ABALOPARATIDE

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Exclusion Criteria	POSTMENOPAUSAL OSTEOPOROSIS: PATIENT HAS RECEIVED A TOTAL OF 24 MONTHS CUMULATIVE TREATMENT WITH ANY PARATHYROID HORMONE THERAPY.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	POSTMENOPAUSAL OSTEOPOROSIS: ONE OF THE FOLLOWING: 1) HIGH RISK FOR FRACTURES DEFINED AS ONE OF THE FOLLOWING: A) HISTORY OF OSTEOPOROTIC (I.E., FRAGILITY, LOW TRAUMA) FRACTURE(S), B) 2 OR MORE RISK FACTORS FOR FRACTURE (E.G., HISTORY OF MULTIPLE RECENT LOW TRAUMA FRACTURES, BMD T-SCORE LESS THAN OR EQUAL TO -2.5, CORTICOSTEROID USE, OR USE OF GNRH ANALOGS), OR C) NO PRIOR TREATMENT FOR OSTEOPOROSIS AND FRAX SCORE OF AT LEAST 20% FOR ANY MAJOR FRACTURE OR OF AT LEAST 3% FOR HIP FRACTURE. 2) UNABLE TO USE ORAL THERAPY (I.E., UPPER GASTROINTESTINAL PROBLEMS UNABLE TO TOLERATE ORAL MEDICATION, LOWER GASTROINTESTINAL PROBLEMS UNABLE TO ABSORB ORAL MEDICATIONS, TROUBLE REMEMBERING TO TAKE ORAL MEDICATIONS OR COORDINATING AN ORAL BISPHOSPHONATE WITH OTHER ORAL MEDICATIONS OR THEIR DAILY ROUTINE). 3) TRIAL OF, INTOLERANCE TO, OR A CONTRAINDICATION TO ONE BISPHOSPHONATE.

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ABATACEPT

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	RHEUMATOID ARTHRITIS (RA), PSORIATIC ARTHRITIS (PSA): 18 YEARS OR OLDER. JUVENILE PSORIATIC ARTHRITIS (JPSA): 2 YEARS OR OLDER.
Prescriber Restrictions	RHEUMATOID ARTHRITIS (RA), POLYARTICULAR COURSE JUVENILE IDIOPATHIC ARTHRITIS (PCJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA) AND JUVENILE PSORIATIC ARTHRITIS (JPSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST.
Coverage Duration	FOR aGVHD: 12 MONTHS. FOR RA, PJIA, PSA, JPSA: INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PJIA: TRIAL OF OR CONTRAINDICATION TO ONE DMARD. FOR ACUTE GRAFT VERSUS HOST DISEASE: ATTESTATION MEMBER IS TAKING IN COMBINATION WITH A CALCINEURIN INHIBITOR AND METHOTREXATE. RENEWAL: RA, PJIA, PSA, JPSA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	No

ABEMACICLIB

Products Affected

- VERZENIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	ADVANCED OR METASTATIC BREAST CANCER: THE PATIENT HAS NOT EXPERIENCED DISEASE PROGRESSION FOLLOWING PRIOR CDK INHIBITOR THERAPY.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ABIRATERONE

Products Affected

- *abiraterone*
- *abirtega*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ABIRATERONE SUBMICRONIZED

Products Affected

- YONSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: ONE OF THE FOLLOWING: 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ACALABRUTINIB

Products Affected

- CALQUENCE
- CALQUENCE (ACALABRUTINIB MAL)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ADAGRASIB

Products Affected

- KRAZATI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC): 1.) EVIDENCE OF KRAS G12C MUTATION AS DETERMINED BY AN APPROVED TEST 2.) PRIOR USE OF AT LEAST ONE SYSTEMIC THERAPY (E.G., ATEZOLIZUMAB, BEVACIZUMAB, CARBOPLATIN, CISPLATIN, NIVOLUMAB, PACLITAXEL, PEMETREXED, ETC.).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ADALIMUMAB

Products Affected

- HUMIRA PEN
- HUMIRA PEN CROHNS-UC-HS START
- HUMIRA PEN PSOR-UVEITS-ADOL HS
- HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML
- HUMIRA(CF) PEDI CROHNS STARTER SUBCUTANEOUS SYRINGE KIT 80 MG/0.8 ML, 80 MG/0.8 ML-40 MG/0.4 ML
- HUMIRA(CF) PEN CROHNS-UC-HS
- HUMIRA(CF) PEN PEDIATRIC UC
- HUMIRA(CF) PEN PSOR-UV-ADOL HS
- HUMIRA(CF) PEN SUBCUTANEOUS PEN INJECTOR KIT 40 MG/0.4 ML, 80 MG/0.8 ML
- HUMIRA(CF) SUBCUTANEOUS SYRINGE KIT 10 MG/0.1 ML, 20 MG/0.2 ML, 40 MG/0.4 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	RA, PJIA, AS: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSA: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CD, UC: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST. UVEITIS: PRESCRIBED BY OR IN CONSULTATION WITH AN OPTHALMOLOGIST. HS: PRESCRIBED BY OR IN CONSULATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): TRIAL OF OR CONTRAINDICATION TO ONE DMARD SUCH AS METHOTREXATE, LEFLUNOMIDE, OR SULFASALAZINE. ANKYLOSING SPONDYLITIS (AS): TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A CORTICOSTEROID (E.G., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE. UVEITIS: NO ISOLATED ANTERIOR UVEITIS. RENEWAL: RA, PJIA, PSA, AS, PSO, HIDRADENITIS SUPPURATIVA (HS), UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AFATINIB DIMALEATE

Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ALECTINIB

Products Affected

- ALECENSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ALPELISIB

Products Affected

- PIQRAY ORAL TABLET 200 MG/DAY (200 MG X 1), 250 MG/DAY (200 MG X1-50 MG X1), 300 MG/DAY (150 MG X 2)
- VIJOICE ORAL GRANULES IN PACKET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AMIKACIN

Products Affected

- *amikacin injection solution 1,000 mg/4 ml, 500 mg/2 ml*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DIAGNOSIS
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ANAKINRA

Products Affected

- KINERET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS (RA) (INITIAL): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	12 Months

PA Criteria	Criteria Details
Other Criteria	RHEUMATOID ARTHRITIS (RA) (INITIAL): ONE OF THE FOLLOWING: A) EITHER A TRIAL AND FAILURE, CONTRAINDICATION, OR INTOLERANCE (TF/C/I) TO TWO OF THE FOLLOWING: ENBREL (ETANERCEPT), HUMIRA (ADALIMUMAB), ORENCIA (ABATACEPT), RINVOQ (UPADACITINIB), XELJANZ/XELJANZ XR (TOFACITINIB), OR ATTESTATION DEMONSTRATING A TRIAL MAY BE INAPPROPRIATE, OR B) FOR CONTINUATION OF PRIOR THERAPY. NEONATAL-ONSET MULTISYSTEM INFLAMMATORY DISEASE (NOMID) (INITIAL): DIAGNOSIS OF NOMID HAS BEEN CONFIRMED BY ONE OF THE FOLLOWING: 1) NLRP-3 (NUCLEOTIDE-BINDING DOMAIN, LEUCINE RICH FAMILY (NLR), PYRIN DOMAIN CONTAINING 3) GENE (ALSO KNOWN AS COLD-INDUCED AUTO-INFLAMMATORY SYNDROME-1 [CIAS1]) MUTATION OR 2) BOTH OF THE FOLLOWING: A) TWO OF THE FOLLOWING CLINICAL SYMPTOMS: URTICARIA-LIKE RASH, COLD/STRESS TRIGGERED EPISODES, SENSORINEURAL HEARING LOSS, MUSCULOSKELETAL SYMPTOMS (E.G., ARTHRALGIA, ARTHRITIS, MYALGIA), CHRONIC ASEPTIC MENINGITIS, OR SKELETAL ABNORMALITIES (E.G., EPIPHYSEAL OVERGROWTH, FRONTAL BOSSING) AND B) ELEVATED ACUTE PHASE REACTANTS (EG, ERYTHROCYTE SEDIMENTATION RATE [ESR], C-REACTIVE PROTEIN [CRP], SERUM AMYLOID A [SAA]). DEFICIENCY OF INTERLEUKIN-1 RECEPTOR ANTAGONIST (DIRA): DIAGNOSIS OF DIRA. RENEWAL: RA, NOMID: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

APALUTAMIDE

Products Affected

- ERLEADA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	INITIAL: NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC): PATIENT HAS HIGH RISK PROSTATE CANCER (I.E., RAPIDLY INCREASING PROSTATE SPECIFIC ANTIGEN [PSA] LEVELS). NMCRPC OR METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC): 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. RENEWAL: DIAGNOSIS OF NMCRPC OR MCSPC.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

APOMORPHINE

Products Affected

- *apomorphine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	RENEWAL: PATIENT HAD IMPROVEMENT WITH MOTOR FLUCTUATIONS DURING OFF EPISODES WITH THE USE OF APOMORPHINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

APOMORPHINE - SL

Products Affected

- KYNMOBI SUBLINGUAL FILM 10 MG, 15 MG, 20 MG, 25 MG, 30 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OR OLDER
Prescriber Restrictions	PARKINSONS DISEASE (PD): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PD: PHYSICIAN HAS OPTIMIZED DRUG THERAPY FOR PARKINSONS DISEASE. RENEWAL: PD: PATIENT IMPROVEMENT WITH MOTOR FLUCTUATIONS DURING OFF EPISODES WITH THE USE OF KYNMOBI.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

APREMILAST

Products Affected

- OTEZLA
- OTEZLA STARTER
- OTEZLA XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 2 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION. FOR BEHCETS DISEASE: NO ADDITIONAL MEDICAL INFORMATION IS REQUIRED.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ARIKAYCE

Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY, OR IN CONSULTATION WITH, AN INFECTIOUS DISEASE SPECIALIST OR PULMONOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ASCIMINIB

Products Affected

- SCEMBLIX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ATOGEPANT

Products Affected

- QULIPTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PREVIOUS TRIAL OF OR CONTRAINDICATION TO ONE ALTERNATIVE FOR PREVENTIVE MIGRAINE TREATMENT: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL. RENEWAL FOR MIGRAINES: THE PATIENT HAS EXPERIENCED A REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY OF AT LEAST 2 DAYS PER MONTH OR A REDUCTION IN MIGRAINE SEVERITY OR MIGRAINE DURATION WITH QULIPTA THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AVACOPAN

Products Affected

- TAVNEOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	POSITIVE TEST FOR ANTI-PR3 OR ANTI-MPO (PROTEINASE 3 OR MYELOPEROXIDASE ANTIBODIES) OR POSITIVE TISSUE BIOPSY.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	COVERED FOR PATIENTS WITH CLINICAL DIAGNOSIS OF ANCA VASCULITIS GPA OR MPA, OR ANCA-POSITIVE VASCULITIS IN COMBINATION WITH STANDARD THERAPY, INCLUDING GLUCOCORTICOIDS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AVAPRITINIB

Products Affected

- AYVAKIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AVATROMBOPAG

Products Affected

- DOPTELET (10 TAB PACK)
- DOPTELET (15 TAB PACK)
- DOPTELET (30 TAB PACK)
- DOPTELET SPRINKLE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CHRONIC LIVER DISEASE (CLD): PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, GASTROENTEROLOGIST, HEPATOLOGIST, IMMUNOLOGIST, ENDOCRINOLOGIST, OR A SURGEON. CHRONIC IMMUNE THROMBOCYTOPENIA (ITP): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
Coverage Duration	CLD: 1 MONTH. CHRONIC ITP: INITIAL: 2 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	CLD: INITIAL: PATIENT HAS A PLANNED PROCEDURE 10 TO 13 DAYS AFTER INITIATION OF DOPTELET. PATIENT IS NOT RECEIVING OTHER THROMBOPOIETIN RECEPTOR AGONISTS (E.G., ROMIPLOSTIM, ELTROMBOPAG, ETC.). CHRONIC ITP: INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING: CORTICOSTEROIDS OR IMMUNOGLOBULINS, OR INSUFFICIENT RESPONSE TO SPLENECTOMY. RENEWAL: PATIENT HAD A CLINICAL RESPONSE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AVUTOMETINIB DEFACTINIB

Products Affected

- AVMAPKI-FAKZYNJA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AXATILIMAB-CSFR

Products Affected

- NIKTIMVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: A) TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CHRONIC GVHD). 2) DOCUMENT: I) FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS (E.G., PREDNISONE, METHOTREXATE, CYCLOSPORINE, TACROLIMUS, MYCOPHENOLATE, BELUMOSUDIL, IBRUTINIB, RUXOLITINIB, IF AVAILABLE), AND II) ACTUAL BODY WEIGHT OF AT LEAST 40KG.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, ONCOLOGIST, OR TRANSPLANT SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AXITINIB

Products Affected

- INLYTA ORAL TABLET 1 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AZACITIDINE

Products Affected

- ONUREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

AZTREONAM LYSINE

Products Affected

- CAYSTON

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	AT LEAST 7 YEARS OLD
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEDAQUILINE FUMARATE

Products Affected

- SIRTURO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 WEEKS
Other Criteria	SIRTURO USED IN COMBINATION WITH AT LEAST 3 OTHER ANTIBIOTICS FOR THE TREATMENT OF PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BELIMUMAB

Products Affected

- BENLYSTA INTRAVENOUS
- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SYSTEMIC LUPUS ERYTHEMATOSUS (SLE): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. LUPUS NEPHRITIS (LN): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: SLE: CURRENTLY TAKING CORTICOSTEROIDS, ANTIMALARIALS, NSAIDS, OR IMMUNOSUPPRESSIVE AGENTS. RENEWAL: SLE: PATIENT HAD CLINICAL IMPROVEMENT. LN: CLINICAL IMPROVEMENT IN RENAL RESPONSE COMPARED TO BASELINE OR CLINICAL PARAMETERS (E.G., FLUID RETENTION, USE OF RESCUE DRUGS, GLUCOCORTICOID DOSE).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BELUMOSUDIL MESYLATE

Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BELZUTIFAN

Products Affected

- WELIREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEMPEDOIC ACID

Products Affected

- NEXLETOL
- NEXLIZET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) Diagnosis: A) Treatment of heterozygous familial hypercholesterolemia (HeFH), or B) Treatment of atherosclerotic cardiovascular disease. 2) Document (only for first prescription): A) Patient is using concurrent LDL lowering therapies or prescriber attestation patient is unable to take recommended statin therapy or other LDL lowering therapies, and B) Lipid panel results (baseline LDL-C level must be greater than 70 mg/dL).
Age Restrictions	18 years of age or older
Prescriber Restrictions	1) Cardiologist, 2) Endocrinologist, 3) Internist, 4) Lipid Disorders Specialist or 5) Vascular Surgeon
Coverage Duration	12 MONTHS
Other Criteria	1) Atherosclerotic cardiovascular disease (CVD) can be considered as: acute coronary syndromes (ACS), stroke, myocardial infarction, transient ischemic attack, stable or unstable angina, peripheral arterial disease, coronary or arterial revascularization, or myocardial revascularization procedures (CABG or PCI). 2) FDA recommends avoiding concomitant use of simvastatin in doses greater than 20 mg, and pravastatin in doses greater than 40 mg.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BENRALIZUMAB

Products Affected

- FASENRA PEN
- FASENRA SUBCUTANEOUS SYRINGE 30 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ASTHMA: BLOOD EOSINOPHIL LEVEL GREATER THAN OR EQUAL TO 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
Age Restrictions	
Prescriber Restrictions	ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE.
Coverage Duration	12 MONTHS

PA Criteria	Criteria Details
Other Criteria	ASTHMA: INITIAL: 1) PRIOR THERAPY WITH A MEDIUM, HIGH-DOSE, OR MAXIMALLY-TOLERATED DOSE OF AN INHALED CORTICOSTEROID AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 2) ONE OF THE FOLLOWING: (A) PATIENT EXPERIENCED AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) PATIENT HAS POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NOT CONCURRENTLY RECEIVING XOLAIR, DUPIXENT OR OTHER ANTI-IL5 BIOLOGICS WHEN THESE ARE USED FOR THE TREATMENT OF ASTHMA. RENEWAL: CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: 1) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, 2) DECREASED UTILIZATION OF RESCUE MEDICATIONS, 3) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE, OR 4) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEROTRALSTAT

Products Affected

- ORLADEYO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEREDITARY ANGIOEDEMA (HAE): DIAGNOSIS CONFIRMED BY COMPLEMENT TESTING.
Age Restrictions	
Prescriber Restrictions	HAE: PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST, IMMUNOLOGIST, OR HEMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	HAE: INITIAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE, RENEWAL: IMPROVEMENT COMPARED TO BASELINE IN HAE ATTACKS (I.E., REDUCTIONS IN ATTACK FREQUENCY OR ATTACK SEVERITY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEVACIZUMAB

Products Affected

- AVASTIN INTRAVENOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINIDCATION TO ZIRABEV WHERE INDICATIONS ALIGN. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEVACIZUMAB-AWWB

Products Affected

- MVASI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINDICATION TO ZIRABEV WHERE INDICATIONS ALIGN. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEVACIZUMAB-BVZR

Products Affected

- ZIRABEV

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BEXAROTENE

Products Affected

- *bexarotene*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BINIMETINIB

Products Affected

- MEKTOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BOSUTINIB

Products Affected

- BOSULIF ORAL CAPSULE
- BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CHRONIC, ACCELERATED, OR BLAST PHASE PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA: BCR-ABL MUTATIONAL ANALYSIS CONFIRMING THAT T315I, V299L, G250E, OR F317L MUTATIONS ARE NOT PRESENT.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

BRIGATINIB

Products Affected

- ALUNBRIG ORAL TABLET 180 MG, 30 MG, 90 MG
- ALUNBRIG ORAL TABLETS,DOSE PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

C1 ESTERASE INHIBITOR-CINRYZE, BERINERT

Products Affected

- CINRYZE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEREDITARY ANGIOEDEMA (HAE): DIAGNOSIS CONFIRMED BY COMPLEMENT TESTING.
Age Restrictions	
Prescriber Restrictions	HAE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST, IMMUNOLOGIST, OR ALLERGIST.
Coverage Duration	12 MONTHS
Other Criteria	HAE: CINRYZE: INITIAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE. RENEWAL: IMPROVEMENT COMPARED TO BASELINE IN HAE ATTACKS (I.E., REDUCTIONS IN ATTACK FREQUENCY OR ATTACK SEVERITY). THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

