

PA Criteria

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

5HT-3 ANTAGONISTS

GRANISETRON HCL, ONDANSETRON HCL, ONDANSETRON ODT

All FDA approved indications not otherwise excluded from Part D

A. The patient must be receiving moderate to highly emetogenic chemotherapy, radiation therapy, or post-operative treatment. B. OR the patient is NOT being treated for chemotherapy, radiation therapy, or post-operative treatment AND the patient had a previous trial or contraindication to BOTH promethazine AND prochlorperazine C. Brand name will only be approved with failure on ALL available generic formulations.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

6 months

B vs D coverage determination per CMS guidelines

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

ACNE

AVITA, RETIN-A MICRO, TRETINOIN

All FDA approved indications not otherwise excluded from Part D

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

ACTEMRA

ACTEMRA

All FDA approved indications not otherwise excluded from Part D

Active infection (including TB). Concurrent therapy with other biologic agent(s).

Screening for latent tuberculosis is required. If results are positive, patient must have completed treatment or must currently be receiving treatment for latent tuberculosis.

Evaluate for HBV risk and initiate treatment if appropriate.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Must have an inadequate response to at least one nonbiologic DMARD or intolerance/contraindication to at least 2 nonbiologic DMARDs. Must have an inadequate response or intolerance/contraindication to one TNF antagonist. For renewals, patient must have responded to Actemra therapy (e.g., condition improved or stabilized).

Prior Authorization Group

Drug Names

ACTIMMUNE

ACTIMMUNE

Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Hypersensitivity to E.coli-derived products and/or interferon gamma.
Required Medical Information	Patient has no history of myelosuppression, complete blood count within normal limits, platelet count within normal limits, liver function tests within normal limits. Monitoring of complete blood count, platelet count, and liver function tests every 3 months is required.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	
Prior Authorization Group	ADAGEN
Drug Names	ADAGEN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Severe thrombocytopenia. Use in preparation for or in support of bone marrow transplantation.
Required Medical Information	Bone marrow transplantation failure or patient is not a suitable candidate for bone marrow transplantation.
Age Restrictions	
Prescriber Restrictions	Endocrinologist, ID specialist, Allergist, Immunologist, Clinical or Biochemical Geneticist, Hematologist
Coverage Duration	Plan Year
Other Criteria	Use for direct replacement for deficient enzyme (no benefit achieved in patients with immunodeficiency due to other causes).
Prior Authorization Group	ADCIRCA
Drug Names	ADCIRCA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Nitrate therapy
Required Medical Information	Diagnosis of pulmonary arterial hypertension (PAH), (WHO Group 1). PAH been confirmed by right heart catheterization. If patient is an infant, PAH diagnosed by Doppler echocardiogram.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	AFINITOR
Drug Names	AFINITOR
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	

Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	ALDURAZYME
Drug Names	ALDURAZYME
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Diagnosis confirmed by diagnostic method, enzymatic assay or DNA testing.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	For Scheie syndrome: must have at least 2 moderate to severe symptoms. Must demonstrate improvement in lung function in patients who has previously received at least 26 weeks of Aldurazyme on re-authorization.
Prior Authorization Group	ALPHA1-PROTEINASE INHIBITOR
Drug Names	ARALAST NP
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Patient has IgA deficiency with antibodies against IgA.
Required Medical Information	Alpha1-proteinase inhibitor concentration is less than 11 micromoles per liter. The FEV1 level is between 35% and 60% predicted OR greater than 60% predicted. If the FEV1 is greater than 60% predicted, then the patient has experienced a rapid decline in lung function that warrants treatment.
Age Restrictions	18 years old and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	AMPHETAMINES
Drug Names	AMPHETAMINE/DEXTROAMPHETA, DEXTROAMPHETAMINE SULFATE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	MAOI concurrent use or within the last 14 days except if prescriber is a psychiatrist with experience prescribing both MAOI and amphetamine/dextroamphetamine drugs
Required Medical Information	Sleep studies for narcolepsy diagnosis
Age Restrictions	3 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Consider benefits of use versus the potential risks of serious cardiovascular events
Prior Authorization Group	AMPYRA
Drug Names	AMPYRA
Covered Uses	All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria	Moderate to severe renal impairment, history of seizures, Ampyra at doses exceeding 10 mg twice daily.
Required Medical Information	Patient must be able to walk 25 feet with or without assistance.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	2 months, then plan year upon renewal
Other Criteria	Patient must demonstrate sustained walking impairment prior to starting Ampyra. To continue therapy, the patient must experience an improvement in walking speed or other objective measure of walking ability since starting Ampyra.
Prior Authorization Group	ANABOLIC STEROIDS
Drug Names	OXANDROLONE
Covered Uses	All FDA approved indications not otherwise excluded from Part D, HIV-wasting syndrome
Exclusion Criteria	Known or suspected carcinoma of the prostate or breast (in male patients), carcinoma of the breast in women with hypercalcemia, pregnancy, nephrosis (the nephrotic phase of nephritis), hypercalcemia.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
Prior Authorization Group	ANAGRELIDE
Drug Names	ANAGRELIDE HYDROCHLORIDE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Severe hepatic impairment.

Required Medical Information

A. If the diagnosis is chronic myelogenous leukemia: a. persistent granulocyte count greater than or equal to 50,000/mcL without infection b. absolute basophil count greater than or equal to 100/mcL c. evidence of hyperplasia of the granulocytic line in the bone marrow d. presence of the Philadelphia chromosome e. leukocyte alkaline phosphatase less than or equal to lower limit of the lab range. B. If the diagnosis is polycythemia vera (either all three major criteria or first two major criteria and two minor criteria): major criteria: increase red cell mass (in men, greater than or equal to 36 mL/kg and in women, greater than or equal to 32 mL/kg), normal arterial oxygen saturation (greater than or equal to 92%), splenomegaly. minor criteria: platelet count greater than or equal to 400,000/mcL without iron deficiency or bleeding, white blood cell count greater than or equal to 12,000/mcL without infection, leukocyte alkaline phosphatase greater than or equal to 100 mcL, serum B12 greater than 900 pcg/mL. C. If the diagnosis is thrombocytosis: a. platelet count greater than or equal to 900,000/mcL b. profound megakaryocytic hyperplasia in bone marrow c. absence of Philadelphia chromosome d. normal red cell mass c. normal serum iron and ferritin and normal marrow iron stores d. pre-treatment cardiovascular examination.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Oncologist or Hematologist

6 months

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

ARANESP

ARANESP ALBUMIN FREE

All FDA approved indications not otherwise excluded from Part D

Uncontrolled hypertension, use in patients with nonmetastatic or curable cancer, use in myeloid cancer, hemoglobin at or exceeding 13 g/dL

Required Medical Information

Patients with low iron stores require concomitant iron supplementation, pretreatment hemoglobin level less than 10 g/dL (or less than 11 g/dL with clinical symptoms of anemia). Cancer patients with anemia must be currently receiving myelosuppressive chemotherapy. Patients with myelodysplastic syndrome (MDS) may receive drug for symptomatic anemia provided the MDS is not associated with del(5q) cytogenetic abnormality and serum EPO is less than or equal to 500 mU/mL. Once on therapy, the patient must show an objective clinical response to treatment (ie, rise in hemoglobin or hematocrit from baseline). If hemoglobin increases significantly (eg, more than 1 g/dL in any 2 week period) or exceeds 12 g/dL, the prescriber must reduce dose. Patients must report any signs or symptoms of cardiovascular or thrombotic adverse events to the prescriber.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

12 weeks

Prior Authorization Group	ARCALYST
Drug Names	ARCALYST
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Active or chronic infection. Concurrent therapy with other biologics.
Required Medical Information	
Age Restrictions	12 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	ATYPICAL ODT
Drug Names	FAZACLO
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	For Fazaclo: a. if the patient has any of the following contraindications: agranulocytosis, bone marrow suppression, coma, ileus, leukopenia, myocarditis or neutropenia b. if the patient has CNS depression, dementia-related psychosis or uncontrolled epilepsy.
Required Medical Information	The patient must be unable/unwilling to take tablets or capsules or are high risk for non-compliance AND must not be receiving other tablets or capsules indicating that they can take non-dissolvable tablets.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	AVONEX
Drug Names	AVONEX
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	B VS. D

