or medical reason for not using oral or ophthalmic corticosteroids. Continuation of therapy or reauthorization for MS exacerbation: documentation of symptom improvement and current use of a multiple sclerosis disease modifying agent for maintenance therapy. Continuation of therapy or reauthorization for all other conditions: documented evidence of response to treatment and symptom improvement.
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	MS exacerbation: 1 month. Other conditions: new start for 3 months and reauth end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

COSENTYX

Products Affected

- COSENTYX (300 MG DOSE)
- COSENTYX INTRAVENOUS
- COSENTYX SENSOREADY (300 MG)
- COSENTYX SENSOREADY PEN
- COSENTYX SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 150 MG/ML, 75 MG/0.5ML
- COSENTYX UNOREADY

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	1) Documented diagnosis of ankylosing spondylitis, non-radiographic axial spondyloarthritis, plaque psoriasis, psoriatic arthritis, enthesitis-related arthritis, rheumatoid arthritis or hidradenitis suppurativa. 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CRESEMBA

Products Affected

- CRESEMBA ORAL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documented diagnosis of medically accepted indication.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist or oncologist
Coverage Duration	Request will be authorized for 3 months.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CYSTAGON

Products Affected

- CYSTAGON

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

CYSTARAN

Products Affected

- CYSTARAN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of diagnosis for cystinosis with corneal cystine crystal accumulation.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an ophthalmologist or metabolic disease specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

DALFAMPRIDINE ER

Products Affected

- *dalfampridine er*

PA Criteria	Criteria Details
Exclusion Criteria	History of seizure or moderate/severe renal impairment (CrCl less than or equal to 50 mL/min).
Required Medical Information	For new starts: 1) Attestation that creatinine clearance (CrCl) greater than 50 mL/min was confirmed prior to initiation of therapy, AND 2) Documentation has been provided that member is ambulatory (able to walk at least 25 feet) and has a documented walking impairment, AND 3) For multiple sclerosis, member is currently being treated with a disease modifying agent (e.g. immunomodulator, interferon, etc.) or has a medical reason why member is unable to use a disease modifying agent for their condition. For continuation of therapy or re-authorization requests: 1) Member must experience improvement in walking from baseline due to use of dalfampridine ER.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

DEFERASIROX

Products Affected

- *deferasirox*
- *deferasirox granules*

PA Criteria	Criteria Details
Exclusion Criteria	Creatinine clearance less than 40 mL/min or platelet counts less than 50,000/mm3.
Required Medical Information	For all indications: platelet count greater than or equal to 50,000/mm3 (within 30 days). For chronic iron overload due to transfusions: serum ferritin concentration greater than 1000 mcg/L (lab result within 30 days). For chronic iron overload in non-transfusion-dependent thalassemia syndromes: serum ferritin concentration greater than 300 mcg/L (lab result within 30 days).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For deferasirox granules oral packets, the member must have medical reason for not using deferasirox tablets or oral soluble tablets.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DEFERIPRONE

Products Affected

- *deferiprone*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: 1) serum ferritin level above 1,000 mcg/L and absolute neutrophil count (ANC) greater than 1.5x10^9/L within 30 days of request, and 2) Trial of, contraindication to, or medical reason for not using deferasirox tablets. For continuation of therapy or reauthorization, decrease in serum ferritin from baseline.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DEFLAZACORT

Products Affected

- KYMBEE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	1) Member has a documented diagnosis of Duchenne Muscular Dystrophy as evidenced by one of the following: a) documented mutation of dystrophin gene OR b) muscle biopsy indicating absence of the dystrophin protein AND 2) Patient has trial and failure with, contraindication to or medical reason for not taking prednisone.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DIACOMIT

Products Affected

- DIACOMIT

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For members 2 years and older: Trial of, contraindication to, or medical reason for not using one generic anticonvulsant for appropriate indications. For members under 2 years old: Approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DIHYDROERGOTAMINE NASAL

Products Affected

- *dihydroergotamine mesylate nasal*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: Member has a diagnosis of migraine headaches with or without aura. Prescriber attestation that it will be used for the acute treatment of migraine. For continuation of therapy or reauthorization: Documentation or provider attestation of positive clinical response (e.g., improvement in pain, photophobia, phonophobia).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Requests will be authorized for 12 weeks.
Other Criteria	Trial of, contraindication to, or medical reason (e.g. intolerance or hypersensitivity) for not using a triptan (e.g., rizatriptan, sumatriptan).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DOPTELET

Products Affected

- DOPTELET
- DOPTELET SPRINKLE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for chronic liver disease and chronic immune thrombocytopenia (chronic ITP): documented baseline platelet count of less than 50,000/mcL.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with hematologist, hepatologist or surgeon.
Coverage Duration	For thrombocytopenia with CLD getting procedure: 5 days. For all others: remainder of contract year
Other Criteria	For chronic ITP: trial of, contraindication to, or medical reason for not using a corticosteroid. For thrombocytopenia with chronic liver disease (CLD): approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DOXEPIN CREAM

Products Affected

- *doxepin hcl external*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 1 month.
Other Criteria	Trial of, contraindication to, or medical reason for not using a topical corticosteroid or topical calcineurin inhibitor.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DUPIXENT

Products Affected

- DUPIXENT SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- DUPIXENT SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 200 MG/1.14ML, 300 MG/2ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Chronic spontaneous urticaria (CSU) - Initial: patient has inadequate symptomatic relief despite trial of two weeks of a second-generation oral antihistamine (unless contraindicated). Continuation of therapy: patient has positive clinical response to treatment. Bullous pemphigoid - Initial: patient has had trial of, contraindication to, or medical reason for not using a topical corticosteroid, oral corticosteroid, or an immunosuppressive agent. Continuation of therapy: patient has positive clinical response to treatment. Allergic Fungal Rhinosinusitis: Initial (1) patient is 6 years of age and older with history of sino-nasal surgery AND (2) patient has a trial of, contraindication to, or medical reason for not using nasal corticosteroids. Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Initial therapy only: prescribed by or in consultation with specialist for submitted diagnosis.
Coverage Duration	PN: Initial 6 mo. Cont of therapy and all others initial: end of contract year.
Other Criteria	Atopic dermatitis (AD) in patients 6 months of age and older - Initial: (1) patient has diagnosis of moderate to severe AD, AND (2) has had trial of, contraindication to, or medical reason for not using either a topical corticosteroid or topical calcineurin inhibitor. Continuation of therapy: patient has positive clinical response to treatment. Asthma with eosinophilic phenotype - Initial: (1) patient has baseline blood eosinophil count greater than or equal to 150 cells per microliter, AND (2) asthma remains inadequately controlled despite current treatment with or medical reasons for not using BOTH (i) medium to high dose inhaled corticosteroid AND (ii) additional controller (i.e. leukotriene modifier, long acting beta-2-agoinist, long acting muscarinic antagonist, and sustained released theophylline). Asthma, oral corticosteroid dependent - Initial: asthma

PA Criteria	Criteria Details
	remains inadequately controlled despite current treatment with or medical reasons for not using BOTH (i) high dose inhaled corticosteroid AND (ii) additional controller (i.e. leukotriene modifier, long acting beta-2-agoinist, long acting muscarinic antagonist, and sustained released theophylline). Asthma with eosinophilic phenotype or oral corticosteroid dependent - Continuation of therapy: clinical improvement in asthma control (i.e. reduction in frequency and/or severity of exacerbations and symptoms OR reduction in daily maintenance oral corticosteroid dose). Chronic rhinosinusitis with nasal polyps (CRSwNP) - Initial: (1) Dupixent is used as add-on maintenance treatment, AND (2) patient has a trial of, contraindication to, or medical reason for not using nasal corticosteroids. Continuation of therapy: patient has positive clinical response to treatment. Prurigo nodularis (PN): attestation is provided confirming diagnosis. Eosinophilic esophagitis (EoE) - Initial: (1) diagnosis has been confirmed by esophageal biopsy AND (2) patient weighs at least 15 kilograms AND (3) patient experienced an inadequate treatment response, intolerance, or has a contraindication to a topical or oral corticosteroid (i.e. fluticasone propionate or budesonide). Continuation of therapy: patient has positive clinical response to treatment. Chronic Obstructive Pulmonary Disease (COPD) - Initial: (1) documented diagnosis of COPD with an eosinophilic phenotype, AND (2) Dupixent is used as an add-on maintenance treatment, AND (3) documentation that patient's COPD is inadequately controlled. Continuation of therapy: patient has positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EGRIFTA

Products Affected

- EGRIFTA SV
- EGRIFTA WR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of active antiretroviral therapy for at least 8 weeks.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

EMSAM

Products Affected

- EMSAM

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with SSRIs, SNRIs, clomipramine and imipramine, meperidine, tramadol, methadone, pentazocine, and propoxyphene, and the antitussive agent dextromethorphan or carbamazepine
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: Trial of, contraindication to, or medical reason for not using two generic antidepressants.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

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	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For RA: Trial of, medical reason for not using, or contraindication to 1 disease modifying antirheumatic drug (DMARD) (methotrexate, leflunomide, or sulfasalazine). For pJIA: Trial of, medical reason for not using, or contraindication to 1 of the following DMARDs: methotrexate or leflunomide. For PsA or psoriasis: approve. For ankylosing spondylitis: Trial of, medical reason for not using, or contraindication to naproxen. Continuation of therapy: patient has been receiving Enbrel for a minimum of 4 months and has positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ENDARI

Products Affected

- *l-glutamine oral packet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation that two or more painful sickle cell crises have occurred in the past 12 months.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a hematologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using hydroxyurea for at least three months.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ENTYVIO

Products Affected

- ENTYVIO PEN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ulcerative colitis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EPIDIOLEX

Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using one generic anticonvulsant for appropriate indications.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EPRONTIA

Products Affected

- *topiramate oral solution 25 mg/ml*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	The request will be authorized until the end of the contract year.
Other Criteria	Documented trial of, contraindication to, or medical reason for not using topiramate sprinkle capsule or topiramate oral tablet.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ERYTHROPOIETIN STIMULATING AGENTS

Products Affected

- ARANESP (ALBUMIN FREE) INJECTION SOLUTION 100 MCG/ML, 200 MCG/ML, 25 MCG/ML, 40 MCG/ML, 60 MCG/ML
- ARANESP (ALBUMIN FREE) INJECTION SOLUTION PREFILLED SYRINGE
- EPOGEN INJECTION SOLUTION 10000 UNIT/ML, 2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML, 4000 UNIT/ML
- PROCRIT
- 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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for all indications: Hgb within compendia range for treatment of the requested medical condition. For continuation of therapy or re-authorization: Hgb must not exceed 10 g/dL (anemia related to cancer), 11 g/dL (anemia of CKD), 12 g/dL (zidovudine-related anemia in members with HIV and ribavirin-induced anemia), 13 g/dL (elective, noncardiac, nonvascular surgery needing red blood cell allogeneic transfusion reduction).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 6 months.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ERZOFRI

Products Affected

- ERZOFRI INTRAMUSCULAR SUSPENSION PREFILLED SYRINGE 117 MG/0.75ML, 156 MG/ML, 234 MG/1.5ML, 351 MG/2.25ML, 39 MG/0.25ML, 78 MG/0.5ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented history of receiving oral paliperidone or oral risperidone without any clinically significant side effects.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For schizophrenia: Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least two of the following: Abilify Maintena, Abilify Asimtufii, Invega Sustenna, Invega Trinza, Invega Hafyera, or Risperidone Microsphere ER. For schizoaffective disorder: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EUCRISA

Products Affected

- EUCRISA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a dermatologist, immunologist or an allergist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For patients 2 years of age and older: Trial of, contraindication to, or medical reason for not using topical tacrolimus or pimecrolimus. For patients less than 2 years of age: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EVRYSDI