C1 ESTERASE INHIBITOR-HAEGARDA, RUCONEST

Products Affected

- HAEGARDA SUBCUTANEOUS RECON SOLN 2,000 UNIT, 3,000 UNIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEREDITARY ANGIOEDEMA (HAE): DIAGNOSIS CONFIRMED BY COMPLEMENT TESTING.
Age Restrictions	
Prescriber Restrictions	HAE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST, IMMUNOLOGIST, OR ALLERGIST.
Coverage Duration	12 MONTHS
Other Criteria	HAE: HAEGARDA: INITIAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE. RENEWAL: IMPROVEMENT COMPARED TO BASELINE IN HAE ATTACKS (I.E., REDUCTIONS IN ATTACK FREQUENCY OR ATTACK SEVERITY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CABOZANTINIB

Products Affected

- COMETRIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CABOZANTINIB S-MALATE - CABOMETYX

Products Affected

- CABOMETYX ORAL TABLET 20 MG, 40 MG, 60 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CANNABIDIOL

Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	DRAVET SYNDROME (DS), LENNOX-GASTAUT SYNDROME (LGS), TUBEROUS SCLEROSIS COMPLEX (TSC): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: LENNOX-GASTAUT SYNDROME (LGS): TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING: CLOBAZAM, TOPIRAMATE, LAMOTRIGINE. RENEWAL: DS, LGS, TSC: CONFIRMATION OF DIAGNOSIS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CAPIVASERTIB

Products Affected

- TRUQAP

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS OF HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, LOCALLY ADVANCED OR METASTATIC BREAST CANCER: A) EVIDENCE OF PIK3CA/AKT1/PTEN MUTATIONS, B) PROGRESSION ON AT LEAST ONE ENDOCRINE-BASED REGIMEN IN THE METASTATIC SETTING OR RECURRENCE ON OR WITHIN 12 MONTHS IF USED AS ADJUVANT THERAPY, AND C) WILL BE USED IN COMBINATION WITH FULVESTRANT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CAPLACIZUMAB YHDP

Products Affected

- CABLIVI INJECTION KIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	CABLIVI WAS PREVIOUSLY INITIATED AS PART OF THE FDA APPROVED TREATMENT REGIMEN IN COMBINATION WITH PLASMA EXCHANGE AND IMMUNOSUPPRESSIVE THERAPY WITHIN AN INPATIENT SETTING. THE PATIENT HAS NOT EXPERIENCED MORE THAN TWO RECURRENCES OF ATTP WHILE ON CABLIVI THERAPY (I.E., NEW DROP IN PLATELET COUNT REQUIRING REPEAT PLASMA EXCHANGE DURING 30 DAYS POST-PLASMA EXCHANGE THERAPY [PEX] AND UP TO 28 DAYS OF EXTENDED THERAPY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CAPMATINIB

Products Affected

- TABRECTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CARFILZOMIB

Products Affected

- KYPROLIS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER.
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	INITIAL: 1) Diagnosis of Multiple Myeloma relapsed or refractory, as assessed by one of the following is the past 30 days: A) Serum M-protein greater than or equal to 0.5 g/dL, OR B) Urine M-protein greater than or equal to 200mg/24h, OR C) Serum free light chain (FLC) assay with involved FLC level greater than or equal to 10 mg/dL (100 mg/L) provided serum FLC ratio is abnormal, OR D) Member has progressive disease assessed within 60 days following the last dose of last anti-myeloma drug regimen received. 2) For Multiple Myeloma relapsed or refractory, it is used: A) In combination with dexamethasone or with lenalidomide plus dexamethasone in patients who have received one to three lines of therapy, or B) As a single agent in patients who have received one or more lines of therapy, OR C) In combination with Darzalex (daratumumab) or Darzalex Faspro (daratumumab/hyaluronidase-fihj) and dexamethasone in patients who have received one to three lines of therapy, OR D) In combination with Sarclisa (isatuximab-irfc) and dexamethasone in patients who have received one to three lines of therapy. 3) Request meets one of the following: A) Monotherapy: dose does not exceed 56 mg/m2 twice weekly each 28-day cycle, B) With dexamethasone and Revlimid: dose does not exceed 27 mg/m2 twice weekly 3 out of 4 weeks for twelve 28-day cycles, then 27 mg/m2 twice weekly 2 out of 4 weeks for the next six 28-day cycles for up to a total of 18 cycles, C) With dexamethasone with or without Darzalex dose does not exceed: i) 70 mg/m2 once weekly each 28-day cycle, OR ii) 56 mg/m2 twice weekly each 28-day cycle, D) With dexamethasone and Sarclisa: 56 mg/m2 twice weekly each 28-day cycle. 5) Prescribed carfilzomib (in combination with rituximab and dexamethasone) in the management of Waldenstrom macroglobulinemia in patients who have not previously received bortezomib or rituximab. RENEWAL: 1) Member is responding positively to therapy.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

CERITINIB

Products Affected

- ZYKADIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CLADRIBINE

Products Affected

- MAVENCLAD (10 TABLET PACK)
- MAVENCLAD (4 TABLET PACK)
- MAVENCLAD (5 TABLET PACK)
- MAVENCLAD (6 TABLET PACK)
- MAVENCLAD (7 TABLET PACK)
- MAVENCLAD (8 TABLET PACK)
- MAVENCLAD (9 TABLET PACK)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION THAT THE PATIENT HAS DEMONSTRATED CLINICAL BENEFIT COMPARED TO PRE TREATMENT BASELINE AND THE PATIENT DOES NOT HAVE LYMPHOPENIA.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	48 WEEKS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CLOBAZAM

Products Affected

- *clobazam oral suspension*
- *clobazam oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINDICATION TO LAMOTRIGINE OR TOPIRAMATE. REQUESTS FOR ORAL SUSPENSION APPROVABLE IF PATIENT IS UNABLE TO SWALLOW OR IS UNDER THE AGE OF 5 YEARS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CLOBAZAM-SYMPAZAN

Products Affected

- SYMPAZAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	LENNOX-GASTAUT SYNDROME (LGS): 1) PATIENT IS UNABLE TO TAKE TABLETS OR SUSPENSION AND 2) TRIAL OF OR CONTRAINDICATION TO A FORMULARY CLOBAZAM AGENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

COBIMETINIB FUMARATE

Products Affected

- COTELLIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CORTICOTROPIN

Products Affected

- ACTHAR
- ACTHAR SELFJECT
- CORTROPHIN GEL

PA Criteria	Criteria Details
Exclusion Criteria	NOT APPROVED FOR DIAGNOSTIC PURPOSES.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MULTIPLE SCLEROSIS (MS): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, ALLERGIST/IMMUNOLOGIST, OPHTHALMOLOGIST, PULMONOLOGIST OR NEPHROLOGIST.
Coverage Duration	INFANTILE SPASMS AND MS: 28 DAYS. OTHER FDA APPROVED INDICATIONS: INITIAL AND RENEWAL: 28 DAYS
Other Criteria	INITIAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS: TRIAL OF OR CONTRAINDICATION TO INTRAVENOUS (IV) CORTICOSTEROIDS. RENEWAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MULTIPLE SCLEROSIS: 1) DEMONSTRATED CLINICAL BENEFIT WHILE ON THERAPY AS INDICATED BY SYMPTOM RESOLUTION AND/OR NORMALIZATION OF LABORATORY TESTS, AND 2) CONTINUES TO POSSESS CONTRAINDICATION TO IV CORTICOSTEROIDS. PART B BEFORE PART D STEP THERAPY, APPLIES ONLY TO BENEFICIARIES IN AN MA-PD PLAN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

CRIZOTINIB

Products Affected

- XALKORI ORAL CAPSULE
- XALKORI ORAL PELLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

CYSTEAMINE HYDROCHLORIDE

Products Affected

- CYSTARAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DABRAFENIB MESYLATE

Products Affected

- TAFINLAR ORAL CAPSULE
- TAFINLAR ORAL TABLET FOR SUSPENSION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DACOMITINIB

Products Affected

- VIZIMPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DALFAMPRIDINE

Products Affected

- *dalfampridine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	MULTIPLE SCLEROSIS (MM): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MM: INITIAL: WALKING DISABILITY SUCH AS MILD TO MODERATE BILATERAL LOWER EXTREMITY WEAKNESS OR UNILATERAL WEAKNESS PLUS LOWER EXTREMITY OR TRUNCAL ATAXIA. RENEWAL: IMPROVEMENT IN WALKING ABILITY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DANZITEN

Products Affected

- DANZITEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DARATUMUMAB

Products Affected

- DARZALEX
- DARZALEX FASPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 Years of age or older
Prescriber Restrictions	Prescribed by or in consultation with an oncologist or hematologist.
Coverage Duration	INTIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: Diagnosis of 1) newly diagnosed Multiple myeloma and one of the following: A) Ineligible for ASCT and in combination with one of the following: i) lenalidomide and dexamethasone OR ii) bortezomib, melphalan, and prednisone. B) Eligible for ASCT in combination with one of the following: i) bortzeomib, thalidomide, and dexamethasone OR ii) bortezomib, lenalidomide, and dexamethasone. 2) Relapsed/refractory Multiple myeloma and one of the following: A) In combination with dexamethasone and either lenalidomide, or bortezomib, or carfilzomib after 1 or more prior therapy. B) In combination with pomalidomide and dexamethasone after 2 or more prior therapies (only Darzalex), including both of the following (i and ii): i) An immunomodulatory agent (e.g. thalidomide, lenalidomide), and ii) A PI (e.g., ixazomib, bortezomib, carfilzomib). 3) Diagnosis of systemic light chain amyloidosis and one of the following: A) Darzalex Faspro in combination with bortezomib, cyclophosphamide, and dexamethasone. RENEWAL: Member is responding positively to therapy.
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	Yes

DAROLUTAMIDE

Products Affected

- NUBEQA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC): PATIENT HAS HIGH RISK PROSTATE CANCER (I.E., RAPIDLY INCREASING PROSTATE SPECIFIC ANTIGEN [PSA] LEVELS) AND ONE OF THE FOLLOWING: 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. METASTATIC HORMONE-SENSITIVE PROSTATE CANCER (MHSPC): 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. RENEWAL: DIAGNOSIS OF NMCRPC OR MHSPC.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DASATINIB

Products Affected

- *dasatinib oral tablet 100 mg, 140 mg, 20 mg, 50 mg, 70 mg, 80 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PREVIOUSLY-TREATED CHRONIC MYELOID LEUKEMIA (CML) REQUIRES BCR-ABL MUTATIONAL ANALYSIS NEGATIVE FOR THE FOLLOWING MUTATIONS: T315I, V299L, T315A, F317L/V/I/C.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DATOPOTAMAB DERUXTECAN

Products Affected

- DATROWAY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: A) UNRESECTABLE OR METASTATIC, HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE (IHC 0 , IHC 1+ OR ICH 2+/ISH-) BREAST CANCER. 2) DOCUMENT: A) FOR BREAST CANCER: I) PREVIOUS TREATMENT WITH ENDOCRINE-BASED THERAPY, II) PREVIOUS TREATMENT WITH AT LEAST ONE ADDITIONAL SYSTEMIC THERAPY FOR UNRESECTABLE OR METASTATIC DISEASE, III) POSITIVE HORMONE RECEPTOR (HR), AND IV) HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE (IHC 0, IHC 1+ OR IHC 2+/ISH-) BREAST CANCER.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DECITABINE/CEDAZURIDINE

Products Affected

- INQOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DEFERASIROX

Products Affected

- *deferasirox*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST OR HEMATOLOGIST/ONCOLOGIST
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS INITIAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 1000 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS). RENEWAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 500 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS). NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) INITIAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS) AND LIVER IRON CONCENTRATION (LIC) OF 5 MG FE/G DRY WEIGHT OR GREATER. RENEWAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS) OR LIC OF 3 MG FE/G DRY WEIGHT OR GREATER. INITIAL FOR ALL INDICATIONS: FORMULARY VERSION OF DEFERASIROX SPRINKLE: TRIAL OF OR CONTRAINDICATION TO A GENERIC EQUIVALENT OF EITHER EXJADE TABLET FOR ORAL SUSPENSION OR A FORMULARY VERSION OF DEFERASIROX TABLET.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

DEFERIPRONE

Products Affected

- *deferiprone*
- FERRIPROX ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST OR HEMATOLOGIST/ONCOLOGIST
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL CRITERIA: TRANSFUSIONAL IRON OVERLOAD DUE TO THALASSEMIA SYNDROMES: (1) TRIAL OF OR CONTRAINDICATION TO A FORMULARY PREFERRED VERSION OF DEFERASIROX OR DEFEROXAMINE, AND (2) ONE OF THE FOLLOWING CRITERIA: A) PATIENT IS EXPERIENCING INTOLERABLE TOXICITIES OR CLINICALLY SIGNIFICANT ADVERSE EFFECTS OR HAS A CONTRAINDICATION TO THESE THERAPIES, OR B) INADEQUATE CHELATION DEFINED BY ONE OF THE FOLLOWING: I) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 2500 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS), OR II) EVIDENCE OF CARDIAC IRON ACCUMULATION (I.E., CARDIAC T2 STAR MRI LESS THAN 10 MILLISECONDS, IRON INDUCED CARDIOMYOPATHY, FALL IN LEFT VENTRICULAR EJECTION FRACTION, ARRHYTHMIA INDICATING INADEQUATE CHELATION). TRANSFUSIONAL IRON OVERLOAD DUE TO SICKLE CELL DISEASE OR OTHER ANEMIAS: TRIAL OF OR CONTRAINDICATION TO A FORMULARY PREFERRED VERSION OF DEFERASIROX OR DEFEROXAMINE. RENEWAL: SERUM FERRITIN LEVELS MUST BE CONSISTENTLY ABOVE 500MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DEFLAZACORT

Products Affected

- *deflazacort oral suspension*
- *deflazacort oral tablet 18 mg, 30 mg, 36 mg, 6 mg*
- EMFLAZA ORAL SUSPENSION
- EMFLAZA ORAL TABLET 18 MG, 30 MG, 36 MG, 6 MG
- *jaythari oral tablet 18 mg, 30 mg, 36 mg, 6 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DUCHENNE MUSCULAR DYSTROPHY (DMD): DIAGNOSIS COMFIRMED BY GENETIC TESTING.
Age Restrictions	
Prescriber Restrictions	DMD: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	DMD: INITIAL: TRIAL OF PREDNISONE OR PREDNISOLONE AND PATIENT MEETS ONE OF THE FOLLOWING: 1) REQUEST DUE TO ADVERSE EFFECTS OF PREDNISONE OR PREDNISOLONE OR 2) REQUEST DUE TO LACK OF EFFICACY OF PREDNISONE OR PREDNISOLONE AND ALL OF THE FOLLOWING: A) PATIENT IS NOT IN STAGE 1 (PRE-SYMPTOMATIC PHASE), B) STEROID MYOPATHY HAS BEEN RULED OUT. RENEWAL: PATIENT HAS MAINTAINED OR DEMONSTRATED A LESS THAN EXPECTED DECLINE IN AMBULATORY ABILITY IN MUSCLE FUNCTION ASSESSMENTS OR OTHER MUSCLE FUNCTION (I.E., PULMONARY OR CARDIAC FUNCTION).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DEGARELIX ACETATE

Products Affected

- FIRMAGON KIT W DILUENT SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DELAFLOXACIN

Products Affected

- BAXDELA ORAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ONE MONTH

PA Criteria	Criteria Details
Other Criteria	ACUTE BACTERIAL SKIN OR SKIN STRUCTURE INFECTION (ABSSSI): ONE OF THE FOLLOWING: 1) PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST, OR 2) ANTIMICROBIAL SUSCEPTIBILITY TESTING SHOWS SUSCEPTIBILITY TO DELAFLOXACIN AND RESISTANCE TO ONE STANDARD OF CARE AGENT FOR ABSSSI (E.G., SULFAMETHOXAZOLE/TRIMETHOPRIM, LEVOFLOXACIN, CLINDAMYCIN, CEPHALEXIN, OR VANCOMYCIN), OR 3) IF SENSITIVITY RESULTS ARE UNAVAILABLE: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED FORMULARY AGENTS FOR ABSSSI: A PENICILLIN, A FLUOROQUINOLONE, A CEPHALOSPORIN, OR A GRAM POSITIVE TARGETING ANTIBIOTIC. COMMUNITY-ACQUIRED BACTERIAL PNEUMONIA (CABP): ONE OF THE FOLLOWING: 1) PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST, OR 2) ANTIMICROBIAL SUSCEPTIBILITY TESTING SHOWS SUSCEPTIBILITY TO DELAFLOXACIN AND RESISTANCE TO AT LEAST TWO STANDARD OF CARE AGENTS FOR CABP (E.G., MACROLIDE, DOXYCYCLINE, LEVOFLOXACIN/MOXIFLOXACIN, BETA-LACTAM, LINEZOLID), OR 3) IF SENSITIVITY RESULTS ARE UNAVAILABLE: TRIAL OF OR CONTRAINDICATION TO AT LEAST TWO STANDARD OF CARE AGENTS FOR CABP (E.G., MACROLIDE, DOXYCYCLINE, LEVOFLOXACIN/MOXIFLOXACIN, BETA-LACTAM, LINEZOLID).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DENOSUMAB-XGEVA

Products Affected

- XGEVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DEUTETRABENAZINE

Products Affected

- AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG
- AUSTEDO XR ORAL TABLET EXTENDED RELEASE 24 HR 12 MG, 18 MG, 24 MG, 30 MG, 36 MG, 42 MG, 48 MG, 6 MG
- AUSTEDO XR TITRATION KT(WK1-4) ORAL TABLET, EXT REL 24HR DOSE PACK 12-18-24-30 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HUNTINGTON DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST. TARDIVE DYSKINESIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR MOVEMENT DISORDER SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	TARDIVE DYSKINESIA: PATIENT HAS A HISTORY OF USING AGENTS THAT CAUSE TARDIVE DYSKINESIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DICHLORPHENAMIDE

Products Affected

- *dichlorphenamide*
- KEVEYIS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 2 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PATIENT DOES NOT HAVE HEPATIC INSUFFICIENCY, PULMONARY OBSTRUCTION, OR A HEALTH CONDITION THAT WARRANTS CONCURRENT USE OF HIGH-DOSE ASPIRIN. RENEWAL: IMPROVEMENT IN SYMPTOMS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DICLOFENAC EPOLAMINE

Products Affected

- *diclofenac epolamine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

DICLOFENAC TOPICAL

Products Affected

- *diclofenac sodium topical gel 3 %*
- *diclofenac sodium topical solution in metered-dose pump*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	DICLOFENAC 2% TOPICAL SOLUTION: TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF DICLOFENAC SODIUM 1% TOPICAL GEL AND A FORMULARY VERSION OF DICLOFENAC SODIUM 1.5% TOPICAL DROPS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DIMETHYL FUMARATE

Products Affected

- *dimethyl fumarate oral capsule,delayed release(dr/ec) 120 mg, 120 mg (14)- 240 mg (46), 240 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DIROXIMEL FUMARATE

Products Affected

- VUMERITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DORDAVIPRONE

Products Affected

- MODEYSO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DRONABINOL

Products Affected

- *dronabinol*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	B VS D COVERAGE CONSIDERATION. PART D COVERAGE CONSIDERATION FOR A DIAGNOSIS OF NAUSEA AND VOMITING ASSOCIATED WITH CANCER CHEMOTHERAPY REQUIRES A TRIAL OF OR CONTRAINDICATION TO CONVENTIONAL ANTIEMETIC THERAPIES. NO ADDITIONAL REQUIREMENTS FOR A DIAGNOSIS OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DROXIDOPA

Products Affected

- *droxidopa*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	BLOOD PRESSURE READINGS WHILE THE PATIENT IS SITTING AND ALSO WITHIN 3 MINUTES OF STANDING FROM A SUPINE (LYING FACE UP) POSITION AT BASELINE AND RENEWAL.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST OR CARDIOLOGIST.
Coverage Duration	INITIAL: 3 MONTHS RENEWAL: 12 MONTHS
Other Criteria	INITIAL: DIAGNOSIS OF ORTHOSTATIC HYPOTENSION AS DOCUMENTED BY A DECREASE OF AT LEAST 20 MMHG IN SYSTOLIC BLOOD PRESSURE OR 10 MMHG DIASTOLIC BLOOD PRESSURE WITHIN THREE MINUTES AFTER STANDING FROM A SITTING POSITION. RENEWAL: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

DUPILUMAB

Products Affected

- DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 200 MG/1.14 ML, 300 MG/2 ML
- DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 100 MG/0.67 ML, 200 MG/1.14 ML, 300 MG/2 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: EOSINOPHILIC ASTHMA: BLOOD EOSINOPHIL LEVEL GREATER THAN OR EQUAL TO 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
Age Restrictions	
Prescriber Restrictions	ATOPIC DERMATITIS: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE. CHRONIC RHINOSINUSITIS WITH NASAL POLYPOSIS (CRSWNP): PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. PN: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: ATOPIC DERM, CRSWNP, EE, PN: 6 MOS, ASTHMA, COPD: 4 MOS. RENEWAL: 12 MOS (ALL INDICATIONS).