Drug Names

ACETYLCYSTEINE, ADRIAMYCIN, ALBUTEROL SULFATE, ALIMTA, AMIFOSTINE, AMINOSYN, AMINOSYN 7%/ELECTROLYTES, AMINOSYN 8.5%/ELECTROLYTE, AMINOSYN II, AMINOSYN II 3.5%/DEXTROSE, AMINOSYN II 3.5/DEXTROSE, AMINOSYN II 4.25/DEXTROSE, AMINOSYN II 5/DEXTROSE 25, AMINOSYN II 8.5%/ELECTROL, AMINOSYN II M 3.5%/DEXTRO, AMINOSYN M, AMINOSYN-HBC, AMINOSYN-HF, AMINOSYN-PF, AMINOSYN-PF 7%, AMIODARONE HCL, AMPHOTERICIN B, ASTRAMORPH, AVASTIN, AZASAN, AZATHIOPRINE, AZATHIOPRINE SODIUM, BICNU, BLEOMYCIN SULFATE, BONIVA, BUDESONIDE, BUSULFEX, CALCITRIOL, CAMPATH, CARBOPLATIN, CELLCEPT, CISPLATIN, CLADRIBINE, CLINIMIX 2.75%/DEXTROSE 5, CLINIMIX 4.25%/DEXTROSE 1, CLINIMIX 4.25%/DEXTROSE 2, CLINIMIX 4.25%/DEXTROSE 5, CLINIMIX 5%/DEXTROSE 15%, CLINIMIX 5%/DEXTROSE 20%, CLINIMIX 5%/DEXTROSE 25%, CLINIMIX E 2.75%/DEXTROSE, CLINIMIX E 4.25%/DEXTROSE, CLINIMIX E 5%/DEXTROSE 15, CLINIMIX E 5%/DEXTROSE 20, CLINIMIX E 5%/DEXTROSE 25, CLINISOL SF 15%, COLISTIMETHATE SODIUM, COSMEGEN, CROMOLYN SODIUM, CUBICIN, CYCLOPHOSPHAMIDE, CYCLOSPORINE, CYCLOSPORINE MODIFIED, CYTARABINE, DACARBAZINE, DAUNORUBICIN HCL, DAUNOXOME, DECAVAC, DEPO-PROVERA, DEXRAZOXANE, DIPHTHERIA/TETANUS TOXOID, DOXIL, DOXORUBICIN HCL, DURAMORPH, ELITEK, ELSPAR, ENGERIX-B, EPIRUBICIN HCL, ETOPOSIDE, FASLODEX, FENTANYL CITRATE, FLUDARABINE PHOSPHATE, FLUOROURACIL, FREAMINE HBC 6.9%, FREAMINE III, FREAMINE III 3%, GAMASTAN S/D, GANCICLOVIR, GEMCITABINE HCL, GENGRAF, GRANISETRON HCL, HECTOROL, HEPARIN SODIUM, HEPARIN SODIUM/D5W, HEPARIN SODIUM/NACL 0.45%, HEPARIN SODIUM/SODIUM CHL, HEPATAMINE, HEPATASOL, HERCEPTIN, HYCAMTIN, HYDROMORPHONE HCL, IDARUBICIN HCL, IFEX, IFOSFAMIDE, IFOSFAMIDE/MESNA, INTRALIPID, INTRON-A, INTRON-A W/DILUENT, IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE/ALBUT, IRINOTECAN, ISTODAX, LEUCOVORIN CALCIUM, LEVALBUTEROL, LEVOCARNITINE, LIDOCAINE/PRILOCAINE, LIPOSYN II, LIPOSYN III, MELPHALAN HYDROCHLORIDE, MESNA, METHOTREXATE SODIUM, MIACALCIN, MITOMYCIN, MITOXANTRONE HCL, MORPHINE SULFATE, MUSTARGEN, MYCOPHENOLATE MOFETIL, MYFORTIC, NEORAL, NEPHRAMINE, NOVAMINE, NULOJIX, ONCASPAR, ONDANSETRON HCL, ONTAK, OXALIPLATIN, PACLITAXEL, PENTOSTATIN, PHOTOFRIN, PREMASOL, PROCALAMINE, PROGRAF, PROLEUKIN, PROSOL, RAPAMUNE, RECOMBIVAX HB, RENAMIN, SANDIMMUNE, TACROLIMUS, TAXOTERE, TETANUS TOXOID ADSORBED, TETANUS/DIPHTHERIA TOXOID, TOBI, TOPOSAR, TOPOTECAN HCL, TPN ELECTROLYTES, TRAVASOL, TREANDA, TRELSTAR DEPOT MIXJECT, TRELSTAR LA MIXJECT, TRELSTAR MIXJECT, TRISENOX, TROPHAMINE,

VANCOMYCIN HCL, VELCADE, VIDAZA, VINBLASTINE SULFATE, VINCASAR PFS, VINCRISTINE SULFATE, VINOURELBINE TARTRATE, XOPENEX, ZEMPLAR, ZORTRESS

Covered Uses

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.

Exclusion Criteria

Required Medical Information

Age Restrictions

Prescriber Restrictions

Coverage Duration

NA

Other Criteria

Prior Authorization Group

BANZEL

Drug Names

BANZEL

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

The patient is diagnosed with familial short QT Syndrome.

Required Medical Information

The patient has seizures associated with Lennox-Gastaut Syndrome.

Age Restrictions

Prescriber Restrictions

Neurologist or affiliated with neurology practice

Coverage Duration

Plan Year

Other Criteria

Prior Authorization Group

BETASERON

Drug Names

BETASERON

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Albumin hypersensitivity, concurrent use of any of the following medications: Interferon-beta therapy (Avonex, Extavia, or Rebif), glatiramer acetate, or mitoxantrone

Required Medical Information

MRI has been performed and has features suggestive of MS (evidence of lesion)

Age Restrictions

Prescriber Restrictions

Neurologist

Coverage Duration

Plan Year

Other Criteria

Patients with previous use (12 or more months) of Betaseron must demonstrate 1 of the following clinical responses: decrease in the frequency of relapses, slowing of disease progression, MRI lesions have diminished with therapy, OR patient is stable on therapy.

Prior Authorization Group

BUPRENORPHINE

Drug Names

BUPRENORPHINE HCL, SUBOXONE

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Required Medical Information

Age Restrictions

16 years of age and older

Prescriber Restrictions	Prescribers must be registered with the Substance Abuse and Mental Health Services Administration
Coverage Duration	Buprenorphine - one month (40 weeks if pregnant). Buprenorphine-naloxone - 12 months.
Other Criteria	Buprenorphine and buprenorphine-naloxone should be part of an overall treatment program. The patient should be monitored periodically.
Prior Authorization Group	BYETTA
Drug Names	BYETTA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	A. The patient is diagnosed as having type-2 diabetes with an HbA1c level greater than 7. B. The patient has a creatinine clearance of greater than 30mL/minute or normal kidney function. C. The patient has had an inadequate treatment response, intolerance or contraindication to metformin or a sulfonyleurea medication. D. If the patient has received previous Byetta therapy for at least 3 months, the patient demonstrated a reduction in HbA1c since initiating Byetta therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	CAMPRAL
Drug Names	CAMPRAL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Renal failure.
Required Medical Information	A. Clinical diagnosis for alcohol dependence. B. AND clinical evidence indicated that the patient will be abstinent at least 5 days prior to treatment initiation. C. AND a trial of naltrexone (oral/injectable) has been attempted, at clinically significant dosage and duration. Or therapy is documented to be clinically inappropriate (hepatic insufficiency, chronic pain medication use). D. AND medication administration should be part of a comprehensive psychosocial treatment program.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
Prior Authorization Group	CAYSTON
Drug Names	CAYSTON
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	

Required Medical Information Diagnosis of cystic fibrosis is confirmed by appropriate diagnostic or genetic testing. Confirmation of P. aeruginosa in cultures of the airways. Upon renewal, patients 6 years of age or older who have diminished pulmonary function tests by greater than 10% while receiving Cayston therapy should have a clinical reason to continue Cayston therapy. Upon renewal, patients younger than 6 years of age should have a clinical reason to continue Cayston therapy.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

CELEBREX

CELEBREX

All FDA approved indications not otherwise excluded from Part D

Post-operative pain following CABG surgery.

6 months for FAP and JRA, 12 months for dysmenorrhea, OA, RA, AS, 1 month for acute pain

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

CEREZYME

CEREZYME

All FDA approved indications not otherwise excluded from Part D

Concurrent therapy with Zavesca.

Diagnosis confirmed by bone marrow histology, DNA testing or measurement of b-glucocerebrosidase enzyme activity less than 30 percent. Must have at least one of following conditions: anemia, thrombocytopenia, bone disease, hepatomegaly or splenomegaly. Must demonstrate a decrease in liver and spleen volume and/or increase in platelet count and/or increase in Hgb concentration in patients who has previously received 24 months of Cerezyme therapy.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

Age Restrictions

CHANTIX

CHANTIX

All FDA approved indications not otherwise excluded from Part D

Concurrent Zyban use

Prescriber Restrictions	
Coverage Duration	12 weeks initial, 12 weeks additional upon renewal
Other Criteria	
Prior Authorization Group	CIMZIA
Drug Names	CIMZIA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Active infection (including TB).
Required Medical Information	Screening for latent TB infection and assessment for Hep B risk and be treated if positive.
Age Restrictions	18 years of age and older
Prescriber Restrictions	Gastroenterologist or Rheumatologist
Coverage Duration	Plan Year
Other Criteria	Rheumatoid arthritis - At least 8-week maximum tolerated dose trial/failure or contraindication /intolerance to at least one nonbiologic DMARD and trial/failure of either Enbrel or Humira. Crohn's Disease - Trial/failure or contraindication/intolerance to at least one oral corticosteroid and Humira. For re-authorization, demonstrate improvement in clinical symptoms.
Prior Authorization Group	COPAXONE
Drug Names	COPAXONE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Mannitol hypersensitivity, concurrent use of any of the following medications: Interferon-beta therapy (Avonex, Betaseron, Extavia, or Rebif), or mitoxantrone.
Required Medical Information	MRI has been performed and has features suggestive of MS.
Age Restrictions	
Prescriber Restrictions	Neurologist
Coverage Duration	Plan Year
Other Criteria	Patients with previous use (12 or more months) of Copaxone must demonstrate one of the following clinical responses: decrease in the frequency of relapses, slowing of disease progression, MRI lesions have diminished with therapy, OR patient is stable on therapy.
Prior Authorization Group	DIFFERIN
Drug Names	ADAPALENE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	

Prior Authorization Group	DRONABINOL
Drug Names	DRONABINOL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	A. The diagnosis is documented as anorexia associated with weight loss in a patient with AIDS a. AND the patient has had an involuntary weight loss of greater than 10% of pre-illness baseline body weight or a body mass index (BMI) less than 20kg/m ² in the absence of a concurrent illness or medical condition other than HIV that may cause weight loss b. AND the patient has failed to respond to a 30-day drug regimen of megestrol (Megace) c. AND if the participant has received previous dronabinol therapy, he/she must show a positive response to therapy by maintaining or increasing their initial weight and/or muscle mass before initiating dronabinol therapy. B. The diagnosis is documented as nausea and vomiting associated with cancer chemotherapy in a cancer patient a. AND the participant is receiving a chemotherapy or radiation regimen b. AND if dronabinol is NOT being used as a full therapeutic replacement for an intravenous anti-emetic drug (e.g., ondansetron) c. AND if dronabinol is being used as a full therapeutic replacement for an intravenous anti-emetic drug (e.g., ondansetron) BUT dronabinol will NOT be within 48 hours of cancer therapy d. AND the patient has had a full trial and failure through at least one cycle of chemotherapy with IV ondansetron AND at least one of the following oral anti-emetic agents: metoclopramide, promethazine, prochlorperazine, meclizine, trimethobenzamide, oral 5-HT ₃ receptor antagonists e. AND if the participant has received previous dronabinol therapy, he/she must show a positive response by showing a reduced incidence or emesis and/or nausea.