Products Affected

- EVRYSDI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts, all of the following must be included: 1) Documentation of genetic testing confirming diagnosis AND 2) Documentation of baseline motor function or motor milestone achievement [e.g. CHOP Infant Test of Neuromuscular Disorders (CHOP-INTEND) or Hammersmith Infant Neurological Examination (HINE) for Type 1 or Hammersmith Functional Motor Scale Expanded Scores (HFMSE), or Total Motor Function Measure 32 (MFM-32) for Type II and Type III, or 6 minute walk test in subjects able to walk]. For continuation of therapy or reauthorization, documentation of clinical response has been submitted (e.g. improvement in motor function/motor milestone achievement scores using CHOP-INTEND or HFMSE, MFM-32, 6 minute walk test or HINE improvement in more categories of motor milestones than worsening).
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

FABHALTA

Products Affected

- FABHALTA

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with another complement inhibitor for the treatment of PNH (i.e. Empaveli, Soliris, or Ultomiris).
Required Medical Information	PNH - Initial: patient has documented diagnosis of paroxysmal nocturnal hemoglobinuria (PNH) confirmed by (1) flow cytometry analysis confirming presence of PNH clones AND (2) patient has signs and symptoms of PNH (i.e. anemia, abdominal pain, dyspnea, kidney disease, pulmonary hypertension, hemolysis/hemoglobinuria, etc.). Continuation of therapy: patient has documented positive clinical response to treatment (i.e. decrease in LDH, increased or stabilization of hemoglobin levels, reduction in transfusions, increased reticulocyte count, etc.). Reduction of proteinuria in adults with immunoglobulin A (IgA) nephropathy - Initial: patient has documented diagnosis of IgA nephropathy AND IgA nephropathy at risk of rapid disease progression (i.e. clinical evidence of rapid disease progression generally a urine protein-to-creatinine ratio or UPCR greater or equal to 1.5g/g OR other clinically relevant tests). Continuation of therapy: patient has documented positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hematologist, nephrologist or oncologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

FASENRA

Products Affected

- FASENRA
- FASENRA PEN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	New starts for severe asthma with an eosinophilic phenotype: 1)Baseline blood eosinophil count greater than or equal to 150 cells per microliter AND 2) symptoms persist with at least 1 exacerbation in the last 12 months requiring additional treatment (e.g. oral systemic steroids) while on a high dose inhaled corticosteroid with an additional controller medication (ie. long-acting B2 agonist). Continuation of therapy or re-authorization for severe asthma with an eosinophilic phenotype: clinical benefit from use of the drug. New starts for eosinophilic granulomatosis with polyangiitis (EGPA)- Initial: patient has a documented history or the presence of an eosinophil count of more than 1000 cells per microliter or a blood eosinophil level of greater than 10 percent AND trial of, contraindication to, or medical reason for not using one of the following medications: cyclophosphamide or methotrexate. Continuation of therapy: patient has a beneficial response to treatment with the requested drug (i.e. a reduction in the frequency of relapses, decrease in the daily oral corticosteroid dose, or no active vasculitis).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

FILSPARI

Products Affected

- FILSPARI

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with renin-angiotensin-aldosterone system (RAAS) inhibitors, endothelin receptor antagonists, or aliskiren
Required Medical Information	For new starts for IgAN: Documentation is provided that the member has diagnosis of primary immunoglobulin A nephropathy (IgAN) at risk of rapid disease progression. Member has proteinuria. For new starts for focal segmental glomerulosclerosis (FSGS): Documentation is provided that the member has a diagnosis of FSGS confirmed by renal biopsy OR documented genetic mutation in a podocyte protein associated with FSGS AND (2) used to reduce proteinuria AND (3) patient is not currently in nephrotic syndrome (i.e. has proteinuria greater than 3.5 g/day in adults or UPCR greater than 2.0 g/g in pediatric patients, serum albumin less than 3.0 g/dL, and edema). For continuation of therapy or reauthorization: Documentation of positive clinical response (ie. decrease in urine protein-to-creatinine ratio (UPCR))
Age Restrictions	FSGS: 8 years of age and older
Prescriber Restrictions	Prescribed by or in consultation with a nephrologist.
Coverage Duration	New starts will be authorized for 9 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

FINTEPLA

Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using one generic anticonvulsant for appropriate indications.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

FIRDAPSE

Products Affected

- FIRDAPSE

PA Criteria	Criteria Details
Exclusion Criteria	History of seizures.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

FLUCYTOSINE

Products Affected

- *flucytosine oral*

PA Criteria	Criteria Details
Exclusion Criteria	Complete dihydropyrimidine dehydrogenase (DPD) enzyme deficiency
Required Medical Information	Attestation member is taking in combination with amphotericin B.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

GALAFOLD

Products Affected

- GALAFOLD

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed must be a geneticist, cardiologist, nephrologist or specialist experienced in the treatment of Fabry disease.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

GATTEX

Products Affected

- GATTEX

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: attestation of 1) Colonoscopy and upper gastrointestinal endoscopy with removal of polyps within six months prior to starting treatment for adults or 2) Fecal occult blood testing within six months prior to starting treatment for pediatric patients and 3) Baseline laboratory assessments (bilirubin, alkalinephosphatase, lipase, and amylase). For continuation of therapy or reauthorization: Documentation is provided that the member has obtained a clinical benefit (e.g. reduction in parenteral fluid volume, reduction in number of days receiving parenteral nutrition).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

GLP-1 AGONISTS

Products Affected

- *liraglutide*
- MOUNJARO SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- OZEMPIC
- 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 SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	The member has an indication of weight loss/obesity only or type 1 diabetes. The member has concurrent use of any GLP-1 receptor agonist.
Required Medical Information	The member has a documented diagnosis via chart notes of type 2 diabetes.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Diabetes: patient has diagnosis of type 2 diabetes mellitus. All other indications: patient must have medically accepted indication.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

GNRH AGONISTS

Products Affected

- ELIGARD
- FIRMAGON (240 MG DOSE)
- FIRMAGON SUBCUTANEOUS SOLUTION RECONSTITUTED 80 MG
- *leuprolide acetate (3 month)*
- *leuprolide acetate injection*
- LUPRON DEPOT (1-MONTH)
- LUPRON DEPOT (3-MONTH)
- LUPRON DEPOT (4-MONTH)
- LUPRON DEPOT (6-MONTH)
- LUTRATE DEPOT
- TRELSTAR MIXJECT

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Fibroids and endometriosis: 6 months. All other indications: end of contract year.
Other Criteria	If the medication request is for the treatment of prostate cancer and if the request is for any other GnRH agonist other than Eligard or leuprolide, the patient must have a documented trial of, contraindication to, or medical reason for not using Eligard or leuprolide to treat their prostate cancer.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

GOCOVRI

Products Affected

- GOCOVRI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	New starts: trial of, contraindication to, or medical reason for not using generic amantadine. Continuation of therapy or reauthorization: Member demonstrates clinical benefit (i.e. improvement in levodopa-induced dyskinesia or decreased off episodes).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

GROWTH HORMONES

Products Affected

- GENOTROPIN MINIQUICK SUBCUTANEOUS PREFILLED SYRINGE
- GENOTROPIN SUBCUTANEOUS CARTRIDGE
- NGENLA
- OMNITROPE SUBCUTANEOUS SOLUTION CARTRIDGE
- OMNITROPE SUBCUTANEOUS SOLUTION RECONSTITUTED
- SKYTROFA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for growth hormone deficiency: Documentation showing bone age testing, height, weight, and Growth Hormone Stimulation Test results OR Insulin Growth Factor 1 level. For continuation of therapy or reauthorization for growth hormone deficiency: documentation (medical records) showing positive response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an endocrinologist or nephrologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts for growth hormone deficiency: 1) If the request is not for Genotropin, trial of, contraindication to, or medical reason for not using Genotropin. For requests for all other medically accepted indications other than growth hormone deficiency, the request will be approved for products other than Skytrofa.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

HEPCLUDEX

Products Affected

- HEPCLUDEX

PA Criteria	Criteria Details
Exclusion Criteria	Patients with decompensated cirrhosis
Required Medical Information	N/A
Age Restrictions	18 years of age or older
Prescriber Restrictions	Prescribed by or in consultation with a hepatologist, gastroenterologist, or infectious disease specialist
Coverage Duration	Request will be authorized until the end of the contract year
Other Criteria	Initial therapy: (1) documented diagnosis of viral hepatitis D infection confirmed with a positive HDV antibody test or HDV RNA PCR 6 months prior to the start of treatment AND (2) current positive HDV RNA PCR result AND (3) documentation of elevated pretreatment ALT AND (4) patient does not have cirrhosis OR has compensated cirrhosis. Continuation of therapy: patient demonstrates clinical benefit (e.g. undetectable HDV RNA or decrease from baseline, ALT decline, decrease in liver stiffness, etc.).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

HEREDITARY ANGIOEDEMA AGENTS

Products Affected

- CINRYZE
- HAEGARDA
- ORLADEYO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an allergist, immunologist, rheumatologist or hematologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For continuation of therapy or reauthorization: Documentation has been provided that patient has clinically benefited from medication.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK MEDICATION

Products Affected

- *benztropine mesylate oral*
- *cyproheptadine hcl oral syrup 2 mg/5ml*
- *cyproheptadine hcl oral tablet*
- *diphenoxylate-atropine oral liquid*
- *diphenoxylate-atropine oral tablet 2.5-0.025 mg*
- *dipyridamole oral*
- *ergotamine-caffeine*
- *glyburide micronized oral tablet 3 mg, 6 mg*
- *glyburide oral tablet 1.25 mg, 2.5 mg, 5 mg*
- *glyburide-metformin oral tablet 1.25-250 mg*
- *glyburide-metformin oral tablet 2.5-500 mg, 5-500 mg*
- *guanfacine hcl er oral tablet extended release 24 hour 1 mg, 2 mg, 3 mg, 4 mg*
- *guanfacine hcl oral*
- *hydroxyzine hcl oral syrup 10 mg/5ml*
- *hydroxyzine hcl oral tablet*
- *hydroxyzine pamoate oral*
- *indomethacin oral capsule 25 mg, 50 mg*
- *ketorolac tromethamine oral*
- *nifedipine oral*
- *promethazine hcl oral solution 6.25 mg/5ml*
- *promethazine hcl oral tablet*
- *promethazine hcl rectal suppository 12.5 mg, 25 mg*
- *promethazine-phenylephrine*
- *promethegan rectal suppository 50 mg*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: 1) the benefits of treatment with the drug outweigh the potential risks and 2) the risks and side effects have been discussed and will be monitored.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

HIGH RISK MEDICATION - PROTECTED CLASS DRUGS

Products Affected

- *amitriptyline hcl oral*
- *amoxapine*
- *clomipramine hcl oral*
- *nortriptyline hcl oral*
- *perphenazine-amitriptyline*
- *phenobarbital oral elixir*
- *phenobarbital oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: 1) the benefits of treatment with the drug outweigh the potential risks and 2) the risks and side effects have been discussed and will be monitored.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK MEDICATION, BUTALBITAL

Products Affected

- *bac (butalbital-acetamin-caff)*
- *butalbital-acetaminophen oral tablet 50-325 mg*
- *butalbital-apap-caff-cod oral capsule 50-325-40-30 mg*
- *butalbital-apap-caffeine oral capsule 50-325-40 mg*
- *butalbital-apap-caffeine oral solution*
- *butalbital-apap-caffeine oral tablet 50-325-40 mg*
- *butalbital-asa-caff-codeine*
- *butalbital-aspirin-caffeine oral capsule*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: 1) the benefits of treatment with the drug outweigh the potential risks and 2) the risks and side effects have been discussed and will be monitored.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using an oral NSAID.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

HIGH RISK MEDICATION, MEGESTROL

Products Affected

- *megestrol acetate oral suspension 40 mg/ml, 400 mg/10ml, 625 mg/5ml*
- *megestrol acetate oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: the benefits of treatment with the drug outweigh the potential risks identified for people 65 years old and older.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK MEDICATION, SHORT TERM MUSCLE RELAXANT

Products Affected

- *carisoprodol oral*
- *chlorzoxazone oral tablet 500 mg*
- *cyclobenzaprine hcl oral tablet 10 mg, 5 mg*
- *metaxalone oral tablet 800 mg*
- *methocarbamol oral tablet 500 mg, 750 mg*
- *orphenadrine citrate er*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: 1) the benefits of treatment with the drug outweigh the potential risks identified for people 65 years old and older, and 2) the risks and side effects have been discussed and will be monitored.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	New starts will be authorized for 30 days. Continuation of therapy or reauth will be for 90 days.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK MEDICATION, SLEEP AGENTS