PA Criteria	Criteria Details
Other Criteria	INITIAL: ATOPIC DERMATITIS: 1) ATOPIC DERMATITIS INVOLVING AT LEAST 10% OF BODY SURFACE AREA (BSA) OR ATOPIC DERMATITIS AFFECTING THE FACE, HEAD, NECK, HANDS, FEET, GROIN, OR INTERTRIGINOUS AREAS, 2) INTRACTABLE PRURITUS OR CRACKING/OOZING/BLEEDING OF AFFECTED SKIN, AND 3) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING: TOPICAL CORTICOSTEROIDS, TOPICAL CALCINEURIN INHIBITORS, OR TOPICAL PDE4 INHIBITORS. ASTHMA: 1) PRIOR THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY-TOLERATED DOSE OF AN INHALED CORTICOSTEROID AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, 2) ONE OF THE FOLLOWING: (A) PATIENT EXPERIENCED AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) PATIENT HAS POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NOT CONCURRENTLY RECEIVING XOLAIR OR OTHER ANTI-IL5 BIOLOGICS WHEN THESE ARE USED FOR THE TREATMENT OF ASTHMA. CRSWNP: 1) EVIDENCE OF NASAL POLYPS BY DIRECT EXAMINATION, ENDOSCOPY OR SINUS CT SCAN, 2) INADEQUATELY CONTROLLED DISEASE AS DETERMINED BY THE USE OF SYSTEMIC STEROIDS IN THE PAST 2 YEARS OR ENDOSCOPIC SINUS SURGERY, AND 3) PRESCRIBED AS AN ADD-ON THERAPY TO CURRENT THERAPY. PN: DOCUMENT PRIOR FAILURE OR CONTRAINDICATIONS TO MEDIUM TO HIGH POTENCY TOPICAL CORTICOSTEROIDS. RENEWAL: ATOPIC DERMATITIS, CRSWNP: IMPROVEMENT WHILE ON THERAPY.
	ASTHMA: CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: 1) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, 2) DECREASED UTILIZATION OF RESCUE MEDICATIONS, 3) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE, OR 4) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

DUVELISIB

Products Affected

- COPIKTRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

EFGARTIGIMOD

Products Affected

- VYVGART
- VYVGART HYTRULO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	GENERALIZED MYASTHENIA GRAVIS (GMG): 1) POSITIVE SEROLOGICAL TEST FOR ANTI-ACETYLCHOLINE RECEPTOR (ANTI-ACHR) ANTIBODIES, 2) DOCUMENTATION THAT INDIVIDUAL HAS A MYASTHENIA GRAVIS ACTIVITIES OF DAILY LIVING (MG-ADL) SCORE OF AT LEAST 5 OR HIGHER.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	GMG: 1) TRIAL AND INADEQUATE RESPONSE AFTER 3 MONTHS OR INTOLERANCE TO AN ACETYLCHOLINESTERASE INHIBITOR (PYRIDOSTIGMINE BROMIDE), IS ON A STABLE DOSE OF AN ACETYLCHOLINESTERASE INHIBITOR, OR CONTRAINDICATION TO ACETYLCHOLINESTERASE INHIBITORS. 2) TRIAL AND INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE IMMUNOSUPPRESSIVE AGENTS (INCLUDING BUT NOT LIMITED TO SYSTEMIC CORTICOSTEROIDS (PREDNISONE, PREDNISOLONE) FOR AT LEAST 3 MONTHS OR NON-STEROIDAL IMMUNOSUPPRESSANTS (AZATHIOPRINE, MYCOPHENOLATE) FOR NO MORE THAN 90 DAYS, IS ON A STABLE DOSE OF ONE OR MORE IMMUNOSUPPRESSIVE AGENTS (INCLUDING BUT NOT LIMITED TO SYSTEMIC CORTICOSTEROIDS OR NON-STEROIDAL IMMUNOSUPPRESSANTS), OR CONTRAINDICATION TO SYSTEMIC CORTICOSTEROIDS AND NON-STEROIDAL IMMUNOSUPPRESSANTS. RENEWAL: GMG: 1) REDUCTION IN SIGNS OR SYMPTOMS THAT IMPACT DAILY FUNCTION. 2) DOCUMENTATION IS PROVIDED TO SHOW IMPROVEMENT IN SYPMTOM SEVERITY AND DAILY FUNCTIONING COMPARED TO PRE-TREATMENT. 3) CONTINUED TREATMENT REQUIRED TO MAINTAIN RESPONSE OR TO REGAIN CLINICALLY MEANINGFUL RESPONSE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

EFLAPEGRASTIM

Products Affected

- ROLVEDON

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

EFLORNITHINE

Products Affected

- IWILFIN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 Months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ELACESTRANT

Products Affected

- ORSERDU

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THE PATIENT HAS NOT EXPERIENCED DISEASE PROGRESSION FOLLOWING PRIOR CDK INHIBITOR THERAPY
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ELAGOLIX SODIUM

Products Affected

- ORILISSA ORAL TABLET 150 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: PREVIOUS TRIAL OF OR CONTRAINDICATION TO NSAID AND PROGESTIN-CONTAINING PREPARATION. RENEWAL: MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: IMPROVEMENT IN PAIN ASSOCIATED WITH ENDOMETRIOSIS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ELEXACAFTOR-TEZACAFTOR-IVACAFTOR

Products Affected

- TRIKAFTA ORAL GRANULES IN PACKET, SEQUENTIAL
- TRIKAFTA ORAL TABLETS, SEQUENTIAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CYSTIC FIBROSIS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME.
Other Criteria	RENEWAL: MAINTAINED, IMPROVED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN FEV1 OR BODY MASS INDEX (BMI), OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ELIGLUSTAT TARTRATE

Products Affected

- CERDELGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ELTROMBOPAG

Products Affected

- *eltrombopag olamine oral powder in packet 12.5 mg, 25 mg*
- *eltrombopag olamine oral tablet 12.5 mg, 25 mg, 50 mg, 75 mg*
- PROMACTA ORAL POWDER IN PACKET 12.5 MG, 25 MG
- PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	ITP: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST
Coverage Duration	ITP: INITIAL: 2 MO. RENEW: 12 MO. HCV: 12 MO. SEVERE APLASTIC ANEMIA: 12 MO.
Other Criteria	INITIAL: CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA PURPURA (ITP): TRIAL OF OR CONTRAINDICATION TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR AN INSUFFICIENT RESPONSE TO SPLENECTOMY. ALL INDICATIONS: APPROVAL FOR PROMACTA ORAL SUSPENSION PACKETS REQUIRES A TRIAL OF PROMACTA TABLETS OR PATIENT IS UNABLE TO TAKE TABLET FORMULATION. ITP: RENEWAL: PATIENT HAS SHOWN A CLINICAL RESPONSE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ENASIDENIB

Products Affected

- IDHIFA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ENCORAFENIB

Products Affected

- BRAFTOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ENDOTHELIN RECEPTOR ANTAGONISTS

Products Affected

- *ambrisentan* SUSPENSION
- OPSUMIT
- TRACLEER ORAL TABLET
- TRACLEER ORAL TABLET FOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS II-IV SYMPTOMS
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	INITIAL: MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) 3 WOOD UNITS OR GREATER. AMBRISENTAN: PATIENT DOES NOT HAVE IDIOPATHIC PULMONARY FIBROSIS (IPF). FORMULARY VERSION OF BOSENTAN: PATIENT DOES NOT HAVE ELEVATED LIVER ENZYMES (ALT, AST) MORE THAN 3 TIMES UPPER LIMIT OF NORMAL (ULN) OR INCREASES IN BILIRUBIN BY 2 OR MORE TIMES ULN. PATIENT IS NOT CONCURRENTLY TAKING CYCLOSPORINE A OR GLYBURIDE. RENEWAL: PATIENT SHOW IMPROVEMENT FROM BASELINE IN THE 6-MINUTE WALK DISTANCE OR PATIENT HAS A STABLE 6-MINUTE WALK DISTANCE WITH A STABLE/IMPROVED WHO FUNCTIONAL CLASS.
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	No

ENTRECTINIB

Products Affected

- ROZLYTREK ORAL CAPSULE 100 MG, 200 MG
- ROZLYTREK ORAL PELLETS IN PACKET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ENZALUTAMIDE

Products Affected

- XTANDI ORAL CAPSULE
- XTANDI ORAL TABLET 40 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS

PA Criteria	Criteria Details
Other Criteria	INITIAL: 1.) DIAGNOSIS: A) METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (mCRPC), B) NON-METASTATIC CASTRATION RESISTANT PROSTATE CANCER (nmCRPC), C) METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (mCSPC), OR D) NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (nmCSPC) WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS. 2) FOR MCRPC, NMCRPC AND MCSPC DOCUMENT: PATIENT IS RECEIVING A GONADOTROPIN-RELEASING HORMONE (GNRH) ANALOG CONCURRENTLY (E.G., LEUPROLIDE, GOSERELIN, TRIPTORELIN, OR HISTRELIN), HAD A BILATERAL ORCHIECTOMY, OR CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL). 3) PATIENTS WITH NMCSPC WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS MAY BE TREATED WITH OR WITHOUT A GNRH ANALOG. RENEWAL: DIAGNOSIS: A) METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (mCRPC), B) NON-METASTATIC CASTRATION RESISTANT PROSTATE CANCER (nmCRPC), C) METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (mCSPC), OR D) NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (nmCSPC) WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ERDAFITINIB

Products Affected

- BALVERSA ORAL TABLET 3 MG, 4 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ERENUMAB-AOOE

Products Affected

- AIMOVIG AUTOINJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	MIGRAINE PREVENTION: INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE ALTERNATIVE FOR PREVENTIVE MIGRAINE TREATMENT: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL, RENEWAL: EXPERIENCED A REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY OF AT LEAST 2 DAYS PER MONTH OR A REDUCTION IN MIGRAINE SEVERITY OR MIGRAINE DURATION WITH AIMOVIG THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ERLOTINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ERYTHROPOIESIS STIMULATING AGENTS - RETACRIT

Products Affected

- RETACRIT INJECTION SOLUTION 10,000 UNIT/ML, 2,000 UNIT/ML, 20,000 UNIT/2 ML, 20,000 UNIT/ML, 3,000 UNIT/ML, 4,000 UNIT/ML, 40,000 UNIT/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CHRONIC KIDNEY DISEASE (CKD), ANEMIA RELATED TO ZIDOVUDINE THERAPY, OR CANCER CHEMOTHERAPY: HEMOGLOBIN LEVEL OF LESS THAN 10G/DL. ELECTIVE NON-CARDIAC OR NON-VASCULAR SURGERY: HEMOGLOBIN LEVEL LESS THAN 13G/DL. RENEWAL: CKD: 1) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR 2) HEMOGLOBIN LEVEL HAS REACHED 10G/DL AND DOSE REDUCTION/INTERRUPTION IS REQUIRED TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS. ANEMIA RELATED TO ZIDOVUDINE THERAPY: HEMOGLOBIN LEVEL BETWEEN 10G/DL AND 12G/DL. CANCER CHEMOTHERAPY: HEMOGLOBIN LEVEL OF LESS THAN 10 G/DL OR THAT THE HEMOGLOBIN LEVEL DOES NOT EXCEED A LEVEL NEEDED TO AVOID RBC TRANSFUSION.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ANEMIA FROM CHEMO/CKD WITHOUT DIALYSIS/ZIDOVUDINE:12 MONTHS. SURGERY:1 MONTH.
Other Criteria	RENEWAL: CKD: NOT RECEIVING DIALYSIS TREATMENT. THIS DRUG MAY BE EITHER BUNDLED WITH AND COVERED UNDER END STAGE RENAL DISEASE DIALYSIS RELATED SERVICES OR COVERED UNDER MEDICARE D DEPENDING UPON THE CIRCUMSTANCES. INFORMATION MAY NEED TO BE SUBMITTED DESCRIBING THE USE AND SETTING OF THE DRUG TO MAKE THE DETERMINATION.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

ETANERCEPT

Products Affected

- ENBREL MINI
- ENBREL SUBCUTANEOUS RECON SOLN
- ENBREL SUBCUTANEOUS SOLUTION
- ENBREL SUBCUTANEOUS SYRINGE 25 MG/0.5 ML (0.5), 50 MG/ML (1 ML)
- ENBREL SURECLICK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE OR GENITAL AREA.
Age Restrictions	RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), PSORIATIC ARTHRITIS (PSA): 18 YEARS OR OLDER. JUVENILE PSORIATIC ARTHRITIS (JPSA): 2 YEARS OR OLDER.
Prescriber Restrictions	RA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), AS: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSA AND JPSA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PJIA: TRIAL OF OR CONTRAINDICATION TO ONE DMARD. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. RENEWAL: RA, PJIA, PSA, AS, PSO, JPSA: CONTINUES TO BENEFIT FROM THE MEDICATION.

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

EVEROLIMUS

Products Affected

- *everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *everolimus (antineoplastic) oral tablet for suspension*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED RENAL CELL CARCINOMA (RCC): TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF SUNITINIB OR SORAFENIB.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FECAL MICROBIOTA

Products Affected

- VOWST

PA Criteria	Criteria Details
Exclusion Criteria	ACTIVE CDI TREATMENT OR FIRST CDI OCCURRENCE.
Required Medical Information	1) DIAGNOSIS: PREVENTION OF RECURRENCE OF CLOSTRIDIOIDES DIFFICILE INFECTION (CDI) 2) DOCUMENT: A) DIAGNOSIS OF AT LEAST GREATER THAN OR EQUAL TO 2 RECURRENT EPISODE OF CDI [RECURRENT CDI IS DEFINED AS THE RECURRENCE OF DIARRHEA AND A CONFIRMATORY POSITIVE CDI TEST WITHIN 8 WEEKS AFTER TREATMENT OF AN INITIAL EPISODE (GREATER THAN OR EQUAL TO 3 TOTAL CDI EPISODES)], B) PRESCRIBER DOCUMENTATION INDICATING ANTIBIOTIC REGIMEN THE PATIENT IS RECEIVING AND THAT THE CURRENT EPISODE OF CDI IS CONTROLLED (LESS THAN 3 UNFORMED/LOOSE STOOLS/DAY FOR 2 CONSECUTIVE DAYS), C) POSITIVE STOOL TEST FOR C. DIFFICILE WITHIN 30 DAYS BEFORE PRIOR AUTHORIZATION REQUEST.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, INTERNIST, OR INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	ONE TIME APPROVAL
Other Criteria	ADMINISTRATION WILL OCCUR 48-72 HOURS FOLLOWING COMPLETION OF ANTIBIOTIC COURSE FOR CDI TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FEDRATINIB

Products Affected

- INREBIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: MYELOFIBROSIS: TRIAL OF OR CONTRAINDICATION TO FORMULARY VERSION OF JAKAFI (RUXOLITINIB). RENEWAL: MYELOFIBROSIS: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FENFLURAMINE

Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	RENEWAL: PATIENT HAS SHOWN CONTINUED CLINICAL BENEFIT (E.G. REDUCTION OF SEIZURES, REDUCED LENGTH OF SEIZURES, SEIZURE CONTROL MAINTAINED)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FENTANYL TRANSMUCOSAL AGENTS - FENTANYL CITRATE

Products Affected

- *fentanyl citrate buccal lozenge on a handle*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CANCER RELATED PAIN: CURRENTLY ON A MAINTENANCE DOSE OF CONTROLLED-RELEASE OPIOID PAIN MEDICATION. EITHER A TRIAL OR CONTRAINDICATION TO AT LEAST ONE IMMEDIATE-RELEASE ORAL OPIOID PAIN AGENT OR MEMBER HAS DIFFICULTY SWALLOWING TABLETS/CAPSULES. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FEZOLINETANT