Age Restrictions
Prescriber Restrictions
Coverage Duration
Other Criteria

6 months
 B vs D coverage determination per CMS guidelines

Prior Authorization Group
Drug Names
Covered Uses
Exclusion Criteria
Required Medical Information
Age Restrictions
Prescriber Restrictions
Coverage Duration
Other Criteria

ELAPRASE
 ELAPRASE
 All FDA approved indications not otherwise excluded from Part D
 Diagnosis confirmed by either DNA testing or enzymatic analysis.

Prior Authorization Group
Drug Names
Covered Uses

EMEND
 EMEND
 All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria	Concurrent use of the following medications: cisapride, pimozone.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	For the prevention of nausea and vomiting associated with highly or moderately emetogenic chemotherapy: Emend must be administered in combination with a 5-HT3 antagonist AND a corticosteroid (e.g., dexamethasone). B vs D coverage determination per CMS guidelines.
Prior Authorization Group	EMSAM
Drug Names	EMSAM
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Pheochromocytoma, concurrent use of the following medications: dextromethorphan, St. John's Wort.
Required Medical Information	A. Clinical diagnosis of major depressive disorder not responsive other antidepressants as demonstrated by at least 2 documented trials (clinically sufficient dose and duration of six weeks or longer) of the following: selective serotonin reuptake inhibitors (SSRI), serotonin/norepinephrine reuptake inhibitors (SNRI), bupropion, mirtazapine, or tricyclic/tetracyclic antidepressants B. OR clinical diagnosis of major depressive disorder for those patients who cannot take any oral preparations (including commercially available liquid antidepressants). C. For requests over 6 mg/24 hours, patient must agree to adhere to a tyramine restrictive diet.
Age Restrictions	
Prescriber Restrictions	Psychiatrist or receiving input from a psychiatry practice
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	EPLERENONE
Drug Names	EPLERENONE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	A. Diagnosis of hypertension or post-myocardial infarction with LVEF less than or equal to 40% and clinical evidence of CHF B. AND a serum potassium level less than 5.5 mEq/L. C. For diagnosis of post MI with LVEF less than or equal to 40% and clinical evidence of CHF, the patient must meet the following requirement: creatinine clearance greater than 30 mL/min. D. For the diagnosis of hypertension, the patient must meet the following requirements: the patient does not have type-2 diabetes with microalbuminuria AND the patient has a creatinine clearance greater than 50 mL/min AND the patient has tried and failed maximum tolerated doses of a 60-day trial or had unacceptable toxicity to spironolactone therapy.
Age Restrictions	

Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	EPO
Drug Names	PROCRIT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Uncontrolled hypertension. Red cell aplasia. Hgb greater than 12 g/dL (with the exception of surgery patients with Hgb greater than 13 g/dL).
Required Medical Information	Following labs performed within 30 days of request: Hgb less than or equal to 10 g/dL OR Hct less than or equal to 30% for initial authorization. Hgb less than 12 g/dL OR Hct less than 36% for re-authorization. Transferrin saturation greater than or equal to 20% and ferritin level greater than or equal to 100 ng/mL. For chemo-induced anemia - have serum Epo level less than or equal to 200 mUnits/mL prior to therapy. For anemia secondary to MDS and in HIV-infected patients - have serum Epo level less than or equal to 500 mUnits/mL prior to therapy. Surgery patients - require Hgb level greater than 10 but less than or equal to 13 g/dL.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 weeks
Other Criteria	For chemo-induced anemia - diagnosis is non-myeloid malignancy and receiving concomitant myelosuppressive chemotherapy regimen without an anticipated outcome of cure. For anemia in HIV-infected patient - must be on concurrent anti-retroviral therapy. Surgery patients - patient is at high risk for perioperative blood loss and must receive iron supplementation. Surgery is within 30 days of request. For re-authorization, must have an increase in Hgb of at least 1 g/dL or Hct of at least 3% since the initial Epo treatment.
Prior Authorization Group	EXJADE
Drug Names	EXJADE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Creatinine clearance less than 40 mL/min or serum creatinine more than 2 times the age appropriate upper limit of normal, platelet count less than 50 x 10 ⁹ /L, patients with high-risk MDS with poor performance status or an advanced malignancy, concurrent use of deferoxamine or iron-containing products.
Required Medical Information	The patient must meet all of the following criteria: diagnosis of transfusion-dependent anemia, patient has chronic iron overload due to blood transfusions, pretreatment serum ferritin level within the last 60 days of at least 1,000 mcg/L, and patient will have baseline and monthly monitoring of serum creatinine, creatinine clearance, serum transaminases and bilirubin. For reauthorization, if serum ferritin threshold is less than 500 mcg/L, prescriber should consider interrupting the dose of Exjade.

Age Restrictions	2 years of age and older
Prescriber Restrictions	Hematologist
Coverage Duration	3 months
Other Criteria	
Prior Authorization Group	EXTAVIA
Drug Names	EXTAVIA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Albumin hypersensitivity, concurrent use of any of the following medications: Interferon-beta therapy (Avonex, Betaseron, or Rebif), glatiramer acetate, or mitoxantrone. MRI has been performed and has features suggestive of MS (evidence of lesion).
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	Neurologist
Coverage Duration	Plan Year
Other Criteria	Patients with previous use (12 or more months) of Extavia must demonstrate one of the following clinical responses: decrease in the frequency of relapses, slowing of disease progression, MRI lesions have diminished with therapy, OR patient is stable on therapy.
Prior Authorization Group	FABRAZYME
Drug Names	FABRAZYME
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Diagnosis confirmed with an enzyme assay measuring a deficient activity of alpha-galactosidase enzyme or DNA testing.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	FENTANYL PATCH
Drug Names	FENTANYL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Patients who are not opioid tolerant, patients who do not require continuous opioid analgesia.
Required Medical Information	Assessment for clinical risk of opioid/substance abuse/addiction through CAGE questionnaire, Cyr-Wartman Screen, Skinner Trauma Screen Screener, Opioid Assessment for Patients with Pain (SOAPP) or other assessment tool.
Age Restrictions	2 years of age and older
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
Prior Authorization Group	GILENYA

Drug Names	GILENYA
Covered Uses	All FDA approved uses not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	For new starts, patient had an inadequate response to a trial of a beta interferon agent or Copaxone unless contraindicated or not tolerated.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	For continuation, patient is benefiting from Gilenya therapy.
Prior Authorization Group	GLEEVEC
Drug Names	GLEEVEC
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Chronic myeloid leukemia and acute lymphoblastic leukemia (ALL) must be positive for the Philadelphia chromosome or BCR-ABL gene.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	GONADOTROPIN
Drug Names	CHORIONIC GONADOTROPIN, NOVAREL, PREGNYL W/DILUENT BENZYL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Female. In males: anatomic obstruction, precocious puberty, prostatic carcinoma or other androgen-dependent neoplasm.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	GROWTH HORMONE
Drug Names	NORDITROPIN FLEXPRO, NORDITROPIN NORDIFLEX PEN, SAIZEN, SAIZEN CLICK.EASY, TEV-TROPIN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Malignancy, diabetic retinopathy, acute critical illness, concurrent use with Increlex, closed epiphyses for all pediatric patients, upper airway obstruction for PWS only, and other cause of hypoglycemia have been ruled out for neonates with hypoglycemia, pediatric GHD has been ruled out for ISS with one stimulation test.