Products Affected

- *eszopiclone*
- *temazepam*
- *zaleplon oral capsule 10 mg, 5 mg*
- *zolpidem tartrate er*
- *zolpidem tartrate oral tablet 10 mg*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For patients 65 years old and older the prescriber has documented: 1) the benefits of treatment with the drug outweigh the potential risks identified for people 65 years old and older, and 2) the risks and side effects have been discussed and will be monitored. For zolpidem immediate release 10mg and zolpidem ER: trial of or medical reason for not using zolpidem immediate release 5mg.
Age Restrictions	Prior authorization only applies to members 65 years old or older.
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

HYFTOR

Products Affected

- HYFTOR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: documentation of diagnosis of tuberous sclerosis with facial angiofibroma. For continuation of therapy or reauthorization: documentation that the member has experienced a clinical benefit from treatment (e.g. improvement in size and color of angiofibroma).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist or provider who specializes in the treatment of genetic or dermatologic disorders.
Coverage Duration	New starts: 3 months. Cont. of therapy or reauthorization: until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ICATIBANT

Products Affected

- *icatibant acetate subcutaneous solution prefilled syringe*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an immunologist, allergist, rheumatologist, or hematologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ILARIS

Products Affected

- ILARIS SUBCUTANEOUS SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation was submitted indicating that the member was evaluated for active or latent TB infection (i.e. tuberculin skin test)
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For sJIA: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ILUMYA

Products Affected

- ILUMYA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For psoriasis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

IMBRUVICA

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an oncologist or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts for treatment of graft-versus-host disease (GVHD): Trial of, contraindication to, or medical reason for not using a systemic corticosteroid. For continuation of therapy of for treatment of GVHD: documentation of clinical benefit from use of the drug (i.e. symptom improvement, reduction in corticosteroid dose). For all other indications, approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

INCRELEX

Products Affected

- INCRELEX

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

JAKAFI

Products Affected

- JAKAFI
- JAKAFI XR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an oncologist or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts for treatment of graft-versus-host disease (GVHD): Trial of, contraindication to, or medical reason for not using a systemic corticosteroid. For continuation of therapy of for treatment of GVHD: documentation of clinical benefit from use of the drug (i.e. symptom improvement, reduction in corticosteroid dose). For all other indications, approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

KALYDECO

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	Combination use with Orkambi, Symdeko, or Trikafta.
Required Medical Information	Documentation of CFTR gene that is responsive to ivacaftor treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or an expert in the treatment of cystic fibrosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

KERENDIA

Products Affected

- KERENDIA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: 1) Documentation of diagnosis of chronic kidney disease (CKD) due to type 2 diabetes mellitus OR heart failure (HF) with left ventricular ejection fraction (LVEF) greater than or equal to 40% AND AND 2) Documentation of serum potassium levels less than or equal to 5 mEq/L AND 3) eGFR greater than or equal to 25ml/min/1.73 m2 AND 4) Documentation that member is taking Kerendia in combination with an angiotensin-converting enzyme inhibitor (ACEi) or angiotensin receptor blocker (ARB) at maximum tolerated doses or documentation has been provided that the member is unable to tolerate ACEi or ARB. For continuation of therapy or reauthorization for patients with CKD due to type 2 diabetes mellitus: 1) Documentation of serum potassium levels less than or equal to 5.5 mEq/L AND 2) Documentation that member is taking Kerendia in combination with an ACEi or ARB at maximum tolerated doses or documentation has been provided that the member is unable to tolerate ACEi or ARB. For continuation of therapy or reauthorization for patients with HF with LVEF greater than or equal to 40%: 1) Documentation of serum potassium levels less than 6 mEq/L AND 2) Documentation that member is taking Kerendia in combination with an ACEi or ARB at maximum tolerated doses or documentation has been provided that the member is unable to tolerate ACEi or ARB.
Indications	All Medically-accepted Indications.

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

KEVZARA

Products Affected

- KEVZARA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For polymyalgia rheumatica (PMR): Trial of, medical reason for not using, or contraindication to corticosteroids. For pJIA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

KINERET

Products Affected

- KINERET SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For cryopyrin-associated periodic syndromes or deficiency of interleukin-1 receptor antagonist: Approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LEQEMBI IQLIK

Products Affected

- LEQEMBI IQLIK
- LEQEMBI IQLIK (500 MG DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist, geriatric psychiatrist, or geriatrician who specializes in treating dementia
Coverage Duration	Initial: 6 months. Continuation of therapy: end of the contract year.
Other Criteria	Initiation: (1) the patient has documentation of 18 months of treatment with intravenous Leqembi prior to initiation of subcutaneous maintenance therapy & (2) prior to initation with intravenous Leqembi, the patient has an initial diagnosis of mild cognitive impairment (MCI) caused by Alzheimer's Disease (AD) or mild AD consistent with Stage 3 or Stage 4 Alzheimer's disease & (3) documentation of both of the following: a positive result for the presence of beta-amyloid plaques & baseline Magnetic Resonance Imaging (MRI) scan prior to intiation with intravenous Leqembi. Continuation of therapy: patient has a positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

LEQSELVI

Products Affected

- LEQSELVI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation was submitted indicating that the member was evaluated for active or latent TB infection (i.e. tuberculin skin test).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documentation of confirmed diagnosis and other causes of hair loss have been ruled out.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

LITFULO

Products Affected

- LITFULO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Initial: (1) documented diagnosis via chart notes of severe alopecia areata AND (2) patient is not receiving in combination with either of the following: (i) Targeted immunomodulator (i.e. Olumiant, Enbrel, Cimzia, Simponi, Orencia, adalimumab, Xeljanz, Rinvoq) OR (ii) potent immunosuppressant. Continuation of therapy: documentation of positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documentation of confirmed diagnosis and other causes of hair loss have been ruled out.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

LIVMARLI

Products Affected

- LIVMARLI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist or hepatologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts: 1) Trial of, contraindication to, or medical reason for not using both of the following: cholestyramine AND rifampin. 2) Prescriber attests that the member has cholestasis 3) Baseline serum bile acid level is provided. 4) Documentation of patients weight. For continuation of therapy or reauthorization: 1) Documentation submitted indicating the member has had all of the following: an improvement in pruritis (e.g. improved observed scratching, decreased sleep disturbances/nighttime awakenings due to scratching, etc.) AND reduction in serum bile acid level from baseline. 2) Prescriber attests that patient has had no evidence of hepatic decompensation (e.g. variceal hemorrhage, ascites, hepatic encephalopathy, portal hypertension, etc.). 3) Documentation of patients weight.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LIVTENCITY

Products Affected

- LIVTENCITY

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Cytomegalovirus (CMV): (1) Documented diagnosis of CMV infection AND (2) Member is a recipient of one of the following: (a) hematopoietic stem cell transplant, (b) solid organ transplant AND (3) patient has tried and failed treatment with valganciclovir, ganciclovir, cidofivir, or foscarnet AND (4) patient weighs greater than or equal to 35 kg.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a transplant, oncologist, or infectious disease specialist.
Coverage Duration	Request will be authorized for 8 weeks.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LODOCO

Products Affected

- LODOCO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be, or in consultation with a specialist in the treatment of cardiovascular disease, such as a cardiologist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documentation that patient has established atherosclerotic disease or multiple risk factors for cardiovascular disease AND documentation that patient does not have pre-existing blood dyscrasias (ex. leukopenia, thrombocytopenia) and patient does not have renal failure (CrCl less than 15 ml/min) or severe hepatic impairment
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

LUCEMYRA

Products Affected

- *lofexidine hcl*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 14 days.
Other Criteria	Patient must have trial of, contraindication to, or medical reason for not using clonidine.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LUPKYNIS

Products Affected

- LUPKYNIS

PA Criteria	Criteria Details
Exclusion Criteria	Concurrent use with cyclophosphamide.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be rheumatologist, nephrologist, or other specialist in the treatment of autoimmune disorders.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts: 1) Documentation of urine protein/creatinine ratio (UPCR), 2) Documentation that the member has a baseline eGFR greater than 45 mL/min/1.73m2 or that benefit outweighs risk of using this medication at current eGFR, and 3) Concurrent use of or medical reason for not using background immunosuppressive therapy regimen. For continuation of therapy or reauthorization: Documentation of improvement in renal function (i.e. reduction in UPCR or no confirmed decrease from baseline eGFR greater than or equal to 20%).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

LYBALVI

Products Affected

- LYBALVI

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with opioids.
Required Medical Information	Attestation from the provider that the member has had an opioid-free period of a minimum of 7 days after last use of shorting-acting opioids and 14 days from last use of long-acting opioids before initiating Lybalvi.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documented trial of, contraindication to, or medical reason for not using at least two generic antipsychotics, one of which must be generic olanzapine.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

MAVYRET

Products Affected

- MAVYRET

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Detectable HCV RNA viral load prior to treatment within 6 months of request. In addition, documentation of treatment history, and if cirrhotic, documentation of compensated or decompensated cirrhosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 8-16 weeks as per AASLD-IDSA guidance.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

METHOTREXATE ORAL SOLUTION

Products Affected

- JYLAMVO
- XATMEP

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an oncologist, a rheumatologist, a dermatologist, or other appropriate specialist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Polyarticular Juvenile Idiopathic Arthritis (pJIA) - Initial: (1) patient had been diagnosed with pJIA AND (2) patient had tried, intolerant or has medical reason for not using at least one non-steroidal anti-inflammatory agents (NSAIDs) AND methotrexate.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

METHYLTESTOSTERONE

Products Affected

- *methyltestosterone oral*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

METYROSINE

Products Affected

- *metyrosine*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of one of the following: 1) Concurrent use of alpha adrenergic blockers, 2) Medical reason for being unable to use an alpha adrenergic blocker, OR 3) Patient is not a candidate for surgical resection and requires long term treatment with metyrosine.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

MIFEPRISTONE

Products Affected

- *mifepristone oral tablet 300 mg*

PA Criteria	Criteria Details
Exclusion Criteria	For all members patient must not be currently on simvastatin, lovastatin, cyclosporine, dihydroergotamine, ergotamine, fentanyl, pimozide, quinidine, sirolimus, and tacrolimus.
Required Medical Information	Reviewer will verify available claim history to confirm member is not taking simvastatin, lovastatin, cyclosporine, dihydroergotamine, ergotamine, fentanyl, pimozide, quinidine, sirolimus or tacrolimus concurrently with mifepristone.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

MIGLUSTAT

Products Affected

- *miglustat*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts, documentation of diagnosis for mild to moderate type 1 Gaucher disease. For continuation of therapy or reauthorization: documentation of clinical benefit from use of the drug (i.e. increased platelet count, improvement in anemia, PFT's, improvement in radiographic scans, improved quality of life).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a specialist in treatment of Gaucher's disease
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

MULTIPLE SCLEROSIS AGENTS

Products Affected

- BAFIERTAM
- BETASERON SUBCUTANEOUS KIT
- *cladribine (10 tabs)*
- *cladribine (4 tabs)*
- *cladribine (5 tabs)*
- *cladribine (6 tabs)*
- *cladribine (7 tabs)*
- *cladribine (8 tabs)*
- *cladribine (9 tabs)*
- *cladribine 10 mg tablet therapy pack oral*
- *dimethyl fumarate oral capsule delayed release 120 mg, 240 mg*
- *dimethyl fumarate starter pack oral capsule delayed release therapy pack*
- *fingolimod hcl*
- *glatiramer acetate subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml*
- *glatopa subcutaneous solution prefilled syringe 20 mg/ml, 40 mg/ml*
- KESIMPTA
- MAYZENT
- MAYZENT STARTER PACK
- PONVORY
- PONVORY STARTER PACK
- REBIF REBIDOSE SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- REBIF REBIDOSE TITRATION PACK SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- REBIF SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- REBIF TITRATION PACK SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- TASCENSO ODT
- *teriflunomide*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If the medication request is for glatiramer, Glatopa, or dimethyl fumarate, the request will be approved. If the member is over 17 years of age and the request is not for glatiramer, Glatopa, or dimethyl fumarate for multiple sclerosis, the member must have a documented trial of, contraindication to or a medical reason for not using 2 of the following dimethyl fumarate,