Products Affected

- VEOZAH

PA Criteria	Criteria Details
Exclusion Criteria	A) KNOWN CIRRHOSIS, B) SEVERE RENAL IMPAIRMENT OR END-STAGE RENAL DISEASE, C) CONCOMITANT USE WITH CYP1A2 INHIBITORS
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS DUE TO MENOPAUSE: A) DOCUMENTATION OF CONTRAINDICATION, INTOLERANCE, OR INADEQUATE RESPONSE TO GREATER THAN OR EQUAL TO 1 MENOPAUSAL HORMONAL TREATMENTS (I.E PREMARIN, ESTRADIOL, DUAVEE).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FILGRASTIM

Products Affected

- NIVESTYM
- ZARXIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	NIVESTYM IS THE PREFERRED FILGRASTIM PRODUCT. REQUESTS FOR NIVESTYM DOES NOT REQUIRE A STEP. OTHER FORMULARY VERSIONS OF FILGRASTIM PRODUCTS WILL REQUIRE A TRIAL OF OR CONTRAINDICATION TO NIVESTYM, WHERE INDICATIONS ALIGN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FINGOLIMOD

Products Affected

- *fingolimod*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FLUTAMIDE

Products Affected

- EULEXIN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FOSTAMATINIB

Products Affected

- TAVALISSE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF A CLINICAL RESPONSE.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FRUQUINTINIB

Products Affected

- FRUZAQLA ORAL CAPSULE 1 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS OF METASTATIC COLORECTAL CANCER (MCRC): A) PRIOR TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-CONTAINING CHEMOTHERAPY, AND ANTI-VEGF THERAPY (E.G., BEVACIZUMAB), AND IF RAS WILD TYPE, AN ANTI-EGFR THERAPY (E.G. ERBITUX, VECTIBIX)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

FUTIBATINIB

Products Affected

- LYTGOBI ORAL TABLET 12 MG/DAY (4 MG X 3), 16 MG/DAY (4 MG X 4), 20 MG/DAY (4 MG X 5)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC ICCA HARBORING FGFR2 GENE FUSIONS OR OTHER REARRANGEMENTS: 1) FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT 2) PREVIOUSLY USED THERAPIES (E.G GEMCITABINE AND CISPLATIN OR DURVALUMAB, GEMCITABINE AND CISPLATIN) 3) DISEASE IS UNRESECTABLE AND LOCALLY ADVANCED OR METASTATIC.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GALCANEZUMAB-GNLM

Products Affected

- EMGALITY PEN
- EMGALITY SYRINGE SUBCUTANEOUS SYRINGE 120 MG/ML, 300 MG/3 ML (100 MG/ML X 3)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: MIGRAINES: 6 MOS. CLUSTER HEADACHE: 3 MOS. RENEWAL (ALL INDICATIONS): 12 MONTHS.
Other Criteria	INITIAL FOR MIGRAINES: PREVIOUS TRIAL OF OR CONTRAINDICATION TO ONE ALTERNATIVE FOR PREVENTIVE MIGRAINE TREATMENT: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL. CLUSTER HEADACHE: NO STEP. RENEWAL FOR MIGRAINES: THE PATIENT HAS EXPERIENCED A REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY OF AT LEAST 2 DAYS PER MONTH OR A REDUCTION IN MIGRAINE SEVERITY OR MIGRAINE DURATION WITH EMGALITY THERAPY. RENEWAL FOR EPISODIC CLUSTER HEADACHE: IMPROVEMENT IN EPISODIC CLUSTER HEADACHE FREQUENCY AS COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GANAXOLONE

Products Affected

- ZTALMY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DIAGNOSIS: TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE 5 (CDKL5) DEFICIENCY DISORDER (CDD). DOCUMENTATION SHOWING GENETIC TESTING CONFIRMING CDLK5 DEFICIENCY
Age Restrictions	2 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULATION BY A NEUROLOGIST, OR GENETICIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GEFITINIB

Products Affected

- *gefitinib*
- IRESSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GILTERITINIB

Products Affected

- XOSPATA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GLASDEGIB

Products Affected

- DAURISMO ORAL TABLET 100 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GLATIRAMER ACETATE

Products Affected

- *glatiramer subcutaneous syringe 20 mg/ml, 40 mg/ml*
- *glatopa subcutaneous syringe 20 mg/ml, 40 mg/ml*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GLECAPREVIR/PIBRENTASVIR

Products Affected

- MAVYRET ORAL PELLETS IN PACKET
- MAVYRET ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (HEPATOLOGIST), OR A SPECIALLY TRAINED GROUP SUCH AS ECHO (EXTENSION FOR COMMUNITY HEALTHCARE OUTCOMES) MODEL.
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. HCV RNA LEVEL WITHIN PAST 6 MONTHS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GLUCAGONLIKE PEPTIDE 1 RECEPTOR AGONIST

Products Affected

- MOUNJARO
- OZEMPIC SUBCUTANEOUS PEN INJECTOR 0.25 MG OR 0.5 MG (2 MG/3 ML), 0.25 MG OR 0.5 MG(2 MG/1.5 ML), 1 MG/DOSE (4 MG/3 ML), 2 MG/DOSE (8 MG/3 ML)
- RYBELSUS
- TRULICITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GLYCEROL PHENYLBUTYRATE

Products Affected

- RAVICTI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: UREA CYCLE DISORDER (UCD): DIAGNOSIS IS CONFIRMED BY ENZYMATIC, BIOCHEMICAL OR GENETIC TESTING
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	INITIAL: UREA CYCLE DISORDER (UCD): TRIAL OF OR CONTRAINDICATION TO SODIUM PHENYLBUTYRATE (BUPHENYL). RENEWAL: UCD: PATIENT HAS CLINICAL BENEFIT FROM BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

GUSELKUMAB

Products Affected

- TREMFYA INTRAVENOUS
- TREMFYA ONE-PRESS
- TREMFYA PEN INDUCTION PK(2PEN)
- TREMFYA PEN SUBCUTANEOUS PEN INJECTOR 200 MG/2 ML
- TREMFYA SUBCUTANEOUS SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. CD, UC: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: TRIAL OF OR CONTRAINDICATION ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. CROHNS DISEASE (CD): TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A CORTICOSTEROID (E.G., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE. RENEWAL: PSO, PSA, CD, UC: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

HIGH CONCENTRATION ORAL OPIOID SOLUTIONS

Products Affected

- *morphine concentrate oral solution*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	OPIOID TOLERANT: 12 MONTHS. HOSPICE, PALLIATIVE CARE OR END OF LIFE CARE: LIFETIME.
Other Criteria	1) OPIOID TOLERANT (I.E. PREVIOUS USE OF 60 MG ORAL MORPHINE PER DAY, 25 MCG TRANSDERMAL FENTANYL PER HOUR, 30 MG ORAL OXYCODONE PER DAY, 8 MG ORAL HYDROMORPHONE PER DAY, 25 MG ORAL OXYMORPHONE PER DAY, 60 MG ORAL HYDROCODONE PER DAY, OR AN EQUIANALGESIC DOSE OF ANOTHER OPIOID) AND HAS TROUBLE SWALLOWING OPIOID TABLETS, CAPSULES, OR LARGE VOLUMES OF LIQUID, OR 2) ENROLLED IN HOSPICE OR PALLIATIVE CARE OR END OF LIFE CARE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

HIGH RISK MEDICATIONS

Products Affected

- *benztropine oral*
- *cyclobenzaprine oral tablet 10 mg, 5 mg*
- *cyproheptadine oral syrup*
- *dicyclomine oral capsule*
- *dicyclomine oral solution*
- *dicyclomine oral tablet*
- *diphenoxylate-atropine oral tablet*
- *disopyramide phosphate oral capsule*
- *hydroxyzine hcl oral tablet*
- *promethazine oral tablet 12.5 mg*
- *scopolamine base*
- *trihexyphenidyl*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PROVIDER ACKNOWLEDGEMENT THAT MEDICATION IS A HRM IN THE ELDERLY AND THAT THE PATIENT HAS FAILED AND/OR TRIED AT LEAST ONE NON-HIGH RISK ALTERNATIVE.
Age Restrictions	PA APPLIES TO PATIENTS 65 YEARS OF AGE OR OLDER.
Prescriber Restrictions	
Coverage Duration	1) FOR CYCLOBENZAPRINE ONLY: 21 DAYS, RENEWALS: 21 DAYS. 2) ALL OTHER DRUGS: 12 MONTHS
Other Criteria	FOR CYCLOBENZAPRINE, DICYCLOMINE, DIGOXIN AND SCOPOLAMINE TRANSDERMAL: ONLY PROVIDER ATTESTATION OF RISKS AND BENEFITS OF THERAPY IS REQUIRED.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

HIGH RISK MEDICATIONS 2

Products Affected

- *paroxetine hcl oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PROVIDER ACKNOWLEDGEMENT THAT MEDICATION IS A HRM IN THE ELDERLY AND THAT THE PATIENT HAS FAILED AND/OR TRIED AT LEAST ONE NON-HIGH RISK ALTERNATIVE.
Age Restrictions	PA APPLIES TO PATIENTS 65 YEARS OF AGE OR OLDER.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

HRM BENZO

Products Affected

- *alprazolam oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg*
- *chlordiazepoxide hcl*
- *clonazepam oral tablet 0.5 mg, 1 mg, 2 mg*
- *clonazepam oral tablet,disintegrating 0.125 mg, 0.25 mg, 0.5 mg, 1 mg, 2 mg*
- *clorazepate dipotassium*
- *diazepam intensol*
- *diazepam oral solution 5 mg/5 ml (1 mg/ml)*
- *diazepam oral tablet*
- *flurazepam*
- *lorazepam oral tablet 0.5 mg, 1 mg, 2 mg*
- *temazepam oral capsule 15 mg, 30 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Provider acknowledges that the patient might be using this medication in combination with an opioid and the patient has tried and failed at least one non-high-risk alternative. Patients with a diagnosis of cancer, Sickle-cell disease, or receiving hospice or palliative care are excluded from the above requirement.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IBRUTINIB

Products Affected

- IMBRUVICA ORAL CAPSULE 140 MG, 70 MG
- IMBRUVICA ORAL SUSPENSION
- IMBRUVICA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ICATIBANT

Products Affected

- *icatibant*
- *sajazir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN ALLERGIST/IMMUNOLOGIST OR HEMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS OF HEREDITARY ANGIOEDEMA CONFIRMED BY COMPLEMENT TESTING.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IDELALISIB

Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IMATINIB MESYLATE

Products Affected

- *imatinib oral tablet 100 mg, 400 mg*
- IMKELDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ADJUVANT GASTROINTESTINAL STROMAL TUMOR TREATMENT: 36 MONTHS. ALL OTHER DIAGNOSES: 12 MONTHS.
Other Criteria	PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA: PATIENT HAS NOT RECEIVED A PREVIOUS TREATMENT WITH ANOTHER TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

INAVOLISIB

Products Affected

- ITOVEBI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

INFIGRATINIB

Products Affected

- TRUSELTIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

INTERFERON GAMMA-1B

Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CHRONIC GRANULOMATOUS DISEASE (CGD): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST, INFECTIOUS DISEASE SPECIALIST, OR IMMUNOLOGIST. SEVERE MALIGNANT OSTEOPETROSIS (SMO): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN ENDOCRINOLOGIST OR HEMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	RENEWAL: THE PATIENT HAS DEMONSTRATED CLINICAL BENEFIT COMPARED TO BASELINE AND HAS NOT RECEIVED HEMATOPOIETIC CELL TRANSPLANTATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

INTERFERONS FOR MULTIPLE SCLEROSIS

Products Affected

- AVONEX INTRAMUSCULAR PEN INJECTOR KIT
- AVONEX INTRAMUSCULAR SYRINGE KIT
- BETASERON SUBCUTANEOUS KIT
- PLEGRIDY SUBCUTANEOUS PEN INJECTOR 125 MCG/0.5 ML
- PLEGRIDY SUBCUTANEOUS SYRINGE 125 MCG/0.5 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IVACAFTOR

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	HOMOZYGOUS FOR F508DEL MUTATION IN CFTR GENE.
Required Medical Information	CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CYSTIC FIBROSIS
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: LIFETIME
Other Criteria	RENEWAL: MAINTAINED, IMPROVED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN FEV1 OR BODY MASS INDEX (BMI), OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IVOSIDENIB

Products Affected

- TIBSOVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

IXAZOMIB

Products Affected

- NINLARO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LANADELUMAB

Products Affected

- TAKHZYRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEREDITARY ANGIOEDEMA (HAE): DIAGNOSIS CONFIRMED BY COMPLEMENT TESTING.
Age Restrictions	
Prescriber Restrictions	HAE: PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST/IMMUNOLOGIST OR HEMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	HAE: INITIAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE, RENEWAL: IMPROVEMENT COMPARED TO BASELINE IN HAE ATTACKS (I.E., REDUCTIONS IN ATTACK FREQUENCY OR ATTACK SEVERITY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LANREOTIDE

Products Affected

- *lanreotide*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: DIAGNOSIS OF ACROMEGALY: 1) INADEQUATE RESPONSE TO SURGERY OR RADIOTHERAPY, OR SURGERY OR RADIOTHERAPY IS NOT AN OPTION, 2) BASELINE GROWTH HORMONE (GH) AND IGF-I BLOOD LEVELS, 3) WILL NOT BE USED IN COMBINATION WITH ORAL OCTREOTIDE.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: DIAGNOSIS OF CARCINOID SYNDROME: 1) REDUCE THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY. DIAGNOSIS OF GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS): 1) DISEASE IS UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC, WELL OR MODERATELY DIFFERENTIATED. RENEWAL: ALL INDICATIONS: 1) IMPROVED DISEASE RESPONSE FROM BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LAPATINIB

Products Affected

- *lapatinib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LAROTRECTINIB

Products Affected

- VITRAKVI ORAL CAPSULE 100 MG, 25 MG
- VITRAKVI ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	APPROVAL FOR VITRAKVI ORAL SOLUTION: TRIAL OF VITRAKVI CAPSULES OR PATIENT IS UNABLE TO TAKE CAPSULE FORMULATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LAZERTINIB

Products Affected

- LAZCLUZE ORAL TABLET 240 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLSC), 2) DOCUMENT: CONFIRMATION OF EGFR EXON 19 DELETIONS OR EXON 21 L858R SUBSTITUTION MUTATION, 3) PATIENT MUST BE RECEIVING ANTICOAGULANT VTE PROPHYLAXIS FOR THE FIRST 4 MONTHS OF TREATMENT AND 4) PATIENT WILL BE USING LAZCLUZE (LAZERTINIB) IN COMBINATION WITH AMIVANTAMAB.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	1) VALIDATE THAT THE TREATMENT REGIMEN IS FOLLOWING THE MOST UP TO DATE NCCN GUIDELINES RECOMMENDATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LEDIPASVIR-SOFOSBUVIR

Products Affected

- *ledipasvir-sofosbuvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS.
Age Restrictions	
Prescriber Restrictions	GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (HEPATOLOGIST), OR A SPECIALLY TRAINED GROUP SUCH AS ECHO (EXTENSION FOR COMMUNITY HEALTHCARE OUTCOMES) MODEL.
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING: CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, ROSUVASTATIN, SOFOSBUVIR (AS A SINGLE AGENT), OR TIPRANAVIR/RITONAVIR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LENALIDOMIDE

Products Affected

- *lenalidomide*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LENVATINIB

Products Affected

- LENVIMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LETERMOVIR

Products Affected

- PREVYMIS ORAL PELLETS IN PACKET
  120 MG, 20 MG
- PREVYMIS ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	8 MONTHS
Other Criteria	CYTOMEGALOVIRUS (CMV) PROPHYLAXIS IN HEMATOPOIETIC CELL TRANSPLANT RECIPIENTS: THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 28 POST TRANSPLANTATION. CMV PROPHYLAXIS IN KIDNEY TRANSPLANT RECIPIENTS: THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 7 POST TRANSPLANTATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LEVODOPA

Products Affected

- INBRIJA INHALATION CAPSULE, W/INHALATION DEVICE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	PD: INITIAL: 1) PATIENT IS NOT CURRENTLY TAKING MORE THAN 1600MG OF LEVODOPA PER DAY. RENEWAL: PATIENT HAD IMPROVEMENT WITH MOTOR FLUCTUATIONS DURING OFF EPISODES WITH THE USE OF INBRIJA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

L-GLUTAMINE

Products Affected

- ENDARI
- *glutamine (sickle cell)*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SICKLE CELL DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST.
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: LIFETIME.
Other Criteria	INITIAL: PATIENTS 18 YEARS OR OLDER: ONE OF THE FOLLOWING: 1) AT LEAST 2 SICKLE CELL CRISES IN THE PAST YEAR, OR 2) SICKLE-CELL ASSOCIATED SYMPTOMS WHICH ARE INTERFERING WITH ACTIVITIES OF DAILY LIVING, OR 3) HISTORY OF OR HAS RECURRENT ACUTE CHEST SYNDROME. PATIENTS 5 TO 17 YEARS: APPROVED WITHOUT ADDITIONAL CRITERIA. RENEWAL: PATIENT HAS MAINTAINED OR EXPERIENCED REDUCTION IN ACUTE COMPLICATIONS OF SICKLE CELL DISEASE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LIDOCAINE

Products Affected

- *lidocaine hcl mucous membrane solution 4 % (40 mg/ml)*
- *lidocaine topical adhesive patch,medicated 5 %*
- *lidocaine topical ointment*
- *tridacaine ii*
- ZTLIDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

LIDOCAINE PRILOCAINE

Products Affected

- APRIZIO PAK II
- EMPRICAINE-II
- *lidocaine-prilocaine topical cream*
- NUVAKAAN-II
- PRIZOPAK II

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG MAY BE EITHER BUNDLED WITH AND COVERED UNDER END STAGE RENAL DISEASE DIALYSIS RELATED SERVICES OR COVERED UNDER MEDICARE D DEPENDING UPON THE CIRCUMSTANCES. INFORMATION MAY NEED TO BE SUBMITTED DESCRIBING THE USE AND SETTING OF THE DRUG TO MAKE THE DETERMINATION. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

LOMITAPIDE

Products Affected

- JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST, ENDOCRINOLOGIST OR LIPIDOLOGIST.
Coverage Duration	12 MONTHS