Required Medical Information	For neonate with hypoglycemia, patient has pediatric GHD, has a randomly assigned GH level of less than 20 ng/mL. For SBS: patient is receiving specialized nutritional support and patient has not received GH therapy for more than 8 weeks lifetime. For HIV wasting: patient is on antiretroviral therapy, has tried and failed alternative therapies such as dronabinol or megestrol, and alternative causes of wasting have been ruled out. To continue therapy for HIV wasting, BMI has improved or stabilized and it has been at least 4 weeks since completion of last round of GH therapy. For all pediatric patients: patients have short stature and have been evaluated for other causes of growth failure. For pediatric GHD, has delayed bone age and failed 2 stimulation tests. For pediatric GHD with a pituitary or CNS disorder: patient has clinical evidence of GHD and low IGF-1/IGFBP3. For Turner syndrome patient: diagnosis confirmed with karyotyping. For chronic renal insufficiency patients: metabolic, endocrine and nutritional abnormalities have been treated or stabilized, and patient has not had a kidney transplant. For SGA patients: has a low birth weight, and has failed to manifest catch up growth by age 2. For PWS patients: therapy will be discontinued if patient develops severe respiratory impairment. For SHOX patients: diagnosis confirmed by molecular or genetic testing. For adults: assessed for other causes of GHD-like symptoms and failed 2 stimulation tests. For adult GHD with at least 3 pituitary hormone deficiencies or panhypopituitarism: have a low IGF-1. For adult GHD with less than 3 pituitary hormone deficiencies, low IGF-1 and failed one stimulation test. To continue therapy for pediatric patients, growing more than 2 cm per year, open epiphyses, and for PWS only: improved body composition. To continue therapy for adult patients: clinical improvement and IGF-1 to confirm appropriateness of treatment. For Turner syndrome and SGA, 2 years of age and older. For Noonan syndrome and SHOX, 3 years of age and older.
Age Restrictions	
Prescriber Restrictions	Endocrinologist, Pediatric Nephrologist, Gastroenterologist, Nutritional Support Specialist, Infectious Disease Specialist
Coverage Duration	12 weeks for HIV wasting, 8 weeks lifetime for SBS, 12 months for all other indications
Other Criteria	
Prior Authorization Group	HEPSERA
Drug Names	HEPSERA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Renal impairment without dosing adjustment, if the patient is taking/receiving tenofovir or PMPA.

Required Medical Information	A. The patient has been diagnosed with chronic hepatitis B. B. AND the patient has evidence of a positive HBsAg (+ or -) serological marker for greater than 6 months OR evidence by a liver biopsy showing chronic hepatitis. C. AND the patient has a Hepatitis B viral load greater than 20, 000 IU/ml (100,000 copies per ml). D. AND the patient has elevations in liver aminotransferases (ALT or AST) that are two (2) times greater than normal OR normal liver aminotransferase (ALT or AST) levels with evidence of significant disease found on biopsy. E. AND the patient is not receiving duplicate therapy with Intron A. F. AND documented evidence of diagnosis, serological markers or liver biopsy, viral load and liver aminotransferases. G. If the patient has received previous Hepsera treatment, there is documented clinical improvement shown by a drop in viral load or reduction in the patient's liver aminotransferases.
Age Restrictions	12 years and older
Prescriber Restrictions	Gastroenterologist or Infectious Disease Specialist or affiliated with an infectious disease or gastroenterology practice
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	HUMIRA
Drug Names	HUMIRA, HUMIRA PEN-CROHNS DISEASE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Active infection (including TB, sepsis), concurrent use with other biologics.
Required Medical Information	LTBI screening, and if results are positive, patient must have completed treatment or must currently be receiving treatment for LTBI. HBV infection ruled out or treatment initiated. If moderate to severe psoriatic arthritis with predominantly peripheral symptoms or moderate to severe RA or JIA, patient has had at least an 8-week maximum tolerated dose trial and failure or has an intolerance or contraindication to at least 1 nonbiologic DMARD (e.g., methotrexate, cyclosporine, azathioprine, sulfasalazine, leflunomide, hydroxychloroquine). If psoriatic arthritis with predominantly axial symptoms or ankylosing spondylitis, patient has tried and failed to respond to 2 NSAIDs unless patient has a contraindication or intolerance to NSAIDs. For moderate to severe plaque psoriasis, affected area is greater than 10% of BSA OR an area that will affect crucial daily functions (e.g., feet, hands). For moderate to severe plaque psoriasis, patient has tried and failed or has an intolerance or contraindication to at least a 60-day trial of 2 conventional therapies including high potency topical steroid therapy, calcipotriene, phototherapy, retinoids, methotrexate, or cyclosporine. If Crohn's disease, patient has tried and failed or has a contraindication or intolerance to at least a 60-day trial of 2 of the following conventional therapies: sulfasalazine, balsalazide, mesalamine, azathioprine, cyclosporine, methotrexate, mercaptopurine, corticosteroids OR patient has had an inadequate response or intolerance to either Remicade or Cimzia.

Age Restrictions	For PP, 18 years of age and older
Prescriber Restrictions	Rheumatologist, Dermatologist or Gastroenterologist
Coverage Duration	Plan Year
Other Criteria	For continuation of therapy, patient must show an improvement in clinical symptoms or delay in progression of disease.
Prior Authorization Group	INCIVEK
Drug Names	INCIVEK
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Failed previous therapy with a treatment regimen that includes a protease inhibitor (e.g., Incivek, Victrelis). Concomitant administration with a drug that is highly dependent on CYP3A for clearance or strongly induce CYP3A.
Required Medical Information	Hepatitis C virus (HCV) infection confirmed by presence of viral load in serum. HCV Genotype 1. HCV-RNA less than or equal to 1,000 IU/mL at week 4 of treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Initial: 6 weeks. Renewal: Up to 12 weeks.
Other Criteria	Must be given in combination with pegylated interferon (i.e., Pegasys or PegIntron) and ribavirin.
Prior Authorization Group	INCRELEX
Drug Names	INCRELEX
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Benzyl alcohol hypersensitivity, epiphyseal closure, IV administration of Increlex, active malignancy, use in neonates, concurrent use with GH therapy, secondary causes of IGF-1 deficiency.
Required Medical Information	Prior to starting therapy, a height greater than 3 SD below the mean for chronological age and sex, and an IGF-1 level greater than or equal to 3 SD below the mean for chronological age and gender. One stimulation test showing patient has a normal or elevated GH level.
Age Restrictions	Between 2 and 20 years of age
Prescriber Restrictions	Endocrinologist
Coverage Duration	Plan Year
Other Criteria	For continuation of therapy, there is an increase in height velocity by greater than 2.5 cm total growth in one year and patient has open epiphyses.
Prior Authorization Group	INFERGEN
Drug Names	INFERGEN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Decompensated liver failure/disease. Autoimmune disease. Use for nonresponse or relapse.

Required Medical Information Prior to initiating therapy, detectable levels of HCV RNA in the serum. Genotype 1,4: undetectable HCV RNA after 12 weeks of treatment OR at least 2 log decrease in HCV RNA after 12 weeks of therapy and undetectable HCV RNA after 24 weeks of treatment.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

3 to a total of 18 months depending on genotype and initial vs. renewal therapy
Monitored for evidence of depression. If used as monotherapy, patient should have a contraindication or intolerance to ribavirin.

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

INVEGA SUSTENNA

INVEGA SUSTENNA

All FDA approved indications not otherwise excluded from Part D

Treatment of patients with dementia-related psychosis, patients with occurrence of torsade de pointes, prior use of risperidone demonstrated a hypersensitivity reaction.

Required Medical Information

A. Diagnosis is an FDA-approved indication: acute and maintenance treatment of schizophrenia in adults. B. AND the diagnosis is NOT documented as dementia-related psychosis. C. AND Invega Sustenna therapy will not be used if prior use of risperidone demonstrated a hypersensitivity reaction D. AND the patient has a history of non-compliance and/or refuses to utilize oral medication E. AND the patient has received at least ONE of the following: a. three test doses of risperidone b. three test doses of oral Invega c. previous use of Invega Sustenna. F. If the patient is increasing the dose of Invega Sustenna, the patient must have a history of two prior injections.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Psychiatrist or receiving input from a psychiatry practice

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

ITRACONAZOLE

ITRACONAZOLE

All FDA approved indications not otherwise excluded from Part D

A. ventricular dysfunction (e.g., congestive heart failure (CHF) or history of CHF) do not use for onychomycosis. B. If the patient is taking/receiving any of the following: concomitant use with drugs metabolized by CYP3A4 (e.g., dofetilide, pimozide, quinidine).

Required Medical Information Patients with a diagnosis of blastomycosis, pulmonary and extrapulmonary OR patients with a diagnosis of histoplasmosis, including chronic cavitory pulmonary disease and disseminated, non-meningeal histoplasmosis OR patients with a diagnosis of aspergillosis, pulmonary and extrapulmonary OR patients with a diagnosis of onychomycosis of the toenail, with or without fingernail involvement, due to dermatophytes (tinea unguium) OR patients with a diagnosis of onychomycosis of the fingernail due to dermatophytes (tinea unguium), fungal diagnostic test to confirm onychomycosis.

Age Restrictions

Prescriber Restrictions

Coverage Duration 12 weeks

Other Criteria

Prior Authorization Group

IVIG

Drug Names

GAMMAGARD LIQUID, GAMUNEX

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

IgA deficiency with antibody formation. Intolerance to any of the components of immune globulin. Hyperprolinemia in those prescribed Privigen. Presence of risk factor for acute renal failure, unless the patient will receive IGIV products at the minimum concentration available and at the minimum rate of infusion practicable.

Required Medical Information

PID: history of infections with nonsustained response to antimicrobial therapy AND evidence of failed antibody development to established norms for vaccine stimulation. CIDP: presence of objective findings consistent with diagnosis. B-Cell CLL: history of recurrent bacterial infections.

Age Restrictions

Prescriber Restrictions

CIDP diagnosis by a neurologist

Coverage Duration

Plan Year

Other Criteria

B vs D coverage determination per CMS guidelines

Prior Authorization Group

LETAIRIS

Drug Names

LETAIRIS

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

AST/ALT level greater than 3 times ULN, pregnancy for females.

Required Medical Information

Diagnosis of pulmonary arterial hypertension (PAH), (WHO Group 1). NYHA class II or III symptoms. PAH been confirmed by right heart catheterization. If patient is an infant, PAH diagnosed by Doppler echocardiogram.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Plan Year

Other Criteria

IUD or two appropriate contraceptive methods for women of childbearing potential.