PA Criteria	Criteria Details
	glatiramer, Glatopa, teriflunomide, or fingolimod. If the request is for fingolimod and the member is 17 years of age or younger, the request will be approved.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

MYFEMBREE

Products Affected

- MYFEMBREE

PA Criteria	Criteria Details
Exclusion Criteria	Patient has history of osteoporosis or hepatic impairment.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an OB, gynecologist or reproductive endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts for menorrhagia: Trial of, contraindication to, or medical reason for not using an estrogen-progestin contraceptive therapy. For new starts if one of the following drugs has been tried previously, a trial of estrogen-progestin contraceptive therapy is not required: gonadotropin-releasing hormone (GnRH) agonists or tranexamic acid. New starts for endometriosis: Trial of, contraindication to, or medical reason for not using the following concurrently for endometriosis: analgesic pain reliever (e.g. NSAIDs, COX-2 inhibitors) AND either combined estrogen-progestin oral contraceptive, progestin (e.g. medroxyprogesterone acetate, norethindrone), gonadotropin-releasing hormone (GnRH) agonists (e.g. Lupron Depot), OR danazol. For continuation of therapy or reauthorization both of the following are required: 1) Treatment does not exceed the eligible maximum lifetime treatment duration of 2 years, and 2) Documentation has been provided that the member has obtained clinical benefit from medication (e.g. reduced menstrual bleeding from baseline, pain relief).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

MYQORZO

Products Affected

- MYQORZO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist.
Coverage Duration	Initial: 6 months. Continuation of therapy: end of contract year.
Other Criteria	For all new starts, documentation of ALL of the following must be provided: 1) Diagnosis of symptomatic obstructive hypertrophic cardiomyopathy (oHCM) AND 2) Patient has a left ventricular ejection fraction (LVEF) greater than or equal to 55% AND 3) Assessment of Valsalva left ventricular outflow tract (LVOT) gradient AND 4) Trial of, medical reason for not using or contraindication to BOTH of the following: Beta blockers (i.e. metoprolol, propranolol, atenolol) AND Non-dihydropyridine calcium channel blockers (i.e. verapamil, diltiazem). For continuation of therapy or reauthorization, all of the following must be provided: 1) Provider attests that the member has demonstrated a clinically meaningful benefit from therapy AND 2) Member must also have a left ventricular ejection fraction (LVEF) greater than or equal to 40%.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

NASAL ANTISEIZURE AGENTS

Products Affected

- NAYZILAM
- VALTOCO 10 MG DOSE
- VALTOCO 15 MG DOSE NASAL LIQUID THERAPY PACK 2 X 7.5 MG/0.1ML
- VALTOCO 20 MG DOSE NASAL LIQUID THERAPY PACK 2 X 10 MG/0.1ML
- VALTOCO 5 MG DOSE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	Nayzilam: 12 years of age or older. Valtoco: 2 years of age or older.
Prescriber Restrictions	Initial therapy only: prescribed by or in consultation with a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

NITISINONE

Products Affected

- *nitisinone*
- ORFADIN ORAL SUSPENSION

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a geneticist, metabolic specialist, hepatologist, or liver transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Hereditary Tyrosinemia Type 1 (HT-1): diagnosis was confirmed by genetic testing confirming a mutation of the FAH gene OR elevated levels of succinylacetone.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

NON-AMPHETAMINE CENTRAL NERVOUS SYSTEM AGENTS

Products Affected

- *armodafinil*
- *modafinil oral*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

NUCALA

Products Affected

- NUCALA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	New starts for severe asthma: 1) Baseline blood eosinophil count greater than or equal to 150 cells per microliter AND 2) symptoms with equal to or greater than 1 exacerbations in the previous 12 months while on a high-dose inhaled corticosteroid AND 3) the patient has a documented trial of, contraindication to, or medical reason for not using both Dupixent and Fasenra. New starts for eosinophilic granulomatosis with polyangiitis (EGPA): 1) trial of, contraindication to, or medical reason for not using one of the following medications: cyclophosphamide or methotrexate AND 2) the patient has a documented trial of, contraindication to, or medical reason for not using Fasenra. New starts for hypereosinophilic syndrome without an identifiable non-hematologic secondary cause: 1) 2 or more flares within the past 12 months AND 2) trial of, contraindication to, or medical reason for not using oral corticosteroids. New starts for chronic rhinosinusitis with nasal polyps: 1) the patient has a documented trial of, contraindication to, or medical reason for not using both Dupixent and Xolair. Continuation of therapy or re-authorization for all indications: clinical benefit from use of the drug. COPD - initial: (1) documented diagnosis of COPD AND (2) Documentation of blood eosinophil count of greater than or equal to 300 cells/microliter AND (3) Nucala is used as add-on maintenance treatment AND (4) patient has had a trial of, contraindication to, or medical reason for not using at least two of the following: a) long acting beta2-agonist (LABA), b) long acting muscarinic antagonist (LAMA) or c) inhaled

PA Criteria	Criteria Details
	corticosteroid (ICS) AND (5) documentation of COPD exacerbation within the past 12 months. Continuation of therapy: patient has positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

NUEDEXTA

Products Affected

- *dextromethorphan-quinidine*
- NUEDEXTA

PA Criteria	Criteria Details
Exclusion Criteria	Complete atrioventricular (AV) block without implanted pacemaker, or at high risk of complete AV block. History of heart failure. Concomitant use with MAOIs or use of MAOIs within 14 days. Concomitant use with drugs containing quinidine, quinine, or mefloquine. History of quinine-, mefloquine-, dextromethorphan/quinidine-, or quinidine-induced thrombocytopenia, hepatitis, bone marrow depression, or lupus-like syndrome. Non-Part D indications.
Required Medical Information	Confirmation diagnosis is for Part D indication.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist or psychiatrist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

NUPLAZID

Products Affected

- NUPLAZID ORAL CAPSULE
- NUPLAZID ORAL TABLET 10 MG

PA Criteria	Criteria Details
Exclusion Criteria	Patient has a history of dementia-related psychosis.
Required Medical Information	For hallucinations and delusions associated with Parkinson's disease psychosis: documented diagnosis of Parkinson's disease must be made prior to the onset of psychotic symptoms.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

OCTREOTIDE

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for acromegaly: pt meets one of the following (1) inadequate response to surgery and/or radiotherapy OR (2) pt is not an appropriate candidate for surgery and/or radiotherapy OR (3) pt is experiencing negative effects due to tumor size (ex: optic nerve compression). Continuation of therapy or reauthorization: documentation of clinical improvement with therapy.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Continuation of therapy or reauthorization: documentation of clinical improvement with therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

OFEV

Products Affected

- *nintedanib esylate*
- OFEV

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist, rheumatologist, or lung transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For a diagnosis of idiopathic pulmonary fibrosis: 1) Documentation of disease as demonstrated on a high resolution CT scan or through lung biopsy and 2) Documented trial of, contraindication to, or medical reason for not using pirfenidone. For a diagnosis of systemic sclerosis-associated interstitial lung disease (SSc-ILD): documented trial of, contraindication to, or medical reason for not using mycophenolate mofetil or cyclophosphamide. For a diagnosis of chronic fibrosing interstitial lung diseases (ILDs) with a progressive phenotype: documentation is provided confirming diagnosis.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

OPSUMIT

Products Affected

- *macitentan*
- OPSUMIT

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (PAH) World Health Organization (WHO) Group I - Initial: [Note: documentation required] (1) PAH was confirmed by right heart catheterization AND (2) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg AND (3) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg AND (4) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or cardiologist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using sildenafil.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ORAL ANTIPSYCHOTICS

Products Affected

- BYSANTI
- BYSANTI TITRATION PACK A
- BYSANTI TITRATION PACK B
- BYSANTI TITRATION PACK C
- CAPLYTA
- COBENFY
- COBENFY STARTER PACK
- FANAPT
- FANAPT TITRATION PACK A
- FANAPT TITRATION PACK B ORAL TABLET
- FANAPT TITRATION PACK C ORAL TABLET
- OPIPZA ORAL FILM 10 MG, 2 MG, 5 MG
- VRAYLAR ORAL CAPSULE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Schizophrenia: 1) request is for Caplyta, Cobenfy, Fanapt, Opipza, Bysanti, or Vraylar AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Manic or mixed episodes associated with bipolar l disorder: 1) request is for Fanapt, Opipza, Bysanti, or Vraylar AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Major depressive disorder associated with bipolar l or II disorder: 1) request is for Caplyta AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Adjunctive treatment of major depressive disorder: 1) request is for Caplyta, Opipza or Vraylar AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics AND 3) if the request is for Vraylar: provider attestation that the member is concurrently using an antidepressant. Irritability associated with autistic disorder: 1) request is for Opipza AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Treatment of Tourette's disorder: 1) request is

PA Criteria	Criteria Details
	for Opipza AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Bipolar I disorder: 1) request is for Opipza AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics. Depressed bipolar I disorder: 1) request is for Vraylar AND 2) trial of, contraindication to, or medical reason for not using two generic antipsychotics.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ORENCIA

Products Affected

- ORENCIA CLICKJECT
- ORENCIA INTRAVENOUS
- ORENCIA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 125 MG/ML, 50 MG/0.4ML, 87.5 MG/0.7ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For pJIA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For PsA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Xeljanz, Enbrel, Cosentyx, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For acute graft versus host disease: Attestation member is taking in combination with a calcineurin inhibitor and methotrexate.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ORIAHNN

Products Affected

- ORIAHNN

PA Criteria	Criteria Details
Exclusion Criteria	Patient has history of osteoporosis or hepatic impairment.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an OB, gynecologist or reproductive endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: Trial of, contraindication to, or medical reason for not using an estrogen-progestin contraceptive therapy. For new starts if one of the following drugs has been tried previously, a trial of estrogen-progestin contraceptive therapy is not required: gonadotropin-releasing hormone (GnRH) agonists or tranexamic acid. For continuation of therapy or reauthorization both of the following are required: 1) Treatment does not exceed the eligible maximum lifetime treatment duration of 2 years, and 2) Documentation has been provided that the member has obtained clinical benefit from medication (e.g. reduced menstrual bleeding from baseline).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ORILISSA

Products Affected

- ORILISSA

PA Criteria	Criteria Details
Exclusion Criteria	Patient has osteoporosis or severe hepatic impairment.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an OB or gynecologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using the following concurrently for endometriosis: analgesic pain reliever (e.g. NSAIDs, COX-2 inhibitors) AND either combined estrogen-progestin oral contraceptive, progestin (e.g. medroxyprogesterone acetate, norethindrone), gonadotropin-releasing hormone (GnRH) agonists (e.g. Lupron Depot), OR danazol. For continuation of therapy or reauthorization both of the following are required: 1) Treatment does not exceed the eligible maximum lifetime treatment duration of 2 years for 150mg tablet or 6 months for 200mg tablet, and 2) Documentation has been provided that the member has obtained clinical benefit from the medication.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ORKAMBI

Products Affected

- ORKAMBI

PA Criteria	Criteria Details
Exclusion Criteria	Combination use with Kalydeco, Symdeko, or Trikafta.
Required Medical Information	Documentation of CFTR gene that is responsive to lumacaftor-ivacaftor treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or an expert in the treatment of cystic fibrosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

OTEZLA

Products Affected

- OTEZLA
- OTEZLA XR
- OTEZLA/OTEZLA XR INITIATION PK

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For moderate to severe psoriasis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For PsA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Xeljanz, Enbrel, Cosentyx, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For Behcet's Syndrome or mild psoriasis: Approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

OXERVATE

Products Affected

- OXERVATE

PA Criteria	Criteria Details
Exclusion Criteria	Treatment duration greater than 16 weeks per affected eye(s).
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by an ophthalmologist or an optometrist.
Coverage Duration	Initial: 8 weeks. Continuation of therapy: additional 8 weeks.
Other Criteria	Initial: confirmed diagnosis. Continuation of therapy: patient has previously received less than or equal to 8 weeks of treatment per affected eye(s) and the patient has a recurrence of neurotrophic keratitis.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