PA Criteria	Criteria Details
Other Criteria	1) DIAGNOSIS DETERMINED BY A) DEFINITE SIMON BROOME DIAGNOSTIC CRITERIA, OR B) DUTCH LIPID NETWORK CRITERIA SCORE OF 8 OR GREATER, OR C) CLINICAL DIAGNOSIS BASED ON A HISTORY OF AN UNTREATED LDL-C CONCENTRATION GREATER THAN 500 MG/DL TOGETHER WITH EITHER XANTHOMA BEFORE 10 YEARS OF AGE, OR EVIDENCE OF HEFH IN BOTH PARENTS. 2) LDL-C LEVEL GREATER THAN OR EQUAL TO 70MG/DL WHILE ON MAXIMAL DRUG TREATMENT. 3) TRIAL OF EVOLOCUMAB UNLESS THE PATIENT HAS NON-FUNCTIONING LDL RECEPTORS. 4) MEETS ONE OF THE FOLLOWING: A) TAKING A HIGH-INTENSITY STATIN (I.E., ATORVASTATIN 40-80MG DAILY, ROSUVASTATIN 20-40MG DAILY) FOR A DURATION OF AT LEAST 8 WEEKS, B) TAKING A MAXIMALLY TOLERATED DOSE OF ANY STATIN FOR A DURATION OF AT LEAST 8 WEEKS GIVEN THAT THE PATIENT CANNOT TOLERATE A HIGH-INTENSITY STATIN, C) ABSOLUTE CONTRAINDICATION TO STATIN THERAPY (E.G., ACTIVE DECOMPENSATED LIVER DISEASE, NURSING FEMALE, PREGNANCY OR PLANS TO BECOME PREGNANT, HYPERSENSITIVITY REACTIONS), D) STATIN INTOLERANCE, OR E) TRIAL OF ROSUVASTATIN, ATORVASTATIN, OR STATIN THERAPY AT ANY DOSE AND HAS EXPERIENCED SKELETAL-MUSCLE RELATED SYMPTOMS (E.G., MYOPATHY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LOMUSTINE

Products Affected

- GLEOSTINE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LORLATINIB

Products Affected

- LORBRENA ORAL TABLET 100 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LOTILANER

Products Affected

- XDEMVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH OPHTHALMOLOGIST OR OPTOMETRIST
Coverage Duration	6 WEEKS
Other Criteria	DIAGNOSIS OF DEMODEX BLEPHARITIS: A) PRESENCE OF ERYTHEMA IN THE UPPER EYELID MARGIN AND B) PRESENCE OF MITES IN EYELASHES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LUMACAFTOR-IVACAFTOR

Products Affected

- ORKAMBI ORAL GRANULES IN PACKET
- ORKAMBI ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CYSTIC FIBROSIS (CF): CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CF.
Age Restrictions	
Prescriber Restrictions	CF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CF EXPERT.
Coverage Duration	CF: INITIAL: 6 MONTHS, RENEWAL: LIFETIME.
Other Criteria	RENEWAL: CF: PATIENT MAINTAINED, IMPROVED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN FEV1 OR BODY MASS INDEX (BMI), OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

LUSUTROMBOPAG

Products Affected

- MULPLETA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, GASTROENTEROLOGIST, HEPATOLOGIST, IMMUNOLOGIST, ENDOCRINOLOGIST, OR A SURGEON.
Coverage Duration	1 MONTH
Other Criteria	1) PATIENT HAS A PLANNED PROCEDURE 8 TO 14 DAYS AFTER INITIATION OF MULPLETA AND 2) PATIENT IS NOT RECEIVING OTHER THROMBOPOIETIN RECEPTOR AGONISTS (E.G., AVATROMBOPAG, ROMIPLOSTIM, ELTROMBOPAG).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MARIBAVIR

Products Affected

- LIVTENCITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, ONCOLOGIST, INFECTIOUS DISEASE, OR TRANSPLANT SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MEPOLIZUMAB

Products Affected

- NUCALA SUBCUTANEOUS AUTO-INJECTOR
- NUCALA SUBCUTANEOUS RECON SOLN
- NUCALA SUBCUTANEOUS SYRINGE 100 MG/ML, 40 MG/0.4 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ASTHMA: BLOOD EOSINOPHIL LEVEL GREATER THAN OR EQUAL TO 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
Age Restrictions	
Prescriber Restrictions	ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN PULMONARY OR ALLERGY MEDICINE. NASAL POLYPS: PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST.
Coverage Duration	INITIAL: NASAL POLYPS: 6 MO. OTHER INDICATIONS: 12 MO. RENEWAL: NASAL POLYPS, ASTHMA: 12 MO.

PA Criteria	Criteria Details
Other Criteria	INITIAL: ASTHMA: 1) PRIOR THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 2) ONE OF THE FOLLOWING: (A) PATIENT EXPERIENCED AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) PATIENT HAS POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, AND 3) NOT CONCURRENTLY RECEIVING XOLAIR, DUPIXENT OR OTHER ANTI-IL5 BIOLOGICS WHEN THESE ARE USED FOR THE TREATMENT OF ASTHMA. NASAL POLYPS: PREVIOUS 8 WEEK TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID. RENEWAL: ASTHMA: CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: 1) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, 2) DECREASED UTILIZATION OF RESCUE MEDICATIONS, 3) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR 4) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE. NASAL POLYPS: CLINICAL BENEFIT COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MIDOSTAURIN

Products Affected

- RYDAPT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ACUTE MYELOID LEUKEMIA: 6 MONTHS. ADVANCED SYSTEMIC MASTOCYTOSIS: 12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MIFEPRISTONE

Products Affected

- *mifepristone oral tablet 300 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CUSHINGS SYNDROME (CS): DIAGNOSIS CONFIRMED BY ONE OF THE FOLLOWING: 1) 24-HR URINE FREE CORTISOL (2 OR MORE TESTS TO CONFIRM), 2) OVERNIGHT 1MG DEXAMETHASONE TEST, OR 3) LATE NIGHT SALIVARY CORTISOL (2 OR MORE TESTS TO CONFIRM).
Age Restrictions	
Prescriber Restrictions	CS: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	CD: INITIAL: HYPERCORTISOLISM IS NOT A RESULT OF CHRONIC GLUCOCORTICOIDS, RENEWAL: 1) PATIENT CONTINUES TO HAVE IMPROVEMENT OF GLUCOSE TOLERANCE OR STABLE GLUCOSE TOLERANCE (E.G., REDUCED A1C, IMPROVED FASTING GLUCOSE, ETC.), 2) PATIENT CONTINUES TO HAVE TOLERABILITY TO MIFEPRISTONE, AND 3) PATIENT CONTINUES TO NOT BE A CANDIDATE FOR SURGICAL TREATMENT OR HAS FAILED SURGERY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MIGALASTAT

Products Affected

- GALAFOLD

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: FABRY DISEASE: PATIENT IS SYMPTOMATIC OR HAS EVIDENCE OF INJURY FROM GL-3 TO THE KIDNEY, HEART, OR CENTRAL NERVOUS SYSTEM RECOGNIZED BY LABORATORY, HISTOLOGICAL, OR IMAGING FINDINGS.
Age Restrictions	
Prescriber Restrictions	FABRY DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST, CARDIOLOGIST, OR SPECIALIST IN GENETICS OR INHERITED METABOLIC DISORDERS.
Coverage Duration	INITIAL: 6 MOS. RENEWAL: 12 MOS.
Other Criteria	FABRY DISEASE: INITIAL: NOT CONCURRENTLY USING ENZYME REPLACEMENT THERAPY (I.E. FABRAZYME), RENEWAL: PATIENT HAS DEMONSTRATED IMPROVEMENT OR STABILIZATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MIGLUSTAT

Products Affected

- *miglustat*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MILTEFOSINE

Products Affected

- IMPAVIDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MIRDAMETINIB

Products Affected

- GOMEKLI ORAL CAPSULE 1 MG, 2 MG
- GOMEKLI ORAL TABLET FOR SUSPENSION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: TREATMENT OF NEUROFIBROMATOSIS TYPE 1 (NF1), 2) DOCUMENT: A) PATIENT HAS SYMPTOMATIC AND INOPERABLE PLEXIFORM NEUROFIBROMAS (PN), AND B) PATIENTS BODY SURFACE AREA (BSA) OR ACTUAL BODY WEIGHT AND HEIGHT.
Age Restrictions	2 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH GENETICIST, NEUROLOGIST, NEUROSURGEON, HEMATOLOGIST/ONCOLOGIST, OPHTHALMOLOGIST, OR ORTHOPEDIC SURGEON.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MITAPIVAT

Products Affected

- PYRUKYND

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DIAGNOSIS: TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY. FOR INITIAL EVALUATION DOCUMENT: MUTATION IN THE PKLR GENE (MUST HAVE AT LEAST 2 MUTANT ALLELES IN THE PKLR GENE, OF WHICH AT LEAST 1 WAS A MISSENSE MUTATION). CURRENT HEMOGLOBIN LEVEL (MUST BE LESS THAN OR EQUAL TO 10 MG/DL). FOR RENEWALS DOCUMENT: PATIENT HAS EXPERIENCED IMPROVEMENT FROM BASELINE OR REDUCTION IN TRANSFUSION BURDEN.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MOBOCERTINIB

Products Affected

- EXKIVITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MOMELOTINIB

Products Affected

- OJJAARA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

MOSUNETUZUMAB-AXGB

Products Affected

- LUNSUMIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL): 1) PRIOR USE OF AT LEAST TWO LINES OF SYSTEMIC THERAPY (E.G. RITUXIMAB, CYCLOPHOSPHAMIDE, VINCRISTINE, CORTICOSTEROIDS, LENALIDOMIDE, DUVELISIB, ETC.)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NERATINIB MALEATE

Products Affected

- NERLYNX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	EARLY-STAGE TUMOR (STAGE I-III) AND THE MEDICATION IS BEING REQUESTED WITHIN 2 YEARS OF COMPLETING THE LAST TRASTUZUMAB DOSE. ALL OTHER FDA APPROVED INDICATIONS ARE COVERED WITHOUT ADDITIONAL CRITERIA, EXCEPT THOSE CRITERIA IN THE FDA APPROVED LABEL.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NILOTINIB

Products Affected

- NILOTINIB D-TARTRATE ORAL CAPSULE 150 MG, 200 MG, 50 MG
- *nilotinib hcl oral capsule 150 mg, 200 mg, 50 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PREVIOUSLY TREATED CML REQUIRES BCR-ABL MUTATIONAL ANALYSIS NEGATIVE FOR THE FOLLOWING MUTATIONS: T315I, Y253H, E255K/V, F359V/C/I, OR G250E.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NINTEDANIB

Products Affected

- OFEV

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL: IDIOPATHIC PULMONARY FIBROSIS (IPF): NOT APPROVED FOR PATIENTS WITH OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS, SYSTEMIC SCLEROSIS, RHEUMATOID ARTHRITIS, RADIATION, SARCOIDOSIS, BRONCHIOLITIS OBLITERANS ORGANIZING PNEUMONIA, HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION, VIRAL HEPATITIS, OR CANCER). SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD): NOT APPROVED FOR PATIENTS WITH OTHER KNOWN CAUSES OF ILD [E.G., HEART FAILURE/FLUID OVERLOAD, DRUG-INDUCED LUNG TOXICITY (CYCLOPHOSPHAMIDE, METHOTREXATE, ACE-INHIBITORS), RECURRENT ASPIRATION (SUCH AS FROM GERD), PULMONARY VASCULAR DISEASE, PULMONARY EDEMA, PNEUMONIA, CHRONIC PULMONARY THROMBOEMBOLISM, ALVEOLAR HEMORRHAGE OR ILD CAUSED BY ANOTHER RHEUMATIC DISEASE, SUCH AS MIXED CONNECTIVE TISSUE DISEASE (MCTD)].
Required Medical Information	INITIAL: IPF: 1) A USUAL INTERSTITIAL PNEUMONIA (UIP) PATTERN AS EVIDENCED BY HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) ALONE OR VIA A COMBINATION OF SURGICAL LUNG BIOPSY AND HRCT, AND 2) BASELINE FORCED VITAL CAPACITY (FVC) AT LEAST 50% OF PREDICTED VALUE. SSC-ILD: AT LEAST 10% FIBROSIS ON A CHEST HRCT AND BASELINE FVC AT LEAST 40% OF PREDICTED VALUE. PF-ILD: AT LEAST 10% FIBROSIS ON A CHEST HRCT AND BASELINE FVC AT LEAST 45% OF PREDICTED VALUE.
Age Restrictions	INITIAL: IPF, SSC-ILD, PF-ILD: 18 YEARS OR OLDER.

PA Criteria	Criteria Details
Prescriber Restrictions	IPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST. SSC-ILD, PF-ILD: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: SSC-ILD: 6 MOS. IPF, PF-ILD: 12 MOS. RENEWAL: 12 MOS.
Other Criteria	INITIAL: IPF: TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: ESBRIET. SSC-ILD: TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: ACTEMRA SUBQ. PF-ILD: LUNG FUNCTION AND RESPIRATORY SYMPTOMS OR CHEST IMAGING HAVE WORSENED/PROGRESSED DESPITE TREATMENT WITH MEDICATIONS USED IN CLINICAL PRACTICE FOR ILD (NOT ATTRIBUTABLE TO COMORBIDITIES SUCH AS INFECTION, HEART FAILURE). RENEWAL: IPF, SSC-ILD, PF-ILD: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NIRAPARIB

Products Affected

- ZEJULA ORAL CAPSULE
- ZEJULA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER: ZEJULA WILL BE USED AS MONOTHERAPY AND IS STARTED NO LATER THAN 8 WEEKS AFTER THE MOST RECENT PLATINUM-CONTAINING REGIMEN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NIRAPARIB ABIRATERONE

Products Affected

- AKEEGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NIROGASESTAT

Products Affected

- OGSIVEO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS TREATMENT OF PROGRESSING DESMOID TUMORS (DTS): TUMOR PROGRESSION (E.G. A) IMAGING SCANS SUCH AS CT, MRI, OR ULTRASOUND, B) AT LEAST ONE LINE OF THERAPY SUCH AS SURGERY, RADIOTHERAPY, OR SYSTEMIC THERAPY OR C) PRESCRIBER DOCUMENTATION THAT STATES DISEASE PROGRESSION).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

NITISINONE

Products Affected

- *nitisinone*
- NITYR
- ORFADIN ORAL CAPSULE 20 MG
- ORFADIN ORAL SUSPENSION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEREDITARY TYROSINEMIA TYPE 1 (HT-1): DIAGNOSIS CONFIRMED BY ELEVATED URINARY OR PLASMA SUCCINYLACETONE LEVELS OR A MUTATION IN THE FUMARYLACETOACETATE HYDROLASE GENE.
Age Restrictions	
Prescriber Restrictions	HT-1: PRESCRIBED BY OR IN CONSULTATION WITH A PRESCRIBER SPECIALIZING IN INHERITED METABOLIC DISEASES.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	HT-1: INITIAL: ORFADIN SUSPENSION: TRIAL OF OR CONTRAINDICATION TO PREFERRED FORMULARY NITISINONE TABLETS OR CAPSULES, RENEWAL: URINARY OR PLASMA SUCCINYLACETONE LEVELS HAVE DECREASED FROM BASELINE WHILE ON TREATMENT WITH NITISINONE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OCRELIZUMAB

Products Affected

- OCREVUS
- OCREVUS ZUNOVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RELAPSING FORM OF MULTIPLE SCLEROSIS (MS): TRIAL OF TWO AGENTS INDICATED FOR THE TREATMENT OF RELAPSING FORMS OF MS. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OFATUMUMAB-SQ

Products Affected

- KESIMPTA PEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OLANZAPINE/SAMIDORPHAN

Products Affected

- LYBALVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SCHIZOPHRENIA/BIPOLAR I: PRESCRIBED BY OR IN CONSULTATION WITH A PSYCHIATRIST
Coverage Duration	12 MONTHS
Other Criteria	SCHIZOPHRENIA: (1) PATIENT IS AT HIGH RISK OF WEIGHT GAIN AND (2) TRIAL OF OR CONTRAINDICATION TO LATUDA OR ONE OF THE FOLLOWING ORAL ANTIPSYCHOTICS: RISPERIDONE, CLOZAPINE TABLET, OLANZAPINE, IMMEDIATE RELEASE QUETIAPINE FUMARATE, ZIPRASIDONE, ARIPIPRAZOLE. BIPOLAR I: (1) PATIENT IS AT HIGH RISK OF WEIGHT GAIN AND (2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING ORAL ANTIPSYCHOTICS: RISPERIDONE, OLANZAPINE, IMMEDIATE RELEASE QUETIAPINE FUMARATE, ZIPRASIDONE, ARIPIPRAZOLE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OLAPARIB

Products Affected

- LYNPARZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER: 1) MEDICATION WILL BE USED AS MONOTHERAPY, AND 2) PATIENT HAS COMPLETED TWO OR MORE LINES OF PLATINUM-BASED CHEMOTHERAPY. ADVANCED GERMLINE BRCA-MUTATED OVARIAN CANCER: 1) PATIENT HAS COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY. 2) MEDICATION WILL BE USED AS MONOTHERAPY. METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. ALL OTHER FDA APPROVED INDICATIONS ARE COVERED WITHOUT ADDITIONAL CRITERIA, EXCEPT THOSE CRITERIA IN THE FDA APPROVED LABEL.
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	No

OLUTASIDENIB

Products Affected

- REZLIDHIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE IDH1 MUTATION: 1.) DOCUMENTATION OF PRESENCE OF SUSCEPTIBLE IDH1 MUTATION AS DETERMINED BY AN FDA APPROVED TEST.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OMACETAXINE

Products Affected

- SYNRIBO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INDUCTION: 3 MONTHS. POST INDUCTION/RENEWAL: 3 TO 12 MONTHS.
Other Criteria	CML INDUCTION THERAPY: TRIAL OF OR CONTRAINDICATION TO AT LEAST TWO OF THE FOLLOWING AGENTS: GLEEVEC, SPRYCEL, TASIGNA, BOSULIF, OR ICLUSIG. APPROVAL FOR POST-INDUCTION THERAPY DURATION WILL DEPEND ON THE PATIENT'S HEMATOLOGIC RESPONSE, DEFINED AS (1) AN ABSOLUTE NEUTROPHIL COUNT (ANC) GREATER THAN OR EQUAL TO 1.5 X 10^9/L AND PLATELETS GREATER THAN OR EQUAL TO 100 X 10^9/L WITHOUT BLOOD BLASTS OR (2) THE PATIENT HAS BONE MARROW BLASTS AT LESS THAN 5 PERCENT. APPROVAL IS FOR 12 MONTHS IF HEMATOLOGIC RESPONSE IS MET. IF NOT MET, APPROVAL IS FOR 3 MONTHS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OMALIZUMAB

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ASTHMA: POSITIVE SKIN PRICK OR BLOOD TEST (E.G., ELISA, FEIA) TO A PERENNIAL AEROALLERGEN AND A BASELINE IGE SERUM LEVEL GREATER THAN OR EQUAL TO 30 IU/ML.
Age Restrictions	
Prescriber Restrictions	INITIAL AND RENEWAL: CHRONIC IDIOPATHIC URTICARIA (CIU): PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE, DERMATOLOGY OR IMMUNOLOGY. INITIAL: NASAL POLYPS: PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE.
Coverage Duration	INITIAL:ASTHMA: 12 MO. CIU, CRSsNP, IGE-FA: 6 MO. RENEWAL:ASTHMA, CRSsNP, IGE-FA: 12 MO. CIU: 6 MO.