Prior Authorization Group

LEUKINE

Drug Names	LEUKINE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Administration within 24 hours preceding or following chemotherapy or radiotherapy, hypersensitivity to yeast-derived products, treatment of acute afebrile neutropenia, use to increase the chemotherapy dose intensity or dose schedule above established regimens.
Required Medical Information	For use following induction or consolidation chemotherapy in AML: there are less than 10% leukemic myeloid blasts in bone marrow or peripheral blood. For patients with nonmyeloid malignancies receiving myelosuppressive chemotherapy: Leukine may be used for the prevention of chemotherapy-induced neutropenia if the regimen has a 20% or more risk of neutropenia OR the patient experienced febrile neutropenia with a previous chemotherapy cycle. Patients without severe risk for neutropenia may also receive Leukine for prophylaxis if there is a risk for serious medical consequences due to febrile neutropenia and the intent of chemotherapy to prolong survival or cure the disease. Leukine is allowable for the treatment of febrile neutropenia in patients who have received prophylaxis with Leukine (or Neupogen) OR in patients at risk for infection-related complications. Leukine is allowable for patients with neutropenia due to myelodysplastic syndrome if they have a history or recurrent or resistant infections. All patients must receive baseline and regular monitoring of complete blood counts and platelet counts.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	
Prior Authorization Group	LEUPROLIDE DEPOT
Drug Names	LEUPROLIDE ACETATE, LUPRON DEPOT, LUPRON DEPOT-PED
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	For endometriosis, fibroids, and ovarian cancer: pregnancy and breastfeeding.
Required Medical Information	
Age Restrictions	For CPP, patient must be less than 12 years old if female and less than 13 years old if male.
Prescriber Restrictions	Obstetrician/Gynecologist, Oncologist, Endocrinologist. Urologist for diagnosis of prostate cancer.
Coverage Duration	Fibroids - 3 months, endometriosis - 6 months, ovarian cancer, prostate cancer, CPP - 12 months.

Other Criteria	For endometriosis only, patient must have completed a trial and failure of at least 2 of the following therapies: oral contraceptives, medroxyprogesterone, danazol. For ovarian CA, patient has recurrent or stage II to stage IV ovarian cancer. For prostate cancer, orchiectomy or estrogen therapy are unacceptable treatment options. For prostate cancer: used for initial treatment in advanced prostate cancer or when there is intermediate to high risk of recurrence. For prostate cancer, for adjuvant or neoadjuvant therapy.
Prior Authorization Group	LIDODERM
Drug Names	LIDODERM
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	A. The diagnosis is documented as post-herpetic neuralgia B. The skin where the patch is to be applied is intact (not broken or inflamed). C. The patient has completed a documented one month trial and failure of the following two medications: gabapentin or Lyrica D. OR the patient has a contraindication or demonstrated an adverse event to the prerequisite drugs.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Prior Authorization Group	LUMIZYME
Drug Names	LUMIZYME
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Diagnosis of Pompe disease was confirmed by an enzyme assay demonstrating a deficiency of GAA enzyme activity OR by DNA testing that identifies mutations in the GAA gene. Patient has a late (non-infantile) onset Pompe disease with no evidence of cardiac hypertrophy.
Age Restrictions	8 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Appropriate medical support is readily available when Lumizyme is administered in the event of anaphylaxis, severe allergic reaction, or acute cardiorespiratory failure.
Prior Authorization Group	METHYLPHENIDATES
Drug Names	CONCERTA, DEXMETHYLPHENIDATE HCL, METADATE CD, METHYLIN, METHYLPHENIDATE HCL, METHYLPHENIDATE HYDROCHLO, RITALIN LA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	MAOI concurrent use or within the last 14 days
Required Medical Information	Sleep studies for narcolepsy diagnosis

Age Restrictions	6 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Consider benefits of use versus the potential risks of serious cardiovascular events.
Prior Authorization Group	MOZOBIL
Drug Names	MOZOBIL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Concurrent diagnosis of leukemia
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	Mozobil is given in combination with granulocyte-colony stimulating factor
Prior Authorization Group	MYOZYME
Drug Names	MYOZYME
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Diagnosis confirmed by an enzymatic assay showing a deficiency in acid alpha glucosidase or DNA testing.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	NAGLAZYME
Drug Names	NAGLAZYME
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Diagnosis confirmed by an enzymatic assay showing a deficiency in N-acetylgalactosamine activity or DNA testing.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Must have at least one MPS VI symptom. For re-authorization, must demonstrate improvement in walking and/or stair-climbing capacity.
Prior Authorization Group	NEUPOGEN
Drug Names	NEUPOGEN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Administration within 24 hours preceding or following chemotherapy or radiotherapy, E coli hypersensitivity, use in acute afebrile neutropenia, use to increase the chemotherapy dose intensity or dose schedule above labeled use.

Required Medical Information For patients with nonmyeloid malignancies receiving myelosuppressive chemotherapy: Neupogen may be used for the prevention of chemotherapy-induced neutropenia if the regimen is associated with a 20% or more risk of neutropenia OR the patient experienced febrile neutropenia with a previous chemotherapy cycle. Patients without severe risk for neutropenia may receive Neupogen for prophylaxis if there is a risk for serious medical consequences due to febrile neutropenia and the intent of chemotherapy to prolong survival or cure the disease. Neupogen is allowable for the treatment of febrile neutropenia in patients who have received prophylaxis with Neupogen (or Leukine) OR in patients at risk for infection-related complications. Neupogen is allowable for patients with neutropenia due to myelodysplastic syndrome if they have a history or recurrent or resistant infections. All patients must receive baseline and regular monitoring of complete blood counts and platelet counts.

Age Restrictions

Prescriber Restrictions

Coverage Duration

6 months

Other Criteria

Prior Authorization Group

NICOTINE

Drug Names

NICOTROL INHALER, NICOTROL NS

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Required Medical Information

Documentation that the patient is enrolled in a smoking cessation program

Age Restrictions

Prescriber Restrictions

Coverage Duration

6 months

Other Criteria

Prior Authorization Group

NUEDEXTA

Drug Names

NUEDEXTA

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Concomitantly taking other drugs containing quinidine, quinine, mefloquine, monoamine oxidase inhibitors (MAOIs), or drugs that both prolong QT interval and are metabolized by CYP2D6 (e.g., thioridazine and pimozide), patient has a prolonged QT interval, congenital long QT syndrome or a history suggestive of torsades de pointes, or heart failure, patient has complete atrioventricular (AV) block without implanted pacemaker, or is at high risk of complete AV block. Dose in excess of 2 capsules per day.

Required Medical Information

Patient has amyotrophic lateral sclerosis (ALS) OR multiple sclerosis (MS)

Age Restrictions

Prescriber Restrictions

Coverage Duration

Plan Year

Other Criteria

Prior Authorization Group	NUVIGIL
Drug Names	NUVIGIL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	If diagnosis is narcolepsy require sleep lab evaluation, if diagnosis of OSAHS require polysomnography and whether the patient is using CPAP (or CPAP is contraindicated or ineffective). If diagnosis of mild obstructive sleep apnea/hypopnea syndrome, patient may use, and be compliant with, an oral appliance as alternative to CPAP.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	OCTREOTIDE
Drug Names	OCTREOTIDE ACETATE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	ORFADIN
Drug Names	ORFADIN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Confirmation of diagnosis by either biochemical testing (e.g., detection of succinylacetone in urine) and appropriate clinical picture OR DNA testing (mutation analysis).
Age Restrictions	
Prescriber Restrictions	Endocrinologist, Clinical or Biochemical Geneticist, Gastroenterologist, Hepatologist
Coverage Duration	Plan Year
Other Criteria	Protein-restricted diet that is low in phenylalanine and tyrosine.
Prior Authorization Group	OSTEOPOROSIS
Drug Names	FORTEO
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Paget's disease, elevated alkaline phosphatase, pre-existing hypercalcemia, skeletal malignancies, prior radiation of the skeleton, cumulative use of Forteo for more than 24 months lifetime, concurrent bisphosphonate use.

Required Medical Information Patients treated with Forteo meet one of the following criteria for fracture risk: prior fragility fracture OR T score less than or equal to -2.5 and family history of fracture OR inadequate response to a bisphosphonate trial of a minimum of one year (unless bisphosphonate is contraindicated or patient was intolerant to bisphosphonate therapy).

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

OXSORALEN

OXSORALEN ULTRA

All FDA approved indications not otherwise excluded from Part D

Albinism, aphakia, melanoma, porphyria, skin photosensitivity disorder, systemic lupus erythematosus (SLE), xeroderma pigmentosum, invasive squamous cell carcinoma or current skin burns.

Required Medical Information

The patient must be diagnosed with T-cell lymphoma OR psoriasis OR vitiligo AND if the diagnosis is psoriasis, the patient must have previous trial/failure or contraindication to at least one topical steroid.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Dermatologist or Oncologist or affiliated with a dermatologist/oncologist practice

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

PEGASYS

PEGASYS

All FDA approved indications not otherwise excluded from Part D

Current psychosis or a history of psychosis, severe depression, severe thrombocytopenia, decompensated cirrhosis, serious active infection.

Required Medical Information

HCV: HCV genotype, detectable HCV RNA within 90 days prior to starting therapy.