OXYCODONE ER

Products Affected

- OXYCONTIN ORAL TABLET ER 12 HOUR ABUSE-DETERRENT 10 MG, 15 MG, 20 MG, 30 MG, 40 MG, 60 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Members being treated for active cancer diagnoses, sickle cell diagnoses, those in hospice care, or receiving palliative care will be excluded from the concurrent benzodiazepine and muscle relaxant therapy requirement. For new starts, ALL of the following are required: (1) Member has documented history of receiving an immediate-release opioid, (2) Member has a documented trial of, contraindication to, or medical reason for not using long-acting morphine sulfate, (3) If member is on concurrent benzodiazepines and/or muscle relaxant therapy, the prescriber attests that concurrent therapy is medically necessary, (4) Member is not being treated for substance abuse with buprenorphine-containing products. For continuing therapy, ALL of the following are required: (1) Member's pain has been assessed within the last 6 months, (2) Member has demonstrated clinical improvement in pain and function on current medication regimen, (3) If member is on concurrent benzodiazepines and/or muscle relaxant therapy, the prescriber attests that concurrent therapy is medically necessary, (4) Member is not being treated for substance abuse with buprenorphine-containing products.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A

PA Criteria	Criteria Details
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PCSK9 INHIBITORS

Products Affected

- REPATHA
- REPATHA SURECLICK

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For all diagnoses (besides for risk reduction of major cardiovascular events): documented diagnosis of medically accepted indication. For risk reduction of major cardiovascular (CV) events (CV death, myocardial infarction, stroke, unstable angina requiring hospitalization, or coronary revascularization): 1) Patient has an increased risk of major CV events such as death from CV disease, heart attack, stroke, certain types of chest pain conditions (including but not limited to, unstable angina) requiring hospitalization or certain types of heart surgery and (2) tried or has contraindication to high intensity statin.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PEGINTERFERON

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For Hepatitis C: 1) Labs within 3 months of request: liver function tests and detectable HCV RNA viral load. 2) Documentation of genotype, treatment history, and if cirrhotic, documentation of compensated or decompensated cirrhosis. For Hepatitis B: 1) Labs within 3 months of request: ALT/AST, and 2) HBeAg status. For polycythemia vera, approve.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a gastroenterologist, hepatologist, infectious disease doctor or transplant specialist.
Coverage Duration	Request will be authorized for 24 to 48 weeks as defined by compendia.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

PENICILLAMINE

Products Affected

- *penicillamine oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For RA: Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz. For other indications, approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PERSERIS

Products Affected

- PERSERIS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented history of receiving oral risperidone without any clinically significant side effects.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least two of the following: Abilify Maintena, Abilify Asimtufii, Invega Sustenna, Invega Trinza, Invega Hafyera, or Risperidone Microspheres ER.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PHENOXYBENZAMINE

Products Affected

- *phenoxybenzamine hcl oral*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using doxazosin.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PIRFENIDONE

Products Affected

- *pirfenidone*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For idiopathic pulmonary fibrosis, documentation of all of the following: 1) confirmation of diagnosis on high resolution CT scan or through lung biopsy AND 2) FVC greater than or equal to 50% of the predicted value.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or lung transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

POSACONAZOLE

Products Affected

- *posaconazole oral*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Documentation of a consultation with an infectious disease specialist, a transplant specialist, or an oncologist.
Coverage Duration	28 days for oropharyngeal candidiasis, end of contract year for other indications
Other Criteria	For treatment of oropharyngeal candidiasis: trial of, contraindication to, or medical reason for not using fluconazole or itraconazole. For prophylaxis of invasive aspergillus infections due to being severely immunocompromised: trial of, contraindication to, or medical reason for not using voriconazole.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PRETOMANID

Products Affected

- *pretomanid*

PA Criteria	Criteria Details
Exclusion Criteria	MDR-TB that is not treatment-intolerant or nonresponsive to standard therapy
Required Medical Information	Documentation of use in combination with bedaquiline and linezolid.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist.
Coverage Duration	Request will be authorized for 26 weeks.
Other Criteria	Documentation of prior trial of or medical reason for not using first-line TB regimen containing isoniazid and rifampin.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PREVYMIS

Products Affected

- PREVYMIS ORAL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a hematologist, oncologist, infectious disease, or transplant specialist.
Coverage Duration	Request will be authorized for 6 months.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

PROMACTA

Products Affected

- *eltrombopag olamine oral packet 12.5 mg, 25 mg*
- *eltrombopag olamine oral tablet 12.5 mg, 25 mg, 50 mg, 75 mg*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For persistent or chronic immune thrombocytopenia (ITP): Documented baseline platelet count less than 30,000 cells/ microL. For severe aplastic anemia: Documentation of baseline platelet count less than 20,000 cells/microL OR platelet count less than 30,000 cells/microL with bleeding OR reticulocyte count less than 20,000 cells/microL OR absolute neutrophil count less than 500 cells/microL. For thrombocytopenia in patients with Hepatitis C infection: documented baseline platelet count less than 75,000 cells/microL.
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For persistent or chronic immune thrombocytopenia (ITP):Trial of, contraindication to, or medical reason for not using glucocorticosteroids. For severe aplastic anemia: Trial of, contraindication to, or medical reason for not using at least one immunosuppressive agent.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PYRUKYND

Products Affected

- PYRUKYND
- PYRUKYND TAPER PACK

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: 1) documentation of diagnosis and 2) baseline hemoglobin level. For continuation of therapy or reauthorization: documentation of clinical improvement (e.g. reduction in number of blood transfusions, or increase or stabilization in hemoglobin level). If the criteria are not met, may authorize up to 14 days of a Pyrukynd Taper Pack to allow for tapering.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hematologist.
Coverage Duration	New starts: 6 mo. Cont of therapy or reauth: end of contract yr. Denial: 14 days for dose tapering.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

RADICAVA

Products Affected

- RADICAVA ORS
- RADICAVA ORS STARTER KIT

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	New starts: 6 months. Cont. of therapy or reauthorization: until end of contract year.
Other Criteria	For new starts: 1) documentation of ALS functional rating scale (ALSFRS-R) score and 2) documentation that the member has been on riluzole, is beginning therapy as an adjunct to treatment with Radicava, or provider has provided a medical reason why patient is unable to use riluzole. For continuation of therapy or reauthorization: documentation from provider of clinical stabilization in symptoms (e.g. stabilization of ALS functional rating scale (ALSFRS-R) score).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RAVICTI

Products Affected

- *glycerol phenylbutyrate*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Provider is a geneticist, metabolic specialist, gastroenterologist, hepatologist, or liver transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using sodium phenylbutyrate.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RECORLEV

Products Affected

- RECORLEV

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using ketoconazole tablets.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RELISTOR

Products Affected

- RELISTOR ORAL
- RELISTOR SUBCUTANEOUS SOLUTION
- RELISTOR SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For opioid induced constipation, chronic non-cancer pain: Patient must have documented trial of or medical reason for not using at least two the following: 1) lubiprostone, OR 2) lactulose OR 3) Movantik. Additionally, patient must have a medical reason for not being able to use oral Relistor in order to receive Relistor injection. For opioid-induced constipation, in patients with advanced illness or pain caused by active cancer requiring opioid dosage escalation for palliative care: Approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

REVCOVI

Products Affected

- REVCOVI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Patient has diagnosis of adenosine deaminase severe combined immune deficiency (ADA-SCID) confirmed by one of the following: 1) documentation of deficiency or absence of adenosine deaminase OR 2) genetic testing revealing mutations in both alleles of the ADA1 gene
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an immunologist, geneticist, hematologist, oncologist, or provider who specializes in the treatment of ADA-SCID or related disorders.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

REXULTI

Products Affected

- REXULTI

PA Criteria	Criteria Details
Exclusion Criteria	Dementia-related psychosis
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For schizophrenia: trial of, contraindication to, or medical reason for not using two generic antipsychotics. For major depressive disorder: trial of, contraindication to, or medical reason for not using to two generic antidepressants. For agitation due to dementia: 1) patient has documented diagnosis of Alzheimer's disease AND 2) medication will not be used on an "as needed" basis
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

REZDIFFRA

Products Affected

- REZDIFFRA

PA Criteria	Criteria Details
Exclusion Criteria	Patients with decompensated cirrhosis.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a hepatologist, gastroenterologist, endocrinologist or specialist in the treatment of liver disease.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Initial therapy: (1) documented diagnosis of noncirrhotic nonalcoholic steatohepatitis (NASH) with moderate to advanced liver fibrosis, (2) documentation of stage F2 to F3 fibrosis as confirmed by a biopsy or a non-invasive test (NIT), (3) the drug is being prescribed at an FDA approved dose according to the member's weight and (4) prescriber attestation to providing lifestyle counseling on nutrition, exercise and avoiding excessive alcohol intake. For reauthorization: (1) the member has shown clinical benefit from the medication (e.g., the resolution of steatohepatitis and no worsening of liver fibrosis, or at least one stage improvement in liver fibrosis and no worsening of steatohepatitis), (2) the member continues to have a fibrosis stage of 3 or less and (3) the drug continues to be prescribed at an FDA approved dose according to the member's weight.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

REZUROCK

Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a hematologist, oncologist, or transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: documented trial of, contraindication to, or medical reason for not using at least two lines of systemic immunosuppressive therapy (e.g. corticosteroids, tacrolimus, mycophenolate mofetil, Imbruvica, or Jakafi), one of which must be a systemic corticosteroid. For continuation of therapy or re-authorization: documentation of clinical benefit from use of the drug (i.e. symptom improvement, reduction in corticosteroid dose).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RUFINAMIDE

Products Affected

- *rufinamide oral suspension 40 mg/ml*
- *rufinamide oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	History of familial Short QT syndrome
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using one alternative generic anticonvulsant for appropriate indications.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RYKINDO

Products Affected

- RYKINDO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented history of receiving oral risperidone without any clinically significant side effects.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least two of the following: Abilify Maintena, Abilify Asimtufii, Invega Sustenna, Invega Trinza, or Risperidone Microspheres ER.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RYLAZE

Products Affected

- RYLAZE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an oncologist, hematologist, or specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SAPROPTERIN

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: documentation of elevated baseline phenylalanine levels. Continuation of therapy or reauthorization: prescriber attests the member has improvement in phenylalanine levels from baseline.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	New starts will be authorized for 3 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SECUADO

Products Affected

- SECUADO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using to one generic antipsychotics.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SEROSTIM

Products Affected

- SEROSTIM SUBCUTANEOUS SOLUTION RECONSTITUTED 4 MG, 5 MG, 6 MG

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a HIV specialist, gastroenterologist, nutritional support specialist or ID specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For initial starts for HIV wasting/cachexia: 1) Member must be on anti-retroviral therapy and 2) Alternative causes of wasting have been ruled out (diarrhea, malignancies, inadequate caloric intake, etc)
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SIGNIFOR

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Member is not a candidate for surgery or surgery was not curative.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SILDENAFIL ORAL

Products Affected

- *sildenafil citrate oral suspension reconstituted*
- *sildenafil citrate oral tablet 20 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Documentation of concurrent nitrate or Adempas use.
Required Medical Information	Pulmonary Arterial Hypertension: Documentation of pulmonary arterial hypertension (PAH) WHO Group I and PAH Functional Class. Reviewer will verify available patient claim history to confirm patient is not using nitrates or Adempas. Secondary Raynaud's phenomenon- Initial: [Note: documentation required] (1) diagnosis of secondary Raynaud's phenomenon AND (2) diagnosis of primary condition which Raynaud's phenomenon is secondary to (e.g., lupus, scleroderma, rheumatoid arthritis, Sjogren's syndrome, thyroid disease). Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist, cardiologist, or rheumatologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For sildenafil suspension: Documentation of trial of, contraindication to, or medical reason for not using sildenafil tablet.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SILIQ

Products Affected

- SILIQ

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For psoriasis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SIMPONI