PA Criteria	Criteria Details
Other Criteria	INITIAL: CIU: TRIAL OF OR CONTRAINDICATION TO A MAXIMALLY TOLERATED DOSE OF AN H1 ANTI-HISTAMINE AND STILL EXPERIENCES HIVES ON MOST DAYS OF THE WEEK. NASAL POLYPS: PREVIOUS 90 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID. ASTHMA: 1) PRIOR THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, 2) ONE OF THE FOLLOWING: (A) PATIENT EXPERIENCED AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING 3 OR MORE DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) PATIENT HAS POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA, 3) XOLAIR WILL BE USED AS ADD-ON MAINTENANCE TREATMENT, AND 4) NOT CONCURRENTLY RECEIVING DUPIXENT OR OTHER ANTI-IL5 BIOLOGICS WHEN THESE ARE USED FOR THE TREATMENT OF ASTHMA. RENEWAL: CIU: DIAGNOSIS OF CIU. NASAL POLYPS: CLINICAL BENEFIT COMPARED TO BASELINE. ASTHMA: CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: 1) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, 2) DECREASED UTILIZATION OF RESCUE MEDICATIONS, 3) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR 4) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

OSIMERTINIB

Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NSCLC WITH EGFR T790M MUTATION: PATIENT IS NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE-INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PACRITINIB

Products Affected

- VONJO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Initial: 1) Diagnosis of intermediate- or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis in patients, AND 2) Documentation of platelet count below 50 x 10^9/L
Age Restrictions	
Prescriber Restrictions	Prescribed by or in consultation with a hematologist or oncologist
Coverage Duration	Initial: 4 months Renewal: 12 months
Other Criteria	Myelofibrosis: Renewal: Continues to benefit from the medication
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PALBOCICLIB

Products Affected

- IBRANCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	THE PATIENT HAS NOT EXPERIENCED DISEASE PROGRESSION FOLLOWING PRIOR CDK INHIBITOR THERAPY
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PASIREOTIDE DIASPARTATE

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CUSHINGS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS.
Other Criteria	RENEWAL: CD: PATIENT CONTINUES TO HAVE IMPROVEMENT OF CD AND MAINTAINS TOLERABILITY TO SIGNIFOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PAZOPANIB

Products Affected

- *pazopanib*
- VOTRIENT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PCSK9

Products Affected

- PRALUENT PEN
- REPATHA PUSHTRONEX
- REPATHA SURECLICK
- REPATHA SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PDE5 INHIBITORS PAH

Products Affected

- *alyq*
- *sildenafil (pulm.hypertension) oral tablet*
- *tadalafil (pulm. hypertension)*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS II-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	PAH: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	PAH: INITIAL: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA), ANY ORGANIC NITRATES IN ANY FORM, OR GUANYLATE CYCLASE STIMULATORS, AND 2) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) 3 WOOD UNITS OR GREATER, RENEWAL: IMPROVEMENT FROM BASELINE IN THE 6-MINUTE WALK DISTANCE OR A STABLE 6-MINUTE WALK DISTANCE WITH A STABLE/IMPROVED WHO FUNCTIONAL CLASS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEGFILGRASTIM

Products Affected

- FULPHILA
- NEULASTA
- NYVEPRIA
- UDENYCA
- UDENYCA AUTOINJECTOR
- UDENYCA ONBODY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	NYVEPRIA IS THE PREFERRED PEGFILGRASTIM PRODUCT. REQUESTS FOR NYVEPRIA DOES NOT REQUIRE A STEP. OTHER FORMULARY VERSIONS OF PEGFILGRASTIM PRODUCTS WILL REQUIRE A TRIAL OF OR CONTRAINDICATION TO NYVEPRIA, WHERE INDICATIONS ALIGN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEGLOTICASE

Products Affected

- KRYSTEXXA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) Clinical chart notes show submission of laboratory values demonstrating baseline serum uric acid level greater than 6 mg/dL.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Initial: 12 MONTHS. Renewal: 12 MONTHS
Other Criteria	Initial: Chronic Gout: 1) Member has symptomatic gout as defined by one of the following: History of at least 2 gout flares in the previous 12 months, at least 1 gouty tophus, or chronic gouty arthropathy. 2) Member has a history of contraindication, intolerance, or treatment failure (i.e., failure to normalize uric acid to less than 6 mg/dL) after 3 months of therapy (at the maximally medically appropriate dose) with both of the following: allopurinol and febuxostat. 3) Prescriber attestation that serum uric acid levels will be monitored prior to each infusion and that discontinuation of treatment will be considered if preinfusion levels increase above 6 mg/dL. Renewal: 1) You are currently receiving Krystexxa. 2) Documentation of a positive clinical response. 3) Member had two consecutive uric acid levels above 6 mg/dL after initiating treatment.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

PEGVALIASE-PQPZ

Products Affected

- PALYNZIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	RENEWAL: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEGVISOMANT

Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEMBROLIZUMAB

Products Affected

- KEYTRUDA
- KEYTRUDA QLEX SUBCUTANEOUS SOLUTION 395 MG-4,800 UNIT/2.4 ML, 790 MG-9,600 UNIT/4.8 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEMETREXED

Products Affected

- AXTLE
- PEMRYDI RTU

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEMIGATINIB

Products Affected

- PEMAZYRE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PENICILLAMINE

Products Affected

- *penicillamine*
- THIOLA EC
- *tiopronin oral tablet,delayed release (dr/ec)*

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL AND RENEWAL: FORMULARY VERSION OF PENICILLAMINE: RHEUMATOID ARTHRITIS (RA): HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY.
Required Medical Information	INITIAL: WILSONS DISEASE: KNOWN FAMILY HISTORY OF WILSONS DISEASE OR PHYSICAL EXAMINATION CONSISTENT WITH WILSONS DISEASE. CONFIRMATION OF ONE OF THE FOLLOWING: 1) PLASMA COPPER-PROTEIN CERULOPLASMIN IS LESS THAN 20MG/DL, 2) LIVER BIOPSY POSITIVE FOR AN ABNORMALLY HIGH CONCENTRATION OF COPPER (GREATER THAN 250MCG/G DRY WEIGHT) OR THE PRESENCE OF KAYSER-FLEISCHER RINGS, OR 3) CONFIRMATION BY GENETIC TESTING FOR ATP7B MUTATIONS. CYSTINURIA: DIAGNOSIS REQUIRES THE PRESENCE OF NEPHROLITHIASIS AND ONE OR MORE OF THE FOLLOWING: 1) STONE ANALYSIS SHOWING PRESENCE OF CYSTEINE, 2) IDENTIFICATION OF PATHOGNOMONIC HEXAGONAL CYSTINE CRYSTALS ON URINALYSIS, OR 3) POSITIVE FAMILY HISTORY OF CYSTINURIA WITH POSITIVE CYANIDE-NITROPRUSSIDE SCREEN. RENEWAL: WILSONS DISEASE, CYSTINURIA: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION.
Age Restrictions	
Prescriber Restrictions	WILSONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST. CYSTINURIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST. RA: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 12 MONTHS, RENEWAL: LIFETIME.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RA, WILSONS DISEASE: REQUESTS FOR FORMULARY VERSION OF PENICILLAMINE CAPSULE REQUIRE A TRIAL OF OR CONTRAINDICATION TO PENICILLAMINE TABLET (DEPEN). CYSTINURIA: REQUESTS FOR FORMULARY VERSION OF PENICILLAMINE CAPSULE REQUIRES A TRIAL OF OR CONTRAINDICATION TO PENICILLAMINE TABLET (DEPEN) AND A FORMULARY VERSION OF TIOPRONIN (THIOLA)/THIOLA EC. RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. RENEWAL: RA: EXPERIENCED OR MAINTAINED IMPROVEMENT IN TENDER JOINT COUNT OR SWOLLEN JOINT COUNT COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PEXIDARTINIB

Products Affected

- TURALIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PIMAVANSERIN

Products Affected

- NUPLAZID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST, GERIATRICIAN, OR A BEHAVIORAL HEALTH SPECIALIST (SUCH AS A PSYCHIATRIST).
Coverage Duration	INITIAL 12 MONTHS. RENEWAL 12 MONTHS.
Other Criteria	RENEWAL REQUIRES THAT THE PATIENT HAS EXPERIENCED AN IMPROVEMENT IN PSYCHOSIS SYMPTOMS FROM BASELINE AND DEMONSTRATES A CONTINUED NEED FOR TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PIRFENIDONE

Products Affected

- *pirfenidone oral capsule*
- *pirfenidone oral tablet 267 mg, 801 mg*

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL: IDIOPATHIC PULMONARY FIBROSIS (IPF): PATIENTS WITH KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS, SYSTEMIC SCLEROSIS, RHEUMATOID ARTHRITIS, RADIATION, SARCOIDOSIS, BRONCHIOLITIS OBLITERANS ORGANIZING PNEUMONIA, HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION, VIRAL HEPATITIS, OR CANCER).
Required Medical Information	INITIAL: IPF: 1) A USUAL INTERSTITIAL PNEUMONIA (UIP) PATTERN AS EVIDENCED BY HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) ALONE OR VIA A COMBINATION OF SURGICAL LUNG BIOPSY AND HRCT, AND 2) BASELINE FORCED VITAL CAPACITY (FVC) AT LEAST 50% OF PREDICTED VALUE.
Age Restrictions	IPF: 18 YEARS OR OLDER.
Prescriber Restrictions	IPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST.
Coverage Duration	IPF: INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	RENEWAL: IPF: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PIRTOBRUTINIB

Products Affected

- JAYPIRCA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

POMALIDOMIDE

Products Affected

- POMALYST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PONATINIB

Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

POSACONAZOLE

Products Affected

- NOXAFIL ORAL SUSPENSION
- *posaconazole oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	OROPHARYNGEAL CANDIDIASIS (OPC): 3 MONTHS. PROPHYLAXIS: 6 MONTHS. TREATMENT: 12 WEEKS.
Other Criteria	POSACONAZOLE SUSPENSION ONLY: 1) OPC: TRIAL OF OR CONTRAINDICATION TO FLUCONAZOLE OR ITRACONAZOLE. 2) PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTION: INABILITY TO SWALLOW TABLETS. POSACONZOLE TABLETS ONLY: NO EXTRA CRITERIA REQUIRED, ALL FDA APPROVED INDICATION COVERED. CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE REQUIRES NO EXTRA CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PRALSETINIB

Products Affected

- GAVRETO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PRAMLINTIDE

Products Affected

- SYMLINPEN 120
- SYMLINPEN 60

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	TYPE I OR TYPE II DIABETES: REQUIRING INSULIN OR CONTINUOUS INSULIN INFUSION (INSULIN PUMP) FOR GLYCEMIC CONTROL
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PYRIMETHAMINE

Products Affected

- *pyrimethamine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	TOXOPLASMOSIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	TOXOPLASMOSIS: INITIAL: 8 WEEKS. RENEWAL: 6 MOS.
Other Criteria	RENEWAL: CONTINUED TREATMENT OF TOXOPLASMOSIS REQUIRES ONE OF THE FOLLOWING: 1) PERSISTENT CLINICAL DISEASE (HEADACHE, NEUROLOGICAL SYMPTOMS, OR FEVER) AND PERSISTENT RADIOGRAPHIC DISEASE (ONE OR MORE MASS LESIONS ON BRAIN IMAGING) OR 2) CD4 COUNT LESS THAN 200 CELLS/MM3 AND CURRENT ANTI-RETROVIRAL THERAPY IF HIV POSITIVE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

QUININE SULFATE

Products Affected

- *quinine sulfate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

QUIZARTINIB

Products Affected

- VANFLYTA

PA Criteria	Criteria Details
Exclusion Criteria	CONTRAINDICATED IN PATIENTS WITH SEVERE HYPOKALEMIA, SEVERE HYPOMAGNESEMIA, LONG QT SYNDROME, OR IN PATIENTS WITH A HISTORY OF VENTRICULAR ARRHYTHMIAS OR TORSADES DE POINTES.
Required Medical Information	DIAGNOSIS: FOR THE TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 INTERNAL TANDEM DUPLICATION (ITD)-POSITIVE: A) PRESCRIBED IN COMBINATION WITH STANDARD CYTARABINE AND ANTHRACYCLINE (I.E., DAUNORUBICIN) INDUCTION AND CYTARABINE CONSOLIDATION, AND AS MAINTENANCE MONOTHERAPY FOLLOWING CONSOLIDATION CHEMOTHERAPY, B) EVIDENCE OF POSITIVE FLT3 INTERNAL TANDEM DUPLICATION
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	LEUKOSTRAT CDX FLT3 MUTATION ASSAY IS THE FDA-APPROVED TEST FOR SELECTION OF PATIENTS WITH AML FOR VANFLYTA TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

REGORAFENIB

Products Affected

- STIVARGA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RELUGOLIX

Products Affected

- ORGOVYX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

REPOTRECTINIB

Products Affected

- AUGTYRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	12 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	DIAGNOSIS OF LOCALLY ADVANCED OR METASTATIC ROS1-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC): DOCUMENTATION OF ROS1-MUTATED NSCLC. DIAGNOSIS OF SOLID TUMORS THAT NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION, ARE LOCALLY ADVANCED OR METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

REVUMENIB

Products Affected

- REVUFORJ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	DIAGNOSIS OF RELAPSED OR REFRACTORY ACUTE LEUKEMIA WITH A LYSINE METHYLTRANSFERASE 2A GENE (KMT2A) TRANSLOCATION
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RIABNI

Products Affected

- RIABNI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	RHEUMATOID ARTHRITIS (RA), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL), NON-HODGKIN'S LYMPHOMA (NHL): 18 YEARS OR OLDER. GRANULOMATOSIS WITH POLYANGIITIS (GPA) (WEGENER'S GRANULOMATOSIS), MICROSCOPIC POLYANGIITIS (MPA): 2 YEARS OR OLDER.
Prescriber Restrictions	RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. CHRONIC LYMPHOCYTIC LEUKEMIA (CLL), NON-HODGKIN'S LYMPHOMA (NHL): PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST.
Coverage Duration	FOR NHL, GPA, MPA: 12 MONTHS. FOR RA, CLL: INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	FOR GPA, MPA: CONCURRENT USE OF GLUCOCORTICOIDS. INITIAL RA: TRIAL OF OR CONTRAINDICATION TO AT LEAST 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG), SUCH AS METHOTREXATE DOSE GREATER THAN OR EQUAL TO 20MG PER WEEK OR MAXIMALLY TOLERATED DOSE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. FOR CLL: CONCURRRENT CHEMOTHERAPY (E.G., FLUDARABINE, CYCLOPHOSPHAMIDE). RENEWAL: RA: EXPERIENCED OR MAINTAINED A 20 PERCENT OR GREATER IMPROVEMENT IN TENDER JOINT COUNT OR SWOLLEN JOINT COUNT WHILE ON THERAPY.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

RIBOCICLIB

Products Affected

- KISQALI FEMARA CO-PACK ORAL TABLET 200 MG/DAY(200 MG X 1)-2.5 MG, 400 MG/DAY(200 MG X 2)-2.5 MG, 600 MG/DAY(200 MG X 3)-2.5 MG
- KISQALI ORAL TABLET 200 MG/DAY (200 MG X 1), 400 MG/DAY (200 MG X 2), 600 MG/DAY (200 MG X 3)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	THE PATIENT HAS NOT EXPERIENCED DISEASE PROGRESSION FOLLOWING PRIOR CDK INHIBITOR THERAPY
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	REQUIRES A TRIAL OF OR CONTRAINDICATION TO VERZENIO OR IBRANCE WHERE INDICATIONS ALIGN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RIFAXIMIN

Products Affected

- XIFAXAN ORAL TABLET 200 MG, 550 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	TRAVELERS DIARRHEA/HE: 12 MOS. IBS-D: 12 WKS.
Other Criteria	RIFAXIMIN 550 MG TABLETS: HE: TRIAL OF OR CONTRAINDICATION TO LACTULOSE OR CONCURRENT LACTULOSE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RIMEGEPANT

Products Affected

- NURTEC ODT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: ACUTE MIGRAINE TREATMENT: TRIAL OF OR CONTRAINDICATION TO ONE TRIPTAN (E.G., SUMATRIPTAN, RIZATRIPTAN). EPISODIC MIGRAINE PREVENTION: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREVENTIVE MIGRAINE TREATMENTS: DIVALPROEX SODIUM, TOPIRAMATE, PROPRANOLOL, TIMOLOL. RENEWAL: ACUTE MIGRAINE TREATMENT: 1) IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE, OR 2) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS. EPISODIC MIGRAINE PREVENTION: 1) REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY OF AT LEAST 2 DAYS PER MONTH, OR 2) REDUCTION IN MIGRAINE SEVERITY OR MIGRAINE DURATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RIOCIGUAT

Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PULMONARY ARTERIAL HYPERTENSION (PAH): 1) CONFIRMATORY DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION: A) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, B) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) 15 MMHG OR LESS, AND C) PULMONARY VASCULAR RESISTANCE (PVR) 3 WOOD UNITS OR GREATER, AND 2) NYHA-WHO FUNCTIONAL CLASS (FC) II-IV SYMPTOMS. PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH) WHO GROUP 4: NYHA-WHO FUNCTIONAL CLASS II-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	PAH, CTEPH: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PAH: NOT CONCURRENTLY TAKING NITRATES, NITRIC OXIDE DONORS, PHOSPHODIESTERASE INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS. CTEPH: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING NITRATES, NITRIC OXIDE DONORS, OR ANY PDE INHIBITORS, AND 2) NOT A CANDIDATE FOR SURGERY OR HAS INOPERABLE CTEPH OR HAS PERSISTENT OR RECURRENT DISEASE AFTER SURGICAL TREATMENT. RENEWAL: PAH, CTEPH: IMPROVEMENT FROM BASELINE IN THE 6-MINUTE WALK DISTANCE OR A STABLE 6-MINUTE WALK DISTANCE WITH A STABLE/IMPROVED WHO FUNCTIONAL CLASS.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