HBV: HBsAg positive or liver biopsy showing chronic hepatitis AND appropriate HBV DNA levels for HBeAg status AND elevated liver enzymes. CML: unable to tolerate tyrosine kinase inhibitors or post-transplant if not in remission or with relapse. For triple therapy with Pegasys and ribavirin and protease inhibitor (PI): HCV Genotype 1. HCV-RNA less than 100 IU/mL at week 12 of treatment for Victrelis triple therapy. HCV-RNA less than or equal to 1,000 IU/mL at weeks 4 and 12 of treatment for Incivek triple therapy. Undetectable HCV-RNA at week 24 of treatment.

Age Restrictions

Prescriber Restrictions

Coverage Duration

ID specialist, Gastroenterology, Oncologist

HCV=Based on genotype and response 12 to 72 wks for dual tx 6 to 48 wks for triple tx
HBV,CML=12 mos

Other Criteria	HBV: Not receiving duplicate therapy. For reauthorization, clinical improvement. HCV: Retreatment allowed in those who did not receive optimal HCV treatment. For reauthorization at 12 weeks, early virologic response.
Prior Authorization Group	PEGINTRON
Drug Names	PEG-INTRON, PEG-INTRON REDIPEN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Current psychosis or a history of psychosis, severe depression, severe thrombocytopenia, decompensated cirrhosis, serious active infection.
Required Medical Information	HCV: HCV genotype. Detectable HCV RNA within 90 days of initiating therapy. CML: unable to tolerate TKIs or post-transplant if not in remission or with relapse. For triple therapy with PegIntron and ribavirin and protease inhibitor (PI):HCV Genotype 1. HCV-RNA less than 100 IU/mL at week 12 of treatment for Victrelis triple therapy. HCV-RNA less than or equal to 1,000 IU/mL at weeks 4 and 12 of treatment for Incivek triple therapy. Undetectable HCV-RNA at week 24 of treatment.
Age Restrictions	
Prescriber Restrictions	ID specialist, Gastroenterology, Oncology
Coverage Duration	HCV=Based on genotype and response 12 to 72 wks for dual tx, 6 to 48 wks for triple tx. CML=12 mos
Other Criteria	HCV: Retreatment allowed for those who did not receive optimal HCV treatment. For reauthorization at 12 weeks, early virologic response.
Prior Authorization Group	PROLIA
Drug Names	PROLIA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Hypocalcemia
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Patient will be adequately supplemented with calcium and vitamin D.
Prior Authorization Group	PROMACTA
Drug Names	PROMACTA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	

Required Medical Information For new starts, a pretreatment platelet count less than 30,000/microL or a platelet count less than or equal to 50,000/microL with significant mucous membrane bleeding or risk factors for bleeding are required. For continuation of therapy, an increase in platelet count to a level that is sufficient to avoid clinically important bleeding after at least 4 weeks of maximum dose therapy is required. For continuation of therapy, alanine aminotransferase levels must not be greater than or equal to 3 times the upper limit of normal and must not be progressive, persistent, or accompanied by increased bilirubin, symptoms of liver injury, or hepatic decompensation.

Age Restrictions

Prescriber Restrictions

Coverage Duration

6 mo initially, 12 mo renewal w/ platelet response, 3 mo renewal w/out platelet response

Other Criteria

Prior Authorization Group

PROVIGIL

Drug Names

PROVIGIL

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Required Medical Information

If diagnosis is narcolepsy require sleep lab evaluation, if diagnosis of OSAHS require polysomnography and whether the patient is using CPAP (or CPAP is contraindicated or ineffective). If diagnosis of mild obstructive sleep apnea/hypopnea syndrome, patient may use, and be compliant with, an oral appliance as alternative to CPAP.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Plan Year

Other Criteria

Prior Authorization Group

PULMOZYME

Drug Names

PULMOZYME

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Required Medical Information

For continuation of therapy in patients less than 6 years of age, a clinical reason to continue therapy, such as symptomatic improvement, is required. For continuation of therapy in patients more than 6 years of age, no reduction in pulmonary function tests of greater than 10% from baseline or a clinical reason to continue therapy is required.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Plan Year

Other Criteria

B vs D coverage determination per CMS guidelines

Prior Authorization Group

RANEXA

Drug Names

RANEXA

<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Patients with clinically significant hepatic impairment.
<i>Required Medical Information</i>	A. The diagnosis documented as chronic angina with symptoms limiting daily activities. B. AND the patient is NOT receiving a medication that prolongs the QT interval C. AND the patient has tried, failed and/or been intolerant (continues to have angina symptoms that limits daily activities) to a 30-day trial of the following: a. A nitrate b. A beta blocker OR a calcium channel blocker. D. AND if the patient has received prior treatment with Ranexa, the patient must experience a decrease in angina frequency since initiating treatment.
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	Cardiologist or affiliated with a cardiology practice
<i>Coverage Duration</i>	3 months initial, 12 months renewal
<i>Other Criteria</i>	
<i>Prior Authorization Group</i>	REBIF
<i>Drug Names</i>	REBIF, REBIF TITRATION PACK
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	
<i>Required Medical Information</i>	
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	
<i>Coverage Duration</i>	Plan Year
<i>Other Criteria</i>	
<i>Prior Authorization Group</i>	RELISTOR
<i>Drug Names</i>	RELISTOR
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Mechanical gastrointestinal obstruction, known or suspected.
<i>Required Medical Information</i>	A. Relistor is being prescribed for treatment of opioid-induced constipation in patients with advanced illness who are receiving palliative care. B. patient must have previous trial/failure of polyethylene glycol.
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	
<i>Coverage Duration</i>	4 Months
<i>Other Criteria</i>	
<i>Prior Authorization Group</i>	REMICADE
<i>Drug Names</i>	REMICADE
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Active infection (including TB, sepsis), concurrent use with other biologics, moderate to severe HF (NYHA Functional Class III/IV) at doses greater than 5mg/kg, murine protein hypersensitivity.

Required Medical Information

LTBI screening, and if results are positive, patient must have completed treatment or must currently be receiving treatment for LTBI. HBV infection ruled out or treatment initiated. If moderate to severe RA, patient has had at least an 8-week maximum tolerated dose trial and failure or has an intolerance or contraindication to at least 1 nonbiologic DMARD (e.g., methotrexate, cyclosporine, azathioprine, sulfasalazine, leflunomide, hydroxychloroquine) AND patient has had a previous trial and failure to Enbrel or Humira. For moderate to severe RA, patient is receiving MTX concurrently. If moderate to severe psoriatic arthritis with predominantly peripheral symptoms, patient had at least an 8-week maximum tolerated dose trial and failure or has an intolerance or contraindication to at least 1 of the following nonbiologic DMARDs (e.g., methotrexate, cyclosporine, azathioprine, sulfasalazine, leflunomide, hydroxychloroquine) AND patient had a previous trial and failure to Enbrel or Humira. If moderate to severe psoriatic arthritis with predominantly axial symptoms or ankylosing spondylitis, patient has tried and failed to respond to 2 NSAIDs unless patient has a contraindication or intolerance to NSAIDs.

Age Restrictions

For plaque psoriasis, 18 years of age and older.

Prescriber Restrictions

Rheumatologist, Dermatologist and Gastroenterology

Coverage Duration

Plan Year

Other Criteria

If moderate to severe plaque psoriasis, affected area is greater than 10% of body surface area OR an area that will affect crucial daily functions (e.g., feet, hands). For moderate to severe plaque psoriasis, patient has tried and failed (or has an intolerance or contraindication to) at least a 60 day trial of 2 conventional therapies including high potency topical steroid therapy, calcipotriene, phototherapy, retinoids, or methotrexate. If ulcerative colitis: patient has tried and failed (or has a contraindication or intolerance to) at least a 60 day trial of 2 conventional therapies such as sulfasalazine, balsalazide, mesalamine, or corticosteroids. If Crohn's disease: patient has tried and failed (or has an intolerance or contraindication to) 1 conventional therapy (eg, corticosteroids) and 1 biologic therapy ((ie, Humira or Cimzia). For continuation of therapy, patient must show an improvement in clinical symptoms or delay in progression of disease.

Prior Authorization Group

REVATIO

Drug Names

REVATIO

Covered Uses

All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria

Nitrate therapy

Required Medical Information

Diagnosis of pulmonary arterial hypertension (PAH), (WHO Group 1). PAH been confirmed by right heart catheterization. If patient is an infant, PAH diagnosed by Doppler echocardiogram. The patient has had an inadequate response or intolerance to Adcirca.