Products Affected

- SIMPONI SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- SIMPONI SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ankylosing spondylitis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: Enbrel, an adalimumab product, Cosentyx, or Xeljanz, or 2) If utilized within the past 120 days, approve for continuation of therapy. For PsA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Cosentyx, Tremfya, Xeljanz, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For RA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: Enbrel, an adalimumab product, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For adults with UC: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following: an adalimumab product, an ustekinumab product, Tremfya, or Xeljanz or 2) If utilized within the past 120 days, approve for continuation of therapy. For pediatric patients with UC: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

SIRTURO

Products Affected

- SIRTURO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation (consistent with pharmacy claims data, OR for new members to the health plan consistent with medical chart history) that the member is currently taking Sirturo as part of a combination regimen with other antimycobacterial drugs to treat MDR-TB.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist.
Coverage Duration	Request will be authorized for 24 weeks.
Other Criteria	Documentation of prior trial of or medical reason for not using first-line TB regimen containing isoniazid and rifampin.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SODIUM OXYBATE

Products Affected

- *sodium oxybate*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a sleep specialist, pulmonologist, or neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For somnolence associated with narcolepsy in adults: trial of, contraindication to, or medical reason for not using a CNS stimulant (e.g. methylphenidate, modafinil, armodafinil, etc.). For somnolence associated with narcolepsy in pediatric patients: approve. For cataplexy associated with narcolepsy, approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SODIUM PHENYLBUTYRATE

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Provider is a geneticist, metabolic specialist, gastroenterologist, hepatologist, or liver transplant specialist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SOFOSBUVIR/VELPATASVIR

Products Affected

- SOFOSBUVIR-VELPATASVIR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Detectable HCV RNA viral load prior to treatment within 6 months of request. In addition, documentation of treatment history, and if cirrhotic, documentation of compensated or decompensated cirrhosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 12-24 weeks based on AASLD-IDSA guidelines
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SOMAVERT

Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for acromegaly: pt meets one of the following (1) inadequate response to surgery and/or radiotherapy OR (2) pt is not an appropriate candidate for surgery and/or radiotherapy OR (3) pt is experiencing negative effects due to tumor size (ex: optic nerve compression). Continuation of therapy or reauthorization: documentation of clinical improvement with therapy.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SOTYKTU

Products Affected

- SOTYKTU

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For moderate to severe psoriasis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. Active Psoriatic Arthritis- Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Tremfya, Xeljanz, Enbrel, Cosentyx, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SUCRAID

Products Affected

- SUCRAID

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: documentation of diagnosis of congenital sucrase-isomaltase deficiency. For continuation of therapy or reauthorization: Prescriber attests that member has obtained a clinical benefit (e.g. fewer total stools, greater number of hard and formed stools, fewer watery and soft stools, decrease in breath hydrogen output)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SYMDEKO

Products Affected

- SYMDEKO

PA Criteria	Criteria Details
Exclusion Criteria	Combination use with Kalydeco, Orkambi, or Trikafta.
Required Medical Information	Documentation of CFTR gene that is responsive to tezacaftor-ivacaftor treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or an expert in the treatment of cystic fibrosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

SYNAREL

Products Affected

- SYNAREL

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using the following concurrently for endometriosis: analgesic pain reliever (e.g. NSAIDs, COX-2 inhibitors) AND either combined estrogen-progestin oral contraceptive, progestin (e.g. medroxyprogesterone acetate, norethindrone), OR gonadotropin-releasing hormone (GnRH) agonists (e.g. Lupron Depot)
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TADALAFIL

Products Affected

- *tadalafil (pah)*
- TADLIQ

PA Criteria	Criteria Details
Exclusion Criteria	Documentation of concurrent nitrate or Adempas use.
Required Medical Information	Documentation of pulmonary arterial hypertension (PAH) WHO Group I and PAH Functional Class. Reviewer will verify available patient claim history to confirm patient is not using nitrates or Adempas.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or cardiologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For Tadliq: Documentation of trial of, contraindication to, or medical reason for not using tadalafil tablets.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TADALAFIL, BPH

Products Affected

- *tadalafil oral tablet 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of erectile dysfunction
Required Medical Information	Diagnosis of Benign prostatic hyperplasia (BPH) required AND trial of, contraindication to, or medical reason for not using an alpha blocker (e.g. tamsulosin, terazosin).
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TALTZ

Products Affected

- TALTZ SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- TALTZ SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 20 MG/0.25ML, 40 MG/0.5ML, 80 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ankylosing spondylitis: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: Enbrel, an adalimumab product, Cosentyx, or Xeljanz, or 2) If utilized within the past 120 days, approve for continuation of therapy. For non-radiographic axial spondyloarthritis: approve. For psoriasis: Either 1) Trial of, medical reason for not using, or contraindication (e.g., safety concerns, not indicated for patient's age) to 2 of the following therapies: an ustekinumab product, Tremfya, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy. For PsA: Either 1) Trial of, medical reason for not using, or contraindication to 2 of the following therapies: an ustekinumab product, Cosentyx, Tremfya, Xeljanz, Enbrel, or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

TARPEYO

Products Affected

- TARPEYO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: documentation is required that the member has 1) Diagnosis of primary immunoglobulin A nephropathy (IgAN) and 2) at risk of disease progression. Member has proteinuria greater than or equal to 0.5 g/day. For continuation of therapy: documentation that member has been on Tarpeyo for less than 9 months. For reauthorizations: Requests will not be allowed as the safety and efficacy of subsequent courses of Tarpeyo have not been established.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a nephrologist.
Coverage Duration	Request will be authorized for 9 months.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TASIMELTEON

Products Affected

- HETLIOZ LQ
- *tasimelteon*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts of non-24 hour sleep-wake cycle: 1) Member is totally blind with no perception of light, 2) diagnosis of non-24 confirmed by a physiologic circadian phase marker (ex: dim light melatonin onset, assessment of core body temp or measurement of urinary melatonin levels) OR actigraphy with evaluation of sleep logs. For continuation of therapy or reauthorization: documentation of clinical benefit from use of the drug. For night-time sleep disturbances in Smith-Magenis Syndrome (SMS): approve
Age Restrictions	N/A
Prescriber Restrictions	Provider is a sleep specialist or neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TAVNEOS

Products Affected

- TAVNEOS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a rheumatologist or hematologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts: 1) Prescriber attests that Tavneos will be prescribed in combination with corticosteroids AND cyclophosphamide unless there is documented trial of, contraindication to, or medical reason for not using these therapies. 2) Documentation of baseline Birmingham Vasculitis Activity Score (BVAS) score 3) Prescriber attestation that the patient will have liver function tests before treatment (ALT, AST, alkaline phosphate, and total bilirubin) and every 4 weeks after start of therapy for the first 6 months of treatment 4) Prescriber attestation that the patient has been screened for and does not have active hepatitis B virus (HBV) infection at baseline. For continuation of therapy or reauthorization: 1) Documentation of remission (BVAS score of 0) OR improvement in BVAS score 2) Prescriber attestation that patient has no abnormality in liver function tests (abnormality: ALT or AST greater than 3 times the upper limit of normal and bilirubin greater than 2 times the upper limit of normal) 3) Prescriber attestation that patient has no active HBV infection.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

TEPEZZA

Products Affected

- TEPEZZA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of moderate to severe active thyroid eye disease (TED) as evidenced by one or more of the following: a) eyelid retraction greater than or equal to 2 mm, b) moderate or severe soft tissue involvement, c) exophthalmos greater than or equal to 3 mm above normal for race and sex or d) periodic or constant diplopia OR Documentation of chronic TED with one or more of the following: a) a 3 mm or greater increase in exophthalmos from before diagnosis of TED, b) exophthalmos greater than or equal to 3 mm above normal for race and sex.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an ophthalmologist or endocrinologist
Coverage Duration	Request will be authorized for 6 months.
Other Criteria	For active TED: member has had a trial and failure, contraindication to, or medical reason for not using oral or IV glucocorticoids.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TERIPARATIDE

Products Affected

- BONSITY
- *teriparatide subcutaneous solution pen-injector 560 mcg/2.24ml*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation showing patient falls into one of the following categories: a bone mineral density (BMD) value consistent with osteoporosis (i.e., T-scores equal to or less than -2.5) or patient has had an osteoporotic fracture or patient has T-scores from -1.5 to -2.5 at the femoral neck or spine, and a 10-year probability of hip fracture greater than or equal to 3% or a 10-year probability of any major osteoporosis-related fracture greater than or equal to 20% based on the United States-adapted FRAX model.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	In addition, the following criteria is also applicable: 1) Trial of, medical reason for not using, or contraindication to an oral bisphosphonate and a denosumab product.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

THIOLA

Products Affected

- *tiopronin oral*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TIGECYCLINE

Products Affected

- *tigecycline*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	(1) Patient must have documented diagnosis of one of the following infections: (a) complicated skin and skin structure infection, (b) complicated intraabdominal infection, (c) community-acquired pneumonia AND (2) Trial of, contraindication to, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least 2 preferred first-line antibiotics.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an infectious disease specialist.
Coverage Duration	Request will be authorized for 14 days.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TOBI PODHALER

Products Affected

- TOBI PODHALER

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Patient has documented diagnosis of both: 1) cystic fibrosis AND 2) pseudomonas aeruginosa
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOLVAPTAN

Products Affected

- *tolvaptan*
- *tolvaptan (hyponatremia)*

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use with strong CYP3A4 inhibitors (i.e. clarithromycin, ketoconazole, itraconazole, ritonavir, lopinavir-ritonavir, indinavir-ritonavir, indinavir, nelfinavir, saquinavir, nefazodone, conivaptan, and telithromycin).
Required Medical Information	Reviewer will verify available patient claim history to confirm patient is not using a strong CYP3A4 inhibitor (i.e. clarithromycin, ketoconazole, itraconazole, ritonavir, lopinavir-ritonavir, indinavir-ritonavir, indinavir, nelfinavir, saquinavir, nefazodone, conivaptan, and telithromycin).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist, endocrinologist, hepatologist, or nephrologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOPICAL ANTINEOPLASTIC RETINOIDS

Products Affected

- *bexarotene*
- PANRETIN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist, oncologist or specialist for submitted diagnosis
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOPICAL TESTOSTERONE

Products Affected

- *testosterone transdermal gel 1.62 %, 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%)*
- *testosterone transdermal solution*

PA Criteria	Criteria Details
Exclusion Criteria	Patient has history of prostate cancer or breast cancer.
Required Medical Information	New starts of topical testosterone therapy for hypogonadism must have both of the following characteristics of hypogonadism: 1) symptoms associated with hypogonadism (e.g. unexplained mild anemia, low libido, decreased energy, etc.) 2) Two separate instances of low serum total or free testosterone taken in the morning, as defined by the lab reference range.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TRANSDERMAL LIDOCAINE

Products Affected

- *lidocaine external patch 5 %*
- ZTLIDO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documented diagnosis of a medically-accepted indication.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If the request is for the product ZTlido, must provide medical reason for not being able to use generic lidocaine 5% patch
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TREMFYA

Products Affected

- TREMFYA ONE-PRESS SUBCUTANEOUS SOLUTION PEN-INJECTOR
- TREMFYA PEN SUBCUTANEOUS SOLUTION AUTO-INJECTOR 100 MG/ML, 200 MG/2ML
- TREMFYA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 100 MG/ML, 200 MG/2ML
- TREMFYA-CD/UC INDUCTION

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For UC: Approve. Crohn's Disease (CD) - Initial: Trial of, contraindication to, or medical reason (i.e. patient has a diagnosis of severe Crohn's disease) for not using (i.e. severe Crohns disease), or contraindication to 1 of the following: mercaptopurine, azathioprine, methotrexate, sulfasalazine, or corticosteroid (e.g., prednisone, methylprednisolone). Continuation of therapy: patient has been receiving Tremfya for a minimum of 4 months and has a positive clinical response to therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TRIENTINE

Products Affected

- CUVRIOR
- *trientine hcl oral capsule 250 mg*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If the request is for Cuvrior for new starts, member must have trial of, contraindication to, or medical reason for not using trientine hydrochloride.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TRIKAFTA