RIPRETINIB

Products Affected

- QINLOCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RISANKIZUMAB-RZAA

Products Affected

- SKYRIZI SUBCUTANEOUS PEN INJECTOR
- SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML, 75 MG/0.83 ML
- SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PLAQUE PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. RENEWAL: PSO, PSA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RISDIPLAM

Products Affected

- EVRYSDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	SMA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROMUSCULAR SPECIALIST OR NEUROLOGIST.
Coverage Duration	SMA: INITIAL/RENEWAL: 12 MONTHS
Other Criteria	SPINAL MUSCULAR ATROPHY (SMA): INITIAL: DOCUMENTATION OF GENE MUTATION ANALYSIS INDICATING MUTATIONS OR DELETIONS OF BOTH ALLELES OF THE SURVIVAL MOTOR NEURON 1 (SMN1) GENE. FOR PRESYMPTOMATIC PATIENTS: DOCUMENTATION OF UP TO THREE COPIES OF SURVIVAL MOTOR NEURON 2 (SMN2) BASED ON NEWBORN SCREENING. FOR SYMPTOMATIC PATIENTS: 1) DOCUMENTATION OF BASELINE MOTOR FUNCTION ASSESSMENT BY A NEUROMUSCULAR SPECIALIST OR SMA SPECIALIST, 2) IF PREVIOUSLY RECEIVED GENE THERAPY, THE PATIENT HAD LESS THAN EXPECTED CLINICAL BENEFIT. RENEWAL: IMPROVED, MAINTAINED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN MOTOR FUNCTION ASSESSMENTS COMPARED TO BASELINE, OR OTHER MUSCLE FUNCTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ROMOSOZUMAB

Products Affected

- EVENITY SUBCUTANEOUS SYRINGE 105 MG/1.17 ML, 210MG/2.34ML (105MG/1.17MLX2)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	ONE OF THE FOLLOWING: (1) HIGH RISK FOR FRACTURES DEFINED AS ONE OF THE FOLLOWING: A) HISTORY OF OSTEOPOROTIC (I.E., FRAGILITY, LOW TRAUMA) FRACTURE(S). B) 2 OR MORE RISK FACTORS FOR FRACTURE (E.G., HISTORY OF MULTIPLE RECENT LOW TRAUMA FRACTURES, BMD T-SCORE LESS THAN OR EQUAL TO -2.5, CORTICOSTEROID USE, OR USE OF GNRH ANALOGS SUCH AS NAFARELIN, ETC.). C) NO PRIOR TREATMENT FOR OSTEOPOROSIS AND FRAX SCORE OF AT LEAST 20% FOR ANY MAJOR FRACTURE OR OF AT LEAST 3% FOR HIP FRACTURE. (2) UNABLE TO USE ORAL THERAPY (I.E., UPPER GASTROINTESTINAL PROBLEMS UNABLE TO TOLERATE ORAL MEDICATION, LOWER GASTROINTESTINAL PROBLEMS UNABLE TO ABSORB ORAL MEDICATIONS, TROUBLE REMEMBERING TO TAKE ORAL MEDICATIONS OR COORDINATING AN ORAL BISPHOSPHONATE WITH OTHER ORAL MEDICATIONS OR THEIR DAILY ROUTINE). (3) THERAPEUTIC FAILURE OF, INTOLERANCE TO, OR A CONTRAINDICATION TO BISPHOSPHONATES.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	No

ROPEGINTERFERON ALFA-2B-NJFT

Products Affected

- BESREMI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ROZANOLIXIZUMAB-NOLI

Products Affected

- RYSTIGGO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	GENERALIZED MYASTHENIA GRAVIS (GMG): 1) POSITIVE SEROLOGICAL TEST FOR ANTI-ACETYLCHOLINE RECEPTOR (ANTI-ACHR) ANTIBODIES OR ANTI-MUSCLE-SPECIFIC TYROSINE KINASE (MUSK) ANTIBODIES POSITIVE TEST RESULTS, 2) DOCUMENTATION OF PRIOR AND/OR CURRENT TREATMENTS PATIENT HAS USED FOR MYASTHENIA GRAVIS
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH NEUROLOGIST
Coverage Duration	6 Weeks
Other Criteria	PART B VS D EVALUATION ALSO APPLIES. SUBSEQUENT CYCLES CAN BE ADMINISTERED BASED ON CLINICAL EVALUATION. THE SAFETY OF INITIATING SUBSEQUENT CYCLES SOONER THAN 63 DAYS FROM THE START OF THE PREVIOUS TREATMENT CYCLE HAS NOT BEEN ESTABLISHED.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RUCAPARIB

Products Affected

- RUBRACA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) PREVIOUSLY RECEIVED A BILATERAL ORCHIECTOMY, OR 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

RUXOLITINIB

Products Affected

- JAKAFI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MYELOFIBROSIS RENEWAL: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION. ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD): NO RENEWAL CRITERIA.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	MYELOFIBROSIS: INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS. POLYCYTHEMIA VERA, GVHD: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SECUKINUMAB

Products Affected

- COSENTYX (2 SYRINGES)
- COSENTYX PEN (2 PENS)
- COSENTYX SUBCUTANEOUS SYRINGE 75 MG/0.5 ML
- COSENTYX UNOREADY PEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, OR FACE. NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
Age Restrictions	
Prescriber Restrictions	PSO, HS: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR A DERMATOLOGIST. ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, ENTHESITIS-RELATED ARTHRITIS (ERA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. AS, NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID (NON-STEROIDAL ANTI-INFLAMMATORY DRUG). ERA: TRIAL OF OR CONTRAINDICATION TO ONE NSAID, SULFASALAZINE, OR METHOTREXATE. RENEWAL: PSO, PSA, AS, NR-AXSPA, ERA, HS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

SELEXIPAG

Products Affected

- UPTRAVI ORAL TABLET 1,000 MCG, 1,200 MCG, 1,400 MCG, 1,600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG
- UPTRAVI ORAL TABLETS,DOSE PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PULMONARY ARTERIAL HYPERTENSION (PAH): 1) CONFIRMATORY DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION: A) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, B) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) 15 MMHG OR LESS, AND C) PULMONARY VASCULAR RESISTANCE (PVR) 3 WOOD UNITS OR GREATER, AND 2) NYHA-WHO FUNCTIONAL CLASS (FC) II-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	PAH: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PAH: WHO FC II-III SYMPTOMS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING AGENTS FROM DIFFERENT DRUG CLASSES: 1) FORMULARY VERSION OF AN ORAL ENDOTHELIAN RECEPTOR ANTAGONIST (E.G., AMBRISENTAN, BOSENTAN, MACITENTAN), 2) FORMULARY VERSION OF AN ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR (E.G., SILDENAFIL, TADALAFIL), OR 3) FORMULARY VERSION OF AN ORAL CGMP STIMULATOR (E.G., RIOCIGUAT). WHO FC III SYMPTOMS AND EVIDENCE OF RAPID PROGRESSION OR POOR PROGNOSIS, WHO FC IV SYMPTOMS: NO STEP. RENEWAL: PAH: IMPROVEMENT FROM BASELINE IN THE 6-MINUTE WALK DISTANCE TEST OR REMAINED STABLE FROM BASELINE IN THE 6-MINUTE WALK DISTANCE TEST AND WHO FC HAS IMPROVED OR REMAINED STABLE.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No

SELINEXOR

Products Affected

- XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40 MG/WEEK (10 MG X 4), 40 MG/WEEK (40 MG X 1), 40MG TWICE WEEK (40 MG X 2), 60 MG/WEEK (60 MG X 1), 60MG TWICE WEEK (120 MG/WEEK), 80 MG/WEEK (40 MG X 2), 80MG TWICE WEEK (160 MG/WEEK)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SELPERCATINIB

Products Affected

- RETEVMO ORAL CAPSULE 40 MG, 80 MG
- RETEVMO ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SELUMETINIB

Products Affected

- KOSELUGO ORAL CAPSULE 10 MG, 25 MG
- KOSELUGO ORAL CAPSULE, SPRINKLE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SIPONIMOD

Products Affected

- MAYZENT ORAL TABLET 0.25 MG, 1 MG, 2 MG
- MAYZENT STARTER(FOR 1MG MAINT)
- MAYZENT STARTER(FOR 2MG MAINT)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MULTIPLE SCLEROSIS: RENEWAL: 1) DEMONSTRATION OF CLINICAL BENEFIT COMPARED TO PRE-TREATMENT BASELINE AND 2) DOES NOT HAVE LYMPHOPENIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SIROLIMUS PROTEIN-BOUND

Products Affected

- FYARRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 months
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SODIUM OXYBATE

Products Affected

- XYREM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH ONE OF THE FOLLOWING SPECIALISTS: NEUROLOGIST, PSYCHIATRIST, OR SPECIALIST IN SLEEP MEDICINE
Coverage Duration	INITIAL 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	ALL INDICATIONS: INITIAL: THE PATIENT IS NOT CURRENTLY BEING TREATED WITH SEDATIVE HYPNOTIC AGENTS. EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY: FOR PATIENTS 18 YEARS OR OLDER: TRIAL OF OR CONTRAINDICATION TO THE FORMULARY VERSION OF MODAFINIL, ARMODAFINIL, PITOLISANT OR SOLRIAMFETOL AND ONE OTHER GENERIC STIMULANT INDICATED FOR EDS IN NARCOLEPSY. FOR PATIENTS 7 TO 17 YEARS OF AGE: TRIAL OF OR CONTRAINDICATION TO ONE OTHER GENERIC STIMULANT INDICATED FOR EDS IN NARCOLEPSY. RENEWAL: SUSTAINED IMPROVEMENT OF SYMPTOMS COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SOFOSBUVIR/VELPATASVIR

Products Affected

- *sofosbuvir-velpatasvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (HEPATOLOGIST), OR A SPECIALLY TRAINED GROUP SUCH AS ECHO (EXTENSION FOR COMMUNITY HEALTHCARE OUTCOMES) MODEL.
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. HCV RNA LEVEL WITHIN PAST 6 MONTHS. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS NOT RECOMMENDED BY THE MANUFACTURER: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, HIV REGIMEN THAT CONTAINS EFAVIRENZ, ROSUVASTATIN AT DOSES ABOVE 10MG, TIPRANAVIR/RITONAVIR OR TOPOTECAN. PATIENTS WITH DECOMPENSATED CIRRHOSIS REQUIRE CONCURRENT RIBAVIRIN UNLESS RIBAVIRIN INELIGIBLE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SOLRIAMFETOL

Products Affected

- SUNOSI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY: NEUROLOGIST, PSYCHIATRIST, OR SPECIALIST IN SLEEP MEDICINE.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY: THE PATIENT HAS TRIED THE FORMULARY VERSION OF MODAFINIL OR ARMODAFINIL AND ONE OTHER GENERIC STIMULANT INDICATED FOR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY. OBSTRUCTIVE SLEEP APNEA (OSA): THE PATIENT HAS TRIED THE FORMULARY VERSION OF MODAFINIL OR ARMODAFINIL. RENEWAL: SUSTAINED IMPROVEMENT OF SYMPTOMS COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SOMATROPIN - NORDITROPIN

Products Affected

- NORDITROPIN FLEXPRO

PA Criteria	Criteria Details
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES.
Required Medical Information	INITIAL: PEDIATRIC GROWTH HORMONE DEFICIENCY (GHD), IDIOPATHIC SHORT STATURE (ISS), SMALL FOR GESTATIONAL AGE (SGA), TURNER SYNDROME (TS), NOONAN SYNDROME: HEIGHT AT LEAST 2 STANDARD DEVIATIONS (SD) BELOW THE MEAN HEIGHT FOR NORMAL CHILDREN OF THE SAME AGE AND GENDER. PRADER WILLI SYNDROME (PWS): CONFIRMED GENETIC DIAGNOSIS.
Age Restrictions	
Prescriber Restrictions	INITIAL AND RENEWAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: ADULT GHD: GROWTH HORMONE DEFICIENCY ALONE OR ASSOCIATED WITH MULTIPLE HORMONE DEFICIENCIES (HYPOPITUITARISM), AS A RESULT OF PITUITARY DISEASES, HYPOTHALAMIC DISEASE, SURGERY, RADIATION THERAPY, TRAUMA, OR CONTINUATION OF THERAPY FROM CHILDHOOD ONSET GROWTH HORMONE DEFICIENCY. PEDIATRIC GHD, ISS, SGA, TS, NOONAN SYNDROME: OPEN EPIPHYSES. RENEWAL: PEDIATRIC GHD, ISS, SGA, TS, NOONAN SYNDROME: 1) IMPROVEMENT WHILE ON THERAPY (I.E., INCREASED HEIGHT OR INCREASED GROWTH VELOCITY), AND 2) OPEN EPIPHYSES. PWS: IMPROVEMENT IN BODY COMPOSITION.
Indications	All FDA-approved Indications.
Off Label Uses	

PA Criteria	Criteria Details
Part B Prerequisite	No

SOMATROPIN - SEROSTIM

Products Affected

- SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG

PA Criteria	Criteria Details
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES.
Required Medical Information	INITIAL: HIV/WASTING: MEETS ONE OF THE FOLLOWING CRITERIA FOR WEIGHT LOSS: 10% UNINTENTIONAL WEIGHT LOSS OVER 12 MONTHS, OR 7.5% OVER 6 MONTHS, OR 5% BODY CELL MASS (BCM) LOSS WITHIN 6 MONTHS, OR A BCM LESS THAN 35% (MEN) OF TOTAL BODY WEIGHT AND A BODY MASS INDEX (BMI) LESS THAN 27 KG PER METER SQUARED, OR BCM LESS THAN 23% (WOMEN) OF TOTAL BODY WEIGHT AND A BODY MASS INDEX (BMI) LESS THAN 27 KG PER METER SQUARED, OR BMI LESS THAN 18.5 KG PER METER SQUARED.
Age Restrictions	
Prescriber Restrictions	HIV/WASTING: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, NUTRITIONAL SUPPORT SPECIALIST, OR INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	INITIAL AND RENEWAL: 3 MONTHS.
Other Criteria	HIV/WASTING: INITIAL: INADEQUATE RESPONSE TO ONE PREVIOUS THERAPY (E.G., MEGACE, APPETITE STIMULANTS, ANABOLIC STEROIDS), RENEWAL: CLINICAL BENEFIT IN MUSCLE MASS AND WEIGHT. INITIAL AND RENEWAL: CURRENTLY ON HIV ANTIRETROVIRAL THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SONIDEGIB

Products Affected

- ODOMZO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	BASELINE SERUM CREATINE KINASE (CK) AND SERUM CREATININE LEVELS
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SORAFENIB TOSYLATE

Products Affected

- *sorafenib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SOTORASIB

Products Affected

- LUMAKRAS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

STIRIPENTOL

Products Affected

- DIACOMIT ORAL CAPSULE 250 MG, 500 MG
- DIACOMIT ORAL POWDER IN PACKET 250 MG, 500 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	DRAVET SYNDROME: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	RENEWAL: DRAVET SYNDROME: CURRENTLY TREATED WITH CLOBAZAM.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

SUNITINIB MALATE

Products Affected

- *sunitinib malate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	GASTROINTESTINAL STROMAL TUMORS (GIST): TRIAL OF OR CONTRAINDICATION TO GLEEVEC.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TACROLIMUS XR

Products Affected

- ASTAGRAF XL
- ENVARSUS XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TADALAFIL BPH

Products Affected

- *tadalafil oral tablet 2.5 mg, 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	BENIGN PROSTATIC HYPERPLASIA (BPH): DIAGNOSIS OF BPH.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TAFAMIDIS

Products Affected

- VYNDAMAX
- VYNDAQEL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION THAT THE PATIENT HAS NOT PROGRESSED TO NYHA CLASS IV HEART FAILURE.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A CARDIOLOGIST, ATTR SPECIALIST, OR MEDICAL GENETICIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PATIENT HAS NEW YORK HEART ASSOCIATION (NYHA) CLASS I, II, OR III HEART FAILURE. DIAGNOSIS CONFIRMED BY ONE OF THE FOLLOWING: 1) BONE SCAN (SCINTIGRAPHY) STRONGLY POSITIVE FOR MYOCARDIAL UPTAKE OF 99MTCPYP/DPD, OR 2) BIOPSY OF TISSUE OF AFFECTED ORGAN(S) (CARDIAC AND POSSIBLY NON-CARDIAC SITES) TO CONFIRM AMYLOID PRESENCE AND CHEMICAL TYPING TO CONFIRM PRESENCE OF TRANSTHYRETIN (TTR) PROTEIN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TALAZOPARIB

Products Affected

- TALZENNA ORAL CAPSULE 0.1 MG, 0.25 MG, 0.35 MG, 0.5 MG, 0.75 MG, 1 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PATIENT HAS BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING. PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER MUST HAVE ADDITIONAL PRIOR TREATMENT WITH ENDOCRINE THERAPY OR BE CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY. ONLY VERIFICATION OF INDICATION FOR PROSTATE CANCER IS NEEDED. THERE ARE NO EXTRA REQUIREMENTS FOR THIS INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TALETRECTINIB ADIPATE

Products Affected

- IBTROZI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) Diagnosis: Patient has locally advanced or metastatic Non-Small Cell Lung Cancer (NSCLC). 2) Document: Patient has documented ROS1 plus NSCLC. 3) Ibtrozi is given as monotherapy.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST, ONCOLOGIST OR PULMONOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TASIMELTEON

Products Affected

- HETLIOZ LQ
- *tasimelteon*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	LIFETIME
Other Criteria	NON-24 HOUR SLEEP-WAKE DISORDER: PATIENT IS LIGHT-INSENSITIVE OR HAS TOTAL BLINDNESS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TAZEMETOSTAT

Products Affected

- TAZVERIK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TECLISTAMAB-CQYV

Products Affected

- TECVAYLI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	RELAPSED OR REFRACTORY MULTIPLE MYELOMA: 1.) THERAPEUTIC FAILURE TO AT LEAST FOUR PRIOR LINES OF THERAPY, INCLUDING AT LEAST ONE PROTEASOME INHIBITOR (E.G., BORTEZOMIB, CARFILZOMIB, IXAZOMIB), ONE IMMUNOMODULATORY AGENT (E.G., THALIDOMIDE, LENALIDOMIDE, POMALIDOMIDE), AND ONE CD38-DIRECTED MONOCLONAL ANTIBODY (E.G., DARATUMUMAB).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TEDUGLUTIDE