Age Restrictions

Prescriber Restrictions

Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	REVLIMID
Drug Names	REVLIMID
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Pregnancy
Required Medical Information	Diffuse large B-cell lymphoma: relapsed, refractory or progressive disease. Mantle cell lymphoma: as monotherapy for relapsed, refractory or progressive disease. Systemic light chain amyloidosis: use as primary treatment in combination with dexamethasone. Myeloma: FDA approved uses OR palliative treatment OR primary induction in combination with dexamethasone OR maintenance therapy as monotherapy after stem cell transplant or in responders to primary induction therapy. Low or Intermediate-1 Risk MDS: for those with 5q deletion, patients should have transfusion dependent anemia OR clinically significant cytopenias and symptomatic anemia. For those with non-5q deletion and symptomatic anemia, patients should have failed to respond to epoetin alfa or darbepoetin OR have serum erythropoietin levels greater than 500 mU/mL and a low probability of response to immunosuppressive therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Male and female patients of child-bearing potential should be instructed on the importance of proper utilization of appropriate contraceptive methods for Revlimid use. Patients should be monitored for signs and symptoms of thromboembolism.
Prior Authorization Group	RIBAVIRIN
Drug Names	REBETOL, RIBAPAK, RIBASPHERE, RIBAVIRIN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Hemoglobin less than 8.5 g/dL. Hemoglobinopathy. History of unstable heart disease. Creatinine clearance less than 50 mL/minute unless the patient will receive a modified dose of ribavirin. Pregnancy (self or partner).
Required Medical Information	HCV relapse or nonresponse: patient was initially treated with a less than optimal regimen for HCV or there is a clinical reason that suggests that patient will respond to retreatment AND ribavirin will be used in combination with a pegylated interferon AND this is the first time the patient is being retreated with a pegylated IFN and ribavirin. For triple therapy with PEG-IFN and ribavirin and protease inhibitor (PI): Detectable HCV-RNA within 90 days prior to starting therapy. HCV Genotype 1. HCV-RNA less than 100 IU/mL at week 12 of treatment for Victrelis triple therapy. HCV-RNA less than or equal to 1,000 IU/mL at weeks 4 and 12 of treatment for Incivek triple therapy. Undetectable HCV-RNA at week 24 of treatment.
Age Restrictions	

Prescriber Restrictions	
Coverage Duration	Based on genotype and response 12 to 72 weeks for dual therapy and 6 to 48 weeks for triple therapy.
Other Criteria	Patient has been instructed to practice effective contraception during therapy and for six months after stopping ribavirin therapy.
Prior Authorization Group	RISPERDAL CONSTA
Drug Names	RISPERDAL CONSTA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Dementia-related psychosis.
Required Medical Information	A. The patient has a history of non-compliance or refuses to utilize oral medications. B. The patient must have history of 3 test doses of oral risperidone. C. If the patient is increasing the dose of Risperdal Consta they must have a history of two prior injections.
Age Restrictions	
Prescriber Restrictions	Psychiatrist or receiving input from a psychiatry practice
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	RITUXAN
Drug Names	RITUXAN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Hypersensitivity to murine proteins or chimeric monoclonal antibodies, or patient is receiving live vaccines.
Required Medical Information	For rheumatoid arthritis (RA), an inadequate response to a nonbiologic DMARD (8-week trial) and an inadequate response to either Enbrel or Humira are required. For continuation of RA therapy, improvement in clinical symptoms that may include improvement in tender and swollen joint count, mobility, or stiffness, or delay in progression of disease is required.
Age Restrictions	
Prescriber Restrictions	Rheumatologist and Oncologist
Coverage Duration	6 months
Other Criteria	For chronic lymphocytic leukemia and adult acute lymphoblastic leukemia, Rituxan must be used in combination with chemotherapy.
Prior Authorization Group	SABRIL
Drug Names	SABRIL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Patients with or at high risk of vision loss. Patients using other medications associated with serious adverse ophthalmic effects such as retinopathy or glaucoma.
Required Medical Information	Vision is assessed at baseline or will be assessed by an ophthalmologist no longer than 4 weeks after starting Sabril.

Age Restrictions	For initial treatment of infantile spasms (IS), patient is between 1 month to 2 years of age. For initial treatment of complex partial seizures (CPS), patient is 16 years of age or older.
Prescriber Restrictions	
Coverage Duration	Infantile spasms: initial 4 weeks, reauth 6 months. CPS: initial 3 months, reauth 12 months.
Other Criteria	For CPS, Sabril is being used as adjunctive therapy and the patient has failed an adequate regimen with either carbamazepine or phenytoin, unless there is a contraindication or intolerance. For IS, Sabril is being used as monotherapy. For continuation of therapy: patient has shown substantial clinical benefit from Sabril therapy and patient's vision will be assessed by an ophthalmologist every 3 months.
Prior Authorization Group	SANCUSO
Drug Names	SANCUSO
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Documentation showing that the patient has had a previous trial/failure to any oral therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	SANDOSTATIN LAR
Drug Names	SANDOSTATIN LAR DEPOT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	Effective and tolerated prior Sandostatin injection (not depot form) therapy for at least 2 weeks.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	SOMATULINE DEPOT
Drug Names	SOMATULINE DEPOT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	

Prior Authorization Group	SOMAVERT
Drug Names	SOMAVERT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	IV administration of Somavert, concomitant use of Sandostatin or Somatuline.
Required Medical Information	An elevated IGF-1 level or elevated GH level with a glucose tolerance test. Patient has tried and failed at least a 3 month trial of Sandostatin or Somatuline.
Age Restrictions	
Prescriber Restrictions	Endocrinologist
Coverage Duration	Plan Year
Other Criteria	For retreatment, reduction in IGF-1 level from baseline.
Prior Authorization Group	STELARA
Drug Names	STELARA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Active infection (including tuberculosis), concurrent use with other biologics
Required Medical Information	Screening for latent tuberculosis is required. If results are positive, patient must have completed treatment or must currently be receiving treatment for latent tuberculosis.
Age Restrictions	18 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	For renewal, patient's condition must have improved or stabilized.
Prior Authorization Group	STRATTERA
Drug Names	STRATTERA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	MAOI concurrent use or within the last 14 days
Required Medical Information	
Age Restrictions	6 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Monitor for suicidality, clinical worsening, changes in behavior, blood pressure changes, heart rate changes, liver injury.
Prior Authorization Group	SUTENT
Drug Names	SUTENT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Concomitant treatment with interferon-alpha or interleukin-2, clinical manifestations of congestive heart failure.
Required Medical Information	For gastrointestinal stromal tumor (GIST), disease progression while on an at least 30-day regimen of Gleevec or intolerance to Gleevec is required. For continuation of therapy (all covered uses), no evidence of disease progression or tumor growth since initiation of therapy is required.
Age Restrictions	

Prescriber Restrictions	Oncologist
Coverage Duration	Plan Year
Other Criteria	For metastatic thyroid carcinoma, tumors must be at non-central nervous system sites and must be non-radioiodine avid.
Prior Authorization Group	SYLATRON
Drug Names	SYLATRON
Covered Uses	All FDA approved indications not otherwise excluded from Part D, chronic myelogenous leukemia (CML)
Exclusion Criteria	Autoimmune hepatitis, decompensated hepatic disease, uncontrolled major depression or severe mental illness
Required Medical Information	For melanoma, all of the following initial criteria are required: melanoma has microscopic or gross nodal involvement AND Sylatron is used following surgical resection of the tumor and complete lymphadenectomy AND Sylatron is being requested for use within 84 days (12 weeks) of the surgical resection. For CML, the patient meets one of the following criteria: patient is unable to tolerate a tyrosine kinase inhibitor (eg, imatinib, dasatinib, or nilotinib) OR patient is post-transplant without remission or with relapse of CML.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	The patient is monitored and evaluated for signs and symptoms of depression and other psychiatric symptoms throughout treatment with Sylatron
Prior Authorization Group	SYMLIN
Drug Names	SYMLIN, SYMLINPEN 120, SYMLINPEN 60
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Severe hypoglycemia that required assistance during the past 6 months, gastroparesis, patient requires drug therapy to stimulate gastrointestinal motility, the presence of hypoglycemia unawareness (i.e., inability to detect and act upon the signs or symptoms of hypoglycemia).
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	The patient must have inadequate glycemic control (HbA1c greater than 7% but less than 9%) at initiation of therapy, patient currently receiving optimal mealtime insulin therapy, or patient has taken Symlin in previous 6 months.
Prior Authorization Group	TARCEVA
Drug Names	TARCEVA
Covered Uses	All FDA approved indications not otherwise excluded from Part D

Exclusion Criteria	
Required Medical Information	For 1st line therapy of locally advanced metastatic NSCLC, patient should have a known active EGFR mutation and is not currently smoking. For locally advanced, unresectable or metastatic pancreatic cancer, Tarceva will be used in combination with gemcitabine.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	TARGRETIN
Drug Names	TARGRETIN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Pregnancy
Required Medical Information	The patient must be diagnosed with cutaneous manifestations of cutaneous T-cell lymphoma (CTCL) AND the patient must have failed at least one prior systemic therapy OR the patient must be diagnosed with stage 1A or 1B cutaneous T-cell lymphoma with cutaneous manifestations.
Age Restrictions	
Prescriber Restrictions	Dermatologist and Oncologist
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	TERBINAFINE
Drug Names	TERBINAFINE HCL
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	LFTs, fungal diagnostic test (e.g., KOH preparation, positive fungal culture, or nail biopsy)
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	2 months for fingernails only, 3 months if toenail involvement, 6 weeks for tinea infections
Other Criteria	
Prior Authorization Group	TESTOSTERONES
Drug Names	ANDRODERM, ANDROGEL, ANDROGEL PUMP, TESTIM
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Female, prostate cancer, breast cancer.
Required Medical Information	Before the start of testosterone therapy patient has (or patient currently has) a confirmed low testosterone level (i.e. total testosterone less than 300 ng/dL, free or bioavailable, testosterone less than 5 ng/dL) or absence of endogenous testosterone.
Age Restrictions	

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

THALOMID

THALOMID

All FDA approved indications not otherwise excluded from Part D

Pregnancy

Mantle cell lymphoma: relapsed, refractory or progressive disease in combination with rituximab. Waldenstrom's macroglobulinemia: third-line, palliative treatment OR as monotherapy for primary treatment in patients with symptomatic hyperviscosity with plasmapheresis. ENL: if moderate to severe neuritis, Thalomid will not be used as monotherapy. Systemic light chain amyloidosis: use with dexamethasone. Myeloma: advanced, refractory disease OR induction therapy with dexamethasone OR induction therapy in transplant ineligible patients used in combination with melphalan and prednisone OR maintenance therapy as monotherapy after stem cell transplant or in those who responded to primary induction therapy.

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Patients should be informed to be observant for signs and symptoms of thromboembolism. Male and female patients of child-bearing potential should be instructed on importance of proper utilization of appropriate contraceptive methods for Thalomid use.

Prior Authorization Group

Drug Names

Covered Uses

Exclusion Criteria

Required Medical Information

TOPICAL IMMUNOSUPPRESSANT

ELIDEL, PROTOPIC

All FDA approved indications not otherwise excluded from Part D

A. The diagnosis is documented as atopic dermatitis or eczema. B. AND patients must be at least 2 years of age C. AND patients who have completed a documented trial and failure of at least two medium or higher potency topical steroids or have documented intolerance or unresponsiveness to medium or higher potency topical steroids D. AND patients have been advised that Elidel and Protopic should only be used to treat the immediate problem and then should be stopped when the condition improves.

2 years of age and older

Age Restrictions

Prescriber Restrictions

Coverage Duration

Other Criteria

Plan Year

Prior Authorization Group

Drug Names

TOPICAL-ULCERS

REGRANEX

<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Neoplasm(s) at site(s) of application
<i>Required Medical Information</i>	A. Must be used for treatment of lower-extremity diabetic ulcers B. AND the ulcer must extend into subcutaneous tissue or beyond C. AND the tissue must have an adequate blood supply D. AND the patient must have concurrent good ulcer treatment practices including ALL of the following: a. Debridement b. Pressure relief c. Infection relief E. AND the ulcer must be less than 10 cm ² in size.
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	
<i>Coverage Duration</i>	10 weeks
<i>Other Criteria</i>	
<i>Prior Authorization Group</i>	TRACLEER
<i>Drug Names</i>	TRACLEER
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	AST/ALT level greater than 3 times ULN. Pregnancy. Concomitant use of cyclosporine A or glyburide.
<i>Required Medical Information</i>	PAH confirmed by right heart catheterization.
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	
<i>Coverage Duration</i>	Plan Year
<i>Other Criteria</i>	Female patient of childbearing potential must use more than one method of contraception concurrently.
<i>Prior Authorization Group</i>	TYZEKA
<i>Drug Names</i>	TYZEKA
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	
<i>Required Medical Information</i>	A. The patient has been diagnosed with chronic hepatitis B. B. AND the patient has evidence of a positive HBsAg (+ or -) serological marker for greater than 6 months OR evidence by a liver biopsy showing chronic hepatitis. C. AND the patient has a Hepatitis B viral load greater than 20,000 IU/mL (100,000 copies/mL). D. AND the patient has elevations in liver aminotransferases (ALT or AST) that are two (2) times greater than normal OR normal liver aminotransferase (ALT or AST) levels with evidence of significant disease found on biopsy. E. AND the patient has been tested for HIV. (Tyzeka therapy can cause HIV resistance in untreated HIV infection). F. AND if the patient has received previous Tyzeka treatment, there is documented clinical improvement shown by a drop in viral load or reduction in the patient's liver aminotransferases. G. AND the patient is not receiving duplicate therapy that includes Baraclade, Epivir and/or Intron A. H. AND evidence of diagnosis, serological markers or liver biopsy, viral load, and liver aminotransferases is documented in patient's chart.
<i>Age Restrictions</i>	16 years of age and older

Prescriber Restrictions	Infectious Disease Specialist or Gastroenterologist or affiliated with an infectious disease or gastroenterology practice
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	VFEND
Drug Names	VFEND, VFEND IV, VORICONAZOLE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Concurrent use of St. John's Wort (<i>Hypericum perforatum</i>).
Required Medical Information	A. The patient is diagnosed with invasive aspergillosis B. OR the patient is diagnosed with candidiasis and/or candidemia C. AND the patient has previous trial and failure or contraindication to BOTH fluconazole and itraconazole D. OR the patient is diagnosed with <i>Fusarium</i> or <i>Scedosporium</i> sp. E. AND Vfend is being used as salvage therapy due to failure, intolerance or contraindication of other therapies.
Age Restrictions	
Prescriber Restrictions	Infectious Disease Specialist or affiliated with an infectious disease practice
Coverage Duration	1 month
Other Criteria	
Prior Authorization Group	VICTRELIS
Drug Names	VICTRELIS
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Failed previous therapy with a treatment regimen that includes a protease inhibitor (e.g., Incivek, Victrelis). Concomitant administration with a drug that is highly dependent on CYP3A4/5 for clearance or potent CYP3A4/5 inducer.
Required Medical Information	Hepatitis C virus (HCV) infection confirmed by presence of viral load in serum. HCV Genotype 1. HCV-RNA less than 100 IU/mL at week 12 of treatment. Undetectable HCV-RNA at week 24 of treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Initial: 10 weeks. Renewal: Up to 44 weeks.
Other Criteria	Must be given in combination with pegylated interferon (i.e., Pegasys or PegIntron) and ribavirin. Must receive 4 weeks of pegylated interferon and ribavirin prior to starting Victrelis.
Prior Authorization Group	VOTRIENT
Drug Names	VOTRIENT
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Alanine transaminase (ALT) greater than 3 times the upper limit of normal (ULN) and bilirubin greater than 2 times the ULN.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	

Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	VPRIV
Drug Names	VPRIV
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Patients taking miglustat (Zavesca)
Required Medical Information	Diagnosis confirmed by bone marrow histology, DNA testing or measurement of beta-glucocerebrosidase enzyme activity less than 30%.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Patient must have at least one of the following conditions: anemia, thrombocytopenia, bone disease, hepatomegaly, splenomegaly. Patients who have previously received 24 months of VPRIV therapy must have a decrease in liver and spleen volume and/or increase in platelet count and/or increase in hemoglobin concentration for reauthorization.
Prior Authorization Group	XENAZINE
Drug Names	XENAZINE
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	Untreated or inadequately treated depression or actively suicidal, history of hepatic disease or torsade de pointes, use in combination with MAO inhibitors or reserpine (or it has been less than 20 days since reserpine was discontinued).
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	XIFAXAN
Drug Names	XIFAXAN
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	For hepatic encephalopathy, Xifaxan exceeding the recommended dose of two 550mg tablets daily.
Required Medical Information	
Age Restrictions	18 years of age and older
Prescriber Restrictions	
Coverage Duration	Hepatic encephalopathy-6 months
Other Criteria	
Prior Authorization Group	XOLAIR
Drug Names	XOLAIR

<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Patient is a current smoker, hamster protein hypersensitivity, omalizumab hypersensitivity.
<i>Required Medical Information</i>	Patient has evidence of reversible disease (demonstrates at least 20 percent improvement in PEF with a short-acting bronchodilator challenge), patient has experienced 2 or more asthma exacerbations per month within the last 3 months, positive skin test to at least 1 perennial aeroallergen, baseline IgE level at or above 30 IU/mL, asthma is inadequately controlled despite adherent use of inhaled corticosteroids for at least 6 months, inadequate response to a 3 month trial of a leukotriene modifier and long-acting beta2-agonist (unless patient demonstrates intolerance to the therapeutic trial).
<i>Age Restrictions</i>	12 years of age and older
<i>Prescriber Restrictions</i>	Pulmonologist, Allergist or Immunologist
<i>Coverage Duration</i>	Plan Year
<i>Other Criteria</i>	Patients with prior Xolair therapy must demonstrate an improvement in asthma control with use of Xolair.
<i>Prior Authorization Group</i>	XYREM
<i>Drug Names</i>	XYREM
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	If the patient is taking/receiving any of the following: anxiolytics, sedatives, hypnotics, barbiturates, benzodiazepines or ethanol.
<i>Required Medical Information</i>	A. The diagnosis is documented as excessive daytime sleepiness with symptoms that limit their ability to perform normal daily activities. B. AND the diagnosis is documented as cataplexy (a condition characterized by weak or paralyzed muscles) in patients with narcolepsy. C. AND if the patient has received prior treatment with Xyrem, the patient must experience a decrease in daytime sleepiness and/or cataplexy in a narcoleptic patient.
<i>Age Restrictions</i>	
<i>Prescriber Restrictions</i>	
<i>Coverage Duration</i>	3 months
<i>Other Criteria</i>	
<i>Prior Authorization Group</i>	ZAVESCA
<i>Drug Names</i>	ZAVESCA
<i>Covered Uses</i>	All FDA approved indications not otherwise excluded from Part D
<i>Exclusion Criteria</i>	Renal failure. Pregnancy. Labor/obstetric delivery.

Required Medical Information	Diagnosis confirmed by bone marrow histology, DNA testing or measurement of b-glucocerebrosidase enzyme activity less than 30%. Trial and failure of enzyme replacement therapy or is not a therapeutic option. Female patients of childbearing age will be on a form of contraception or have no ability to conceive and been educated on the potential dangers of Zavesca therapy. Must demonstrate a decrease in liver and spleen volume and/or increase in platelet count and/or increase in Hgb concentration in patients who have previously received 24 months of Zavesca therapy.
Age Restrictions	18 years of age and older
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	
Prior Authorization Group	ZYTIGA
Drug Names	ZYTIGA
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	Used in combination with prednisone. Received prior chemotherapy containing docetaxel.
Prior Authorization Group	ZYVOX
Drug Names	ZYVOX
Covered Uses	All FDA approved indications not otherwise excluded from Part D
Exclusion Criteria	If the patient is receiving MAO inhibitors (e.g., phenelzine, isocarboxazid) concomitantly with Zyvox
Required Medical Information	Culture and sensitivity demonstrates that the bacteria are susceptible to Zyvox OR, if unavailable, local bacterial susceptibility patterns support the use of Zyvox
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Plan Year
Other Criteria	