Products Affected

- TRIKAFTA

PA Criteria	Criteria Details
Exclusion Criteria	Combination use with Kalydeco, Orkambi, or Symdeko.
Required Medical Information	Documentation of CFTR gene that is responsive to elexacaftor-tezacaftor-ivacaftor treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or an expert in the treatment of cystic fibrosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

TYMLOS

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation showing patient falls into one of the following categories: a bone mineral density (BMD) value consistent with osteoporosis (i.e., T-scores equal to or less than -2.5) or patient has had an osteoporotic fracture or patient has T-scores from -1.5 to -2.5 at the femoral neck or spine, and a 10-year probability of hip fracture greater than or equal to 3% or a 10-year probability of any major osteoporosis-related fracture greater than or equal to 20% based on the United States-adapted FRAX model.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	The following criteria is also applicable: 1) trial of, contraindication to, or medical reason for not using an oral bisphosphonate and a denosumab product, and 2) therapy does not exceed 2 years.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TYVASO

Products Affected

- TYVASO DPI MAINTENANCE KIT INHALATION POWDER 112 X 32MCG & 112 X64MCG, 112 X 48MCG & 112 X64MCG, 112 X 48MCG & 112 X80MCG, 16 MCG, 32 MCG, 48 MCG, 64 MCG, 80 MCG
- TYVASO DPI TITRATION KIT INHALATION POWDER 16 & 32 & 48 MCG

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or cardiologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For the treatment of pulmonary arterial hypertension (PAH): 1) documentation of PAH WHO Group I classification and PAH Functional Class and 2) trial of, contraindication to, or medical reason for not using a generic phosphodiesterase inhibitor and a generic endothelin receptor antagonist. For the treatment of pulmonary hypertension associated with interstitial lung disease (PH-ILD, WHO Group 3): documentation of PH-ILD and PAH Functional Class.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

UPTRAVI

Products Affected

- UPTRAVI ORAL
- UPTRAVI TITRATION

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Documentation of pulmonary arterial hypertension (PAH) WHO Group I and PAH Functional Class.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist or cardiologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not using a generic phosphodiesterase inhibitor and a generic endothelin receptor antagonist.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

USTEKINUMAB

Products Affected

- IMULDOSA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML
- SELARSDI INTRAVENOUS
- SELARSDI SUBCUTANEOUS SOLUTION
- SELARSDI SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML
- STARJEMZA INTRAVENOUS
- STARJEMZA SUBCUTANEOUS SOLUTION
- STARJEMZA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML
- STELARA INTRAVENOUS
- STELARA SUBCUTANEOUS SOLUTION 45 MG/0.5ML
- STELARA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML
- STEQEYMA INTRAVENOUS
- STEQEYMA SUBCUTANEOUS SOLUTION
- STEQEYMA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML
- *ustekinumab subcutaneous solution*
- *ustekinumab subcutaneous solution prefilled syringe 45 mg/0.5ml, 90 mg/ml*
- *ustekinumab-aauz subcutaneous solution prefilled syringe 45 mg/0.5ml, 90 mg/ml*
- *ustekinumab-aekn subcutaneous solution prefilled syringe 45 mg/0.5ml, 90 mg/ml*
- YESINTEK INTRAVENOUS
- YESINTEK SUBCUTANEOUS SOLUTION
- YESINTEK SUBCUTANEOUS SOLUTION PREFILLED SYRINGE 45 MG/0.5ML, 90 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.

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

UZEDY

Products Affected

- UZEDY SUBCUTANEOUS SUSPENSION PREFILLED SYRINGE 100 MG/0.28ML, 125 MG/0.35ML, 150 MG/0.42ML, 200 MG/0.56ML, 250 MG/0.7ML, 50 MG/0.14ML, 75 MG/0.21ML

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented history of receiving oral risperidone without any clinically significant side effects.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For schizophrenia: Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least two of the following: Abilify Maintena, Abilify Asimtufii, Invega Sustenna, Invega Trinza, or Invega Hafyera. For treatment of bipolar I disorder as monotherapy: Trial and failure of, contraindication, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using at least one of the following: Abilify Maintena or Abilify Asimtufii. For treatment as adjunctive therapy to lithium or valproate for the maintenance treatment of bipolar I disorder: approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VALCHLOR

Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or dermatologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Trial of, contraindication to, or medical reason for not being able to use one of the following: a topical corticosteroids or a topical retinoids.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VEMLIDY

Products Affected

- VEMLIDY

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts: attestation that member has been tested for HIV infection. If member is HIV-positive, Vemlidy is not used alone.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

VEOZAH

Products Affected

- VEOZAH

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Initial: (1) Documented diagnosis of moderate to severe vasomotor symptoms due to menopause AND (2) Trial of, contraindication to, or medical reason (e.g. intolerance, hypersensitivity or contraindication) for not using a hormonal therapy (e.g., estradiol, oral Premarin, Prempro). Reauthorization: (1) Documentation of positive clinical response to therapy (e.g., decrease in frequency or severity of vasomotor symptoms from baseline)
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VIGABATRIN

Products Affected

- *vigabatrin*
- VIGAFYDE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For infantile spasms or West syndrome, the request will be approved. For diagnosis of refractory complex partial seizures: 1) documentation of diagnosis, and 2) attestation the member is currently receiving another antiepileptic drug, and 3) attestation the member has experienced treatment failure from two alternative antiepileptic agents.
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VIJOICE

Products Affected

- VIJOICE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts, all of the following must be included: 1) Documentation of genetic testing confirming diagnosis AND 2) Member has at least one target lesion identified on imaging AND 3) Prescriber attests the patient's condition is severe or life-threatening and necessitates systemic treatment. For continuation of therapy or reauthorization, attestation of a positive clinical response (i.e. reduction in the sum of measurable target lesion volume, absence of progression of non-target lesions, absence of any new lesions, etc.).
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a geneticist, dermatologist, vascular surgeon, hematologist/oncologist, or other specialist in the treatment of PIK3CA-Related Overgrowth Spectrum(PROS).
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

VMAT-2 INHIBITORS

Products Affected

- AUSTEDO
- AUSTEDO XR
- AUSTEDO XR PATIENT TITRATION ORAL TABLET EXTENDED RELEASE THERAPY PACK 12 & 18 & 24 & 30 MG
- INGREZZA ORAL CAPSULE
- INGREZZA ORAL CAPSULE SPRINKLE
- INGREZZA ORAL CAPSULE THERAPY PACK
- *tetrabenazine*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist, clinical geneticist, or psychiatrist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	If the request is for tetrabenazine, Ingrezza or Ingrezza Sprinkle, request will be approved. If the request is for Austedo or Austedo XR, the member must have trial of or medical reason for not using tetrabenazine. Reauthorization: Confirmation of improvement in tardive dyskinesia symptoms or chorea associated with Huntington disease symptoms.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VORICONAZOLE

Products Affected

- *voriconazole intravenous*

PA Criteria	Criteria Details
Exclusion Criteria	Non-Part D indications.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 6 months.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

VOSEVI

Products Affected

- VOSEVI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Detectable HCV RNA viral load prior to treatment within 6 months of request. In addition, documentation of treatment history, and if cirrhotic, documentation of compensated or decompensated cirrhosis.
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized for 12 weeks as per AASLD-IDSA guidance.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

VOWST

Products Affected

- VOWST

PA Criteria	Criteria Details
Exclusion Criteria	Treatment of Clostridioides difficile infection (CDI)
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	If all the criteria are met, the request will be approved for 1 month
Other Criteria	Diagnosis of at least 1 recurrent episode of CDI
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

WEGOVY

Products Affected

- WEGOVY SUBCUTANEOUS SOLUTION AUTO-INJECTOR 0.25 MG/0.5ML, 0.5 MG/0.5ML, 1 MG/0.5ML, 1.7 MG/0.75ML, 2.4 MG/0.75ML

PA Criteria	Criteria Details
Exclusion Criteria	The member has an indication of only weight reduction or maintenance for overweight or obesity. The member has concurrent use of any GLP-1 receptor agonist. The member has a personal history of medullary thyroid carcinoma. The member has Multiple Endocrine Neoplasia syndrome type 2.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Reduction of the risk of major adverse cardiovascular events in adults with established cardiovascular disease and either obesity or overweight- For new starts: The member has an indication for reducing the risk of adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke) in adults with established cardiovascular disease. Documentation demonstrates patient has established cardiovascular disease (e.g., prior myocardial infarction, prior stroke, symptomatic peripheral arterial disease). Documentation is provided that the patient is overweight or obese (defined as a BMI of greater than or equal to 27 kg/m2). Treatment of noncirrhotic metabolic dysfunction-associated steatohepatitis (MASH) - For new starts: The member has a documented diagnosis of MASH with documentation of moderate to advanced liver fibrosis (fibrosis stage 2 or 3) as evidenced by liver biopsy, biomarker testing (AST to platelet ratio index score, fibrosis 4 score, NAFLD fibrosis score, etc.), transient elastography or ultrasonography. For continuation of therapy or reauthorization: Documentation is provided that the patient has a positive clinical response to treatment.

PA Criteria	Criteria Details
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

WEGOVY TABLETS

Products Affected

- WEGOVY ORAL

PA Criteria	Criteria Details
Exclusion Criteria	The member has an indication of only weight reduction or maintenance for overweight or obesity. The member has concurrent use of any GLP-1 receptor agonist. The member has a personal history of medullary thyroid carcinoma. The member has Multiple Endocrine Neoplasia syndrome type 2.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Reduction of the risk of major adverse cardiovascular events in adults with established cardiovascular disease and either obesity or overweight - For new starts: The member has an indication for reducing the risk of adverse cardiovascular events (cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke) in adults with established cardiovascular disease. Documentation demonstrates patient has established cardiovascular disease (i.e., prior myocardial infarction, prior stroke, symptomatic peripheral arterial disease). Documentation is provided that the patient is overweight or obese (defined as a BMI of greater than or equal to 27 kg/m2). The member must have documentation or a medical reason for not being able to use the Wegovy injection. For continuation of therapy or reauthorization: Documentation is provided that the patient has a positive clinical response to treatment.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

WHITE BLOOD CELL STIMULATORS

Products Affected

- FULPHILA
- FYLNETRA
- LEUKINE INJECTION SOLUTION RECONSTITUTED
- NEULASTA ONPRO SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- NEULASTA SUBCUTANEOUS SOLUTION 4 MG/0.4ML
- NEULASTA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- ZARXIO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	For new starts for Neulasta and Fylnetra: documentation of trial of, contraindication to, or medical reason for not using Fulphila. Continuation of therapy or re-authorization criteria: diagnosis of chronic neutropenia or a medical reason for continued need for GCSF.
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	New starts will be authorized for 4 months. Cont of therapy or reauth until end of contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

WINREVAIR

Products Affected

- WINREVAIR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	Pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1)- Initial: [Note: documentation required] (1) PAH was confirmed by right heart catheterization AND (2) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg AND (3) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND (4) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. Continuation of therapy: patient has positive clinical response to treatment.
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Documented trial and failure of, or contraindication to combination therapy including one PDE-5 inhibitor AND one endothelin receptor antagonist. Documentation of platelet count of greater than 50,000/mm3.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

XDEMVY

Products Affected

- XDEMVY

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	N/A
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

XELJANZ

Products Affected

- *tofacitinib citrate er*
- *tofacitinib citrate oral solution*
- *tofacitinib citrate oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For ankylosing spondylitis: Trial of, medical reason for not using, or contraindication to naproxen and 1 of the following: an adalimumab product, Enbrel, or Cosentyx. For pJIA: Trial of, medical reason for not using, or contraindication to 1 of the following DMARDs: methotrexate or leflunomide and 1 of the following: an adalimumab product or Enbrel. For PsA: Trial of, medical reason for not using, or contraindication to 1 of the following: Enbrel, an adalimumab product, or Cosentyx. For RA: Trial of, medical reason for not using, or contraindication to 1 disease modifying antirheumatic drug (DMARD) (methotrexate, leflunomide, or sulfasalazine) and 1 of the following: Enbrel or an adalimumab product. For UC: Approve. Continuation of therapy: patient has been receiving tofacitinib for a minimum of 4 months and has a positive clinical response.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