Products Affected

- GATTEX 30-VIAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PATIENT IS DEPENDENT ON INTRAVENOUS PARENTERAL NUTRITION DEFINED AS REQUIRING PARENTERAL NUTRITION AT LEAST THREE TIMES PER WEEK
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TELISOTUZUMAB VEDOTIN-TLLV

Products Affected

- EMRELIS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TELOTRISTAT

Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TEPOTINIB

Products Affected

- TEPMETKO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TEPROTUMUMAB-TRBW

Products Affected

- TEPEZZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TERIFLUNOMIDE

Products Affected

- *teriflunomide*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TESAMORELIN

Products Affected

- EGRIFTA SV
- EGRIFTA WR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TESTOSTERONE

Products Affected

- *testosterone cypionate*
- *testosterone enanthate*
- *testosterone transdermal gel in metered-dose pump 12.5 mg/ 1.25 gram (1 %), 20.25 mg/1.25 gram (1.62 %)*
- *testosterone transdermal gel in packet 1 % (25 mg/2.5gram), 1 % (50 mg/5 gram)*
- *testosterone transdermal solution in metered pump w/app*
- XYOSTED

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: MALE HYPOGONADISM: CONFIRMED BY: 1) AT LEAST TWO MORNING TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS WHILE IN A FASTED STATE, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 PG/ML.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	PRIMARY OR SECONDARY HYPOGONADISM: 12 MO. ALL OTHER INDICATIONS: LIFETIME OF MEMBERSHIP IN PLAN.
Other Criteria	MALE HYPOGONADISM: RENEWAL: IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TETRABENAZINE

Products Affected

- *tetrabenazine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TEZACAFTOR/IVACAFTOR

Products Affected

- SYMDEKO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CYSTIC FIBROSIS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME
Other Criteria	RENEWAL: MAINTAINED, IMPROVED, OR DEMONSTRATED LESS THAN EXPECTED DECLINE IN FEV1 OR BODY MASS INDEX (BMI), OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

THALIDOMIDE

Products Affected

- THALOMID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TILDRAKIZUMAB

Products Affected

- ILUMYA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER.
Prescriber Restrictions	Prescribed by or in consultation with a dermatologist.
Coverage Duration	Initial: 6 months. Renewal: 12 months
Other Criteria	Initial: Plaque psoriasis: 1) The patient has psoriatic lesions involving greater than or equal to 3% of body surface area (BSA) OR psoriatic lesions affecting the hands, feet, genital area, scalp, or face. 2) The patient had a previous trial of or contraindication to ONE or more forms of conventional therapies such as PUVA (Phototherapy Ultraviolet Light A), UVB (Ultraviolet Light B), topical corticosteroids, calcipotriene, acitretin, methotrexate, or cyclosporine. Renewal: 1) Documentation showing patient has achieved or maintained clear or minimal disease or a decrease in PASI (Psoriasis Area and Severity Index) from baseline.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TISLELIZUMAB

Products Affected

- TEVIMBRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: FOR THE TREATMENT OF UNRESECTABLE OR METASTATIC ESOPHAGEAL SQUAMOUS CELL CARCINOMA (ESCC). 2) DOCUMENT: DISEASE PROGRESSION FOLLOWING AT LEAST ONE SYSTEMIC THERAPY [E.G., FLUOROPYRIMIDINE (FLUOROURACIL OR CAPECITABINE), OXALIPLATIN OR CISPLATIN, ETC.] THAT DID NOT INCLUDE A PD-1 (I.E., NIVOLUMAB, PEMBROLIZUMAB, ETC.), PD-L1 (I.E., DURVALUMAB, ETC.), OR CTLA-4 (I.E., IPILIMUMAB, ETC.) DIRECTED THERAPY FOR RECURRENT LOCALLY ADVANCED OR METASTATIC ESCC.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	PART D VS. PART B EVALUATION ALSO APPLIES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TIVOZANIB

Products Affected

- FOTIVDA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TOFACITINIB

Products Affected

- XELJANZ ORAL SOLUTION
- XELJANZ ORAL TABLET
- XELJANZ XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), POLYARTICULAR COURSE JUVENILE IDIOPATHIC ARTHRITIS (PCJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. PCJIA: TRIAL OF OR CONTRAINDICATION TO ONE DMARD. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. RENEWAL: RA, PSA, AS, PCJIA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TOLVAPTAN

Products Affected

- JYNARQUE ORAL TABLET *sequential*
- JYNARQUE ORAL TABLETS, SEQUENTIAL
- *tolvaptan (polycys kidney dis) oral tablets,*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION THAT PATIENT HAS NOT PROGRESSED TO ESRD/DIALYSIS OR TRANSPLANT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: THE PATIENT MEETS ALL OF THE FOLLOWING: (1) CONFIRMED POLYCYSTIC KIDNEY DISEASE VIA CT, MRI IMAGING, OR ULTRASOUND (2) GENETIC TESTING FOR CAUSATIVE MUTATIONS OR FAMILY HISTORY OF CONFIRMED POLYCYSTIC KIDNEY DISEASE IN ONE OR BOTH PARENTS, AND (3) PATIENT DOES NOT HAVE ESRD.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TOPICAL TRETINOIN

Products Affected

- ALTRENO
- *tretinoin*

PA Criteria	Criteria Details
Exclusion Criteria	COSMETIC INDICATIONS SUCH AS WRINKLES, PHOTOAGING, MELASMA.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	BRAND TOPICAL TRETINOIN REQUIRES TRIAL OF OR CONTRAINDICATION TO A FORMULARY GENERIC TOPICAL TRETINOIN PRODUCT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TOVORAFENIB

Products Affected

- OJEMDA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: FOR THE TREATMENT OF PATIENTS 6 MONTHS OF AGE AND OLDER WITH RELAPSED OR REFRACTORY PEDIATRIC LOW_GRADE GLIOMA (PLGG). 2) DOCUMENT: A) BRAF FUSION OR REARRANGEMENT OR BRAF V600 MUTATION B) DOCUMENTATION OF PATIENT HAS RECEIVED AT LEAST ONE SYSTEMIC THERAPY (TAFINLAR PLUS MEKINIST) PRIOR OR CURRENT TREATMENTS PATIENT HAS USED FOR PLGG AND C) PATIENTS ACTUAL BODY WEIGHT AND BODY SURFACE AREA (DOSING BASED ON BODY SURFACE AREA).
Age Restrictions	6 MONTHS OF AGE AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	1) VALIDATE THAT THE TREATMENT REGIMEN IS FOLLOWING THE MOST UP TO DATE NCCN GUIDELINES RECOMMENDATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TRAMETINIB

Products Affected

- MEKINIST ORAL RECON SOLN
- MEKINIST ORAL TABLET 0.5 MG, 2 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TREMELIMUMAB-ACTL

Products Affected

- IMJUDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	LOCALLY ADVANCED UNRESECTABLE AND/OR METASTATIC HEPATOCELLULAR CARCINOMA (UHCC): 1) PRESCRIBED IN COMBINATION WITH DURVALUMAB AS FIRST-LINE THERAPY ONLY. 2) CHILD-PUGH SCORE CLASS A. 3) PRIOR HISTORY OF PRIOR SYSTEMIC TREATMENT, INCLUDING PD-1/PD-L1 USE, 4) ECOG PERFORMANCE STATUS OF 0-1, 5) RATIONAL OF WHY ATEZOLIZUMAB AND BEVACIZUMAB COMBINATION OR DURVALUMAB AS MONOTHERAPY IS NOT APPROPRIATE AS MONOTHERAPY. METASTATIC NON-SMALL-CELL LUNG CARCINOMA (NSCLC): 1) PRESCRIBED IN COMBINATION WITH DURVALUMAB AND PLATINUM-BASED CHEMOTHERAPY IN PATIENTS WITH NO EGFR OR ALK GENOMIC TUMOR ABERRATIONS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TREOSULFAN

Products Affected

- GRAFAPEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: A) ACUTE MYELOID LEUKEMIA (AML) AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANTATION (ALLOHSCT) B) MYELODYSPLASTIC SYNDROME (MDS) AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANTATION AND II) ACTUAL BODY WEIGHT OR BSA
Age Restrictions	1 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, ONCOLOGIST, OR TRANSPLANT SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TRIENTINE

Products Affected

- *trientine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	WILSONS DISEASE: INITIAL: KNOWN FAMILY HISTORY OF WILSONS DISEASE OR PHYSICAL EXAMINATION CONSISTENT WITH WILSONS DISEASE. CONFIRMATION OF ONE OF THE FOLLOWING: 1) PLASMA COPPER-PROTEIN CERULOPLASMIN LESS THAN 20 MG/DL, 2) LIVER BIOPSY POSITIVE FOR AN ABNORMALLY HIGH CONCENTRATION OF COPPER (GREATER THAN 250 MCG/G DRY WEIGHT) OR THE PRESENCE OF KAYSER-FLEISCHER RINGS, OR 3) CONFIRMATION BY GENETIC TESTING FOR ATP7B MUTATIONS, RENEWAL: PATIENT CONTINUES TO BENEFIT FROM THE MEDICATION.
Age Restrictions	
Prescriber Restrictions	WILSONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST OR GASTROENTEROLOGIST.
Coverage Duration	WILSONS DISEASE: INITIAL: 12 MONTHS, RENEWAL: LIFETIME.
Other Criteria	INITIAL: WILSONS DISEASE: TRIAL OF OR CONTRAINDICATION TO FORMULARY VERSION OF PENICILLAMINE (DEPEN).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TRIFLURIDINE/TIPIRACIL

Products Affected

- LONSURF ORAL TABLET 15-6.14 MG, 20-8.19 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

TUCATINIB

Products Affected

- TUKYSA ORAL TABLET 150 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

UBROGEPANT

Products Affected

- UBRELVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE FORMULARY TRIPTAN. RENEWAL: THE PATIENT HAS EXPERIENCED AN IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE OR THE PATIENT HAS EXPERIENCED CLINICAL IMPROVEMENT AS DEFINED BY ONE OF THE FOLLOWING: 1) ABILITY TO FUNCTION NORMALLY WITHIN 2 HOURS OF DOSE, 2) HEADACHE PAIN DISAPPEARS WITHIN 2 HOURS OF DOSE, 3) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ULTOMIRIS

Products Affected

- ULTOMIRIS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	Prescribed by or in consultation with a hematologist.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 6 MONTHS.
Other Criteria	INITIAL: 1) Diagnosis of Paroxysmal nocturnal hemoglobinuria (PNH) and meets the following: A) Flow cytometry shows detectable GPI-deficient hematopoietic clones or greater than or equal to 5% PNH cells, B) Member has one of the following: i) History of greater than or equal to 1 transfusion in the past 24 months with documentation of hemoglobin less than or equal to 7 g/dL in members without anemia symptoms or hemoglobin less than or equal to 10 g/dL in members with anemia symptoms, or ii) History of thrombosis. RENEWAL: 1) PNH: Member is responding positively to therapy as evidenced by, including but not limited to, improvement in any of the following parameters: Improved measures of intravascular hemolysis (e.g., normalization of lactate dehydrogenase [LDH]), Reduced need for red blood cell transfusions, Increased or stabilization of hemoglobin levels, Less fatigue, Improved health-related quality of life, or Fewer thrombotic events.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

UPADACITINIB

Products Affected

- RINVOQ
- RINVOQ LQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS (RA), GIANT CELL ARTERITIS (GCA) AND ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ATOPIC DERMATITIS (AD): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST, OR IMMUNOLOGIST. FOR CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): GASTROENTEROLOGIST.
Coverage Duration	INITIAL:RA,PSA,AS,CD,UC,NR-AXSPA,PJIA,GCA: 6 MONTHS. AD: 4 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED AND INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS. PSA: INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS. ATOPIC DERMATITIS: TRIAL OF A HIGH OR SUPER HIGH POTENCY TOPICAL CORTICOSTEROID (E.G., TRIAMCINOLONE ACETONIDE, FLUOCINONIDE, CLOBETASOL PROPIONATE, HALOBETASOL PROPIONATE) OR ONE NON-STEROIDAL TOPICAL IMMUNOMODULATING AGENT (E.G., EUCRISA, OPZELURA, PIMECROLIMUS, TACROLIMUS). AS AND NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID (NON-STEROIDAL ANTI-INFLAMMATORY DRUG) AND INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS. CD: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A CORTICOSTEROID (E.G., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE AND INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS.RENEWAL: RA, PSA, UC, AS, GCA AND NR-AXSPA: CONTINUES TO BENEFIT FROM THE MEDICATION. ATOPIC DERMATITIS: EXPERIENCED OR MAINTAINED IMPROVEMENT IN AT LEAST TWO OF THE FOLLOWING: INTRACTABLE PRURITUS, CRACKING AND OOZING/BLEEDING OF AFFECTED SKIN, IMPAIRED ACTIVITIES OF DAILY LIVING.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

URIDINE TRIACETATE

Products Affected

- XURIDEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: DIAGNOSIS CONFIRMED BY 1) GENETIC MUTATION OF URIDINE MONOPHOSPHATE SYNTHASE (UMPS) GENE AND 2) ELEVATED URINE OROTIC ACID PER AGE-SPECIFIC REFERENCE RANGE. RENEWAL: IMPROVEMENT FROM BASELINE OR STABILIZATION OF AGE DEPENDENT HEMATOLOGIC PARAMETERS (E.G., NEUTROPHIL COUNT, NEUTROPHIL PERCENT, WBC COUNT, MEAN CORPUSCULAR VOLUME)
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PRESCRIBER SPECIALIZING IN INHERITED METABOLIC DISEASES
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

USTEKINUMAB

Products Affected

- STELARA INTRAVENOUS
- STELARA SUBCUTANEOUS SOLUTION
- STELARA SUBCUTANEOUS SYRINGE
  45 MG/0.5 ML, 90 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. CD: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL THERAPY SUCH AS A CORTICOSTEROID (E.G., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VALBENAZINE

Products Affected

- INGREZZA
- INGREZZA SPRINKLE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	HUNTINGTON DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST. TARDIVE DYSKINESIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR MOVEMENT DISORDER SPECIALIST.
Coverage Duration	12 Months
Other Criteria	TARDIVE DYSKINESIA: PATIENT HAS A HISTORY OF USING AGENTS THAT CAUSE TARDIVE DYSKINESIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VANDETANIB

Products Affected

- CAPRELSA ORAL TABLET 100 MG, 300 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VEMURAFENIB

Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VENETOCLAX

Products Affected

- VENCLEXTA ORAL TABLET 10 MG, 100 MG, 50 MG
- VENCLEXTA STARTING PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VERICIGUAT

Products Affected

- VERQUVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CHRONIC HEART FAILURE (CHF): PATIENT HAS AN EJECTION FRACTION LESS THAN 45 PERCENT. PATIENT HAS NEW YORK HEART ASSOCIATION (NYHA) CLASS II, III, OR IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 Months
Other Criteria	CHF: ONE OF THE FOLLOWING: A) PATIENT WAS HOSPITALIZED FOR HEART FAILURE WITHIN THE LAST 6 MONTHS, OR B) PATIENT USED OUTPATIENT INTRAVENOUS DIURETICS (E.G., BUMETANIDE, FUROSEMIDE) FOR HEART FAILURE WITHIN THE LAST 3 MONTHS. CHF: TRIAL AND FAILURE, CONTRAINDICATION, OR INTOLERANCE TO TWO OF THE FOLLOWING AT A MAXIMALLY TOLERATED DOSE: A) ONE OF THE FOLLOWING: 1) ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (E.G., CAPTOPRIL, ENALAPRIL), 2) ANGIOTENSIN II RECEPTOR BLOCKER (ARB) (E.G., CANDESARTAN, VALSARTAN), OR 3) ANGIOTENSIN RECEPTOR-NEPRILYSIN INHIBITOR (ARNI) [E.G., ENTRESTO (SACUBITRIL AND VALSARTAN)], B) ONE OF THE FOLLOWING: 1) BISOPROLOL, 2) CARVEDILOL, OR 3) METOPROLOL SUCCINATE EXTENDED RELEASE, C) SODIUM-GLUCOSE CO-TRANSPORTER 2 (SGLT2) INHIBITOR [E.G., JARDIANCE (EMPAGLIFLOZIN), FARXIGA (DAPAGLIFLOZIN), XIGDUO XR (DAPAGLIFLOZIN AND METFORMIN)], OR D) MINERALOCORTICOID RECEPTOR ANTAGONIST (MRA) [E.G., EPLERENONE, SPIRONOLACTONE].

PA Criteria	Criteria Details
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VIGABATRIN

Products Affected

- *vigabatrin*
- *vigadrone oral powder in packet*
- VIGAFYDE
- *vigpoder*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	REFRACTORY COMPLEX PARTIAL SEIZURES (CPS), INFANTILE SPASMS: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	CPS: PATIENT HAS RESPONDED INADEQUATELY TO AT LEAST 2 ANTIEPILEPTIC AGENTS. CPS AND INFANTILE SPASMS: BENEFITS OUTWEIGH THE POTENTIAL FOR VISION LOSS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VIMSELTINIB

Products Affected

- ROMVIMZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: A) SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT).
Age Restrictions	18 YEARS OF AGE AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST/ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VISMODEGIB

Products Affected

- ERIVEDGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VORASIDENIB

Products Affected

- VORANIGO ORAL TABLET 10 MG, 40 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	12 YEARS OF AGE AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	1) DIAGNOSIS: GRADE 2 ASTROCYTOMA OR OLIGODENDROGLIOMA, 2)DOCUMENT: A) SUSCEPTIBLE IDH1 OR IDH2 MUTATION FOLLOWING SURGERY INCLUDING BIOPSY, SUB-TOTAL RESECTION, OR GROSS TOTAL RESECTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

VORICONAZOLE SUSPENSION

Products Affected

- *voriconazole oral suspension for reconstitution*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CANDIDA INFECTIONS: 3 MOS. ALL OTHER INDICATIONS: 6 MOS.
Other Criteria	CANDIDA INFECTIONS: TRIAL OF OR CONTRAINDICATION TO FLUCONAZOLE. ALL INDICATIONS: INABILITY TO SWALLOW TABLETS OR AN INDICATION FOR ESOPHAGEAL CANDIDIASIS. CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE REQUIRES NO EXTRA CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

XANOMELINE/TROSPIUM

Products Affected

- COBENFY
- COBENFY STARTER PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	MUST BE AGE 18 OR OLDER
Prescriber Restrictions	MUST BE PRESCRIBED BY, OR IN CONSULTATION WITH, A SPECIALIST FOR THE CONDITION BEING TREATED
Coverage Duration	12 MONTHS
Other Criteria	TRIAL