XERMELO

Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a gastroenterologist or an oncologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: 1) Attestation that diarrhea is inadequately controlled by stable dose of SSA therapy for at least three months. For continuation of therapy or reauthorization: 1) documentation of positive clinical response to xermelo and 2) Attestation to continue to be used in combination with SSA.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

XIFAXAN

Products Affected

- XIFAXAN

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	For HE: gastroenterologist or hepatologist. For IBS-D: gastroenterologist.
Coverage Duration	For HE: contract year. For IBSD: 14 days (cannot exceed 3 courses of 14 days each). For TD: 3 days.
Other Criteria	For diagnosis of hepatic encephalopathy (HE): trial of, contraindication to, or medical reason for not using lactulose. For diagnosis of irritable bowel syndrome with diarrhea (IBSD): No more than 3 courses of 14 days each. For travelers diarrhea (TD) caused by noninvasive strains of E. Coli (with no bloody stools or fever): patient must be intolerant to or must have had a trial of at least 3 days of one of the following agents: ciprofloxacin, ofloxacin, levofloxacin or azithromycin.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

XOLAIR

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a pulmonologist, allergist, immunologist, dermatologist, or otolaryngologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	New starts for moderate to severe persistent allergic asthma: 1) Evidence of specific allergic sensitivity confirmed by positive skin test (i.e. prick/puncture test) or blood test (i.e. radioallergosorbent test) for a specific IgE or in vitro reactivity to a perennial aeroallergen, AND 2) Pretreatment serum IgE levels greater than 30 IU/mL, AND 3) Symptoms are not adequately controlled with high-dose inhaled corticosteroid (ICS) plus additional controller medication (ie. long-acting B2 agonist) for at least 3 months, or there is a medical reason for not using these drugs. Continuation of therapy or reauthorization criteria for moderate to severe persistent allergic asthma: 1) Reduction in asthma exacerbation resulting in systemic steroid use and/or hospitalization, OR 2) Reduction of rescue inhaler use, OR 3) Documentation of improvement in pulmonary function tests since baseline (prior to initiation of Xolair). New starts for chronic idiopathic urticaria: 1) inadequate symptomatic relief despite trial of two weeks of at least one 2nd generation antihistamine (unless contraindicated), AND 2) disease must be severe enough to warrant short term systemic corticosteroid therapy for management of urticaria. Continuation of therapy or reauthorization criteria for chronic idiopathic urticaria: 1) improvement from baseline of symptoms associated with urticaria within 6 months of Xolair use. New starts for nasal polyps: 1) currently using an intranasal corticosteroid, will be prescribed an intranasal corticosteroid with request, or has a medical reason for not using an intranasal corticosteroid.

PA Criteria	Criteria Details
	Continuation of therapy or reauthorization criteria for nasal polyps: 1) Documentation has been provided that demonstrates a clinical benefit (e.g. improvements in symptom severity, nasal polyp score [NPS], sino-nasal outcome test-22 [SNOT-22], nasal congestion score [NCS]) AND 2) continued use of intranasal corticosteroid, or has a medical reason for not using one. New starts for food allergy: 1) diagnosis of IgE-mediated food allergy 2) Xolair will be used in conjunction with food allergen avoidance. Continuation of therapy or reauthorization criteria for food allergy: 1) Documentation of clinical benefit
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

XOLREMDI

Products Affected

- XOLREMDI

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be an immunologist or a hematologist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: 1) A documented diagnosis of WHIM (warts, hypogammaglobulinemia, infections and myelokathexis) syndrome confirmed by genotype variant of chemokine receptor 4 (CXCR4) and absolute neutrophil count (ANC) of less than or equal to 400 cells/microliter or white blood cells (WBC) less than or equal to 400 cells/microliter and 2) Documentation of baseline ANC and absolute lymphocyte count (ALC). For renewal 1) Documentation or provider attestation of positive clinical response (i.e. improvement from baseline in ANC, WBC and/or ALC or reduced frequency, duration, or severity of infections, fewer warts, or improved or stabilized clinical signs and/or symptoms of WHIM).
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

XYWAV

Products Affected

- XYWAV

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a sleep specialist, pulmonologist or a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For treatment of somnolence associated with narcolepsy, patient must have documentation of either trial of or a medical reason for being unable to use a CNS stimulant (e.g. methylphenidate, modafinil, armodafinil, etc.). For the treatment of cataplexy associated with narcolepsy or idiopathic hypersomnia, approve.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

YORVIPATH

Products Affected

- YORVIPATH

PA Criteria	Criteria Details
Exclusion Criteria	Diagnosis of acute post-surgical hypoparathyroidism.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with an endocrinologist.
Coverage Duration	New starts will be authorized for 6 months. Cont of therapy or reauth until end of contract year.
Other Criteria	For new starts: 1) Documented diagnosis of chronic hypoparathyroidism AND 2) Provider attestation that patient is currently receiving or has medical reason for not receiving calcium supplementation and active vitamin D treatment AND 3) An albumin-corrected serum calcium level of 7.8 mg/dL or greater. For reauthorization: Documentation of improvement in albumin-corrected serum calcium from baseline.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

YUTREPIA

Products Affected

- YUTREPIA

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a cardiologist or a pulmonologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For the treatment of pulmonary arterial hypertension (PAH): 1) documentation of PAH WHO Group I classification and PAH Functional Class and 2) trial of, contraindication to, or medical reason for not using a generic phosphodiesterase inhibitor and a generic endothelin receptor antagonist. For the treatment of pulmonary hypertension associated with interstitial lung disease (PH-ILD, WHO Group 3): documentation of PHILD and PAH Functional Class.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ZEPBOUND

Products Affected

- ZEPBOUND KWIKPEN
- ZEPBOUND SUBCUTANEOUS SOLUTION AUTO-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	The member has an indication of only weight reduction or maintenance for overweight or obesity. The member has concurrent use of any GLP-1 receptor agonist. The member has a personal history of medullary thyroid carcinoma. The member has Multiple Endocrine Neoplasia syndrome type 2.
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a sleep disorder specialist, pulmonologist, ENT, or other provider specializing in obstructive sleep apnea.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For new starts: The member has an indication for moderate to severe obstructive sleep apnea (OSA) in adults with obesity. Documentation of diagnosis of OSA through polysomnography (sleep study) with an apnea-hypopnea index of 15 or more events per hour, or five or more events per hour in the presence of symptoms (e.g., cognitive impairment, fatigue, insomnia, loud snoring) or cardiovascular comorbidities (e.g., hypertension, ischemic heart disease, previous stroke). Documentation is provided that the patient is obese (defined as a BMI of greater than or equal to 30 kg/m2). For continuation of therapy: Documentation of positive response to treatment. Documentation member has achieved and/or maintained a decrease in weight since baseline.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

ZEPOSIA

Products Affected

- ZEPOSIA
- ZEPOSIA 7-DAY STARTER PACK
- ZEPOSIA STARTER KIT ORAL CAPSULE THERAPY PACK 0.23MG &0.46MG 0.92MG(21)

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Specialist for submitted diagnosis.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	For multiple sclerosis: Trial of, contraindication to, or medical reason for not using two of the following: dalfampridine ER, dimethyl fumarate, fingolimod, glatiramer, glatopa, or teriflunomide. For ulcerative colitis: Either 1) Trial of, medical reason for not using, or contraindication to 1 of the following: an ustekinumab product or an adalimumab product or 2) If utilized within the past 120 days, approve for continuation of therapy.
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ZILBRYSQ

Products Affected

- ZILBRYSQ

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescriber must be a neurologist, rheumatologist, or other appropriate specialist
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	Patient has tried and failed, a medical reason for not using, or has a contraindication to two (2) or more conventional therapies (i.e. pyridostigmine, corticosteroids, or non-steroidal immunosuppressive therapies)
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ZTALMY

Products Affected

- ZTALMY

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	N/A
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a neurologist.
Coverage Duration	Request will be authorized until the end of the contract year.
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

ZURZUVAE

Products Affected

- ZURZUVAE

PA Criteria	Criteria Details
Exclusion Criteria	N/A
Required Medical Information	The member has a documented diagnosis of postpartum depression
Age Restrictions	N/A
Prescriber Restrictions	Prescribed by or in consultation with a psychiatrist or obstetrician/gynecologist
Coverage Duration	Request will be authorized until the end of the contract year
Other Criteria	N/A
Indications	All Medically-accepted Indications.
Off-Label Uses	N/A
Part B Prerequisite	No
Prerequisite Therapy Required	No

PART B VERSUS PART D

Products Affected

- *acetylcysteine inhalation solution 10 %, 20 %*
- *acyclovir sodium intravenous solution 50 mg/ml*
- *albuterol sulfate inhalation nebulization solution (2.5 mg/3ml) 0.083%, (5 mg/ml) 0.5%, 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 mg*
- *aprepitant oral capsule therapy pack 80 & 125 mg*
- ASTAGRAF XL ORAL CAPSULE EXTENDED RELEASE 24 HOUR 0.5 MG, 1 MG, 5 MG
- *azathioprine oral tablet 50 mg*
- *budesonide inhalation suspension 0.25 mg/2ml, 0.5 mg/2ml, 1 mg/2ml*
- *clinisol sf intravenous solution 15 %*
- *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*
- *dronabinol oral capsule 10 mg, 2.5 mg, 5 mg*
- EMEND ORAL SUSPENSION RECONSTITUTED 125 MG/5ML
- ENGERIX-B INJECTION SUSPENSION 20 MCG/ML
- ENGERIX-B INJECTION SUSPENSION PREFILLED SYRINGE 10 MCG/0.5ML, 20 MCG/ML
- ENVARSUS 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*
- *formoterol fumarate inhalation nebulization solution 20 mcg/2ml*
- GAMMAGARD ERC INJECTION SOLUTION 10 GM/100ML, 5 GM/50ML
- GAMMAGARD INJECTION SOLUTION 1 GM/10ML, 10 GM/100ML, 2.5 GM/25ML, 20 GM/200ML, 30 GM/300ML, 5 GM/50ML
- GAMMAGARD S/D LESS IGA INTRAVENOUS SOLUTION RECONSTITUTED 10 GM, 5 GM
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- GAMUNEX-C INJECTION SOLUTION 1 GM/10ML, 10 GM/100ML, 2.5 GM/25ML, 20 GM/200ML, 40 GM/400ML, 5 GM/50ML
- *gengraf oral capsule 100 mg, 25 mg*
- *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*
- *levalbuterol hcl inhalation nebulization solution 0.31 mg/3ml, 0.63 mg/3ml, 1.25 mg/3ml*
- *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 %
- OCTAGAM INTRAVENOUS SOLUTION 1 GM/20ML, 10 GM/100ML, 10 GM/200ML, 2 GM/20ML, 2.5 GM/50ML, 20 GM/200ML, 30 GM/300ML, 5 GM/100ML, 5 GM/50ML
- *ondansetron hcl oral solution 4 mg/5ml*
- *ondansetron hcl oral tablet 24 mg, 4 mg, 8 mg*
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- *pentamidine isethionate inhalation solution reconstituted 300 mg*
- *plenamine intravenous solution 15 %*
- PRIVIGEN INTRAVENOUS SOLUTION 10 GM/100ML, 20 GM/200ML, 40 GM/400ML, 5 GM/50ML
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- PROGRAF ORAL PACKET 0.2 MG, 1 MG
- PULMOZYME INHALATION SOLUTION 2.5 MG/2.5ML
- RABAVERT INTRAMUSCULAR SUSPENSION RECONSTITUTED
- 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*
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- *tacrolimus er oral capsule extended release 24 hour 0.5 mg, 1 mg, 5 mg*
- *tacrolimus intravenous solution 5 mg/ml*
- *tacrolimus oral capsule 0.5 mg, 1 mg, 5 mg*
- TENIVAC INTRAMUSCULAR INJECTABLE 5-2 LFU (INJECTION)
- TENIVAC INTRAMUSCULAR SUSPENSION 5-2 LF/0.5ML
- *tobramycin inhalation nebulization solution 300 mg/4ml, 300 mg/5ml*

Details

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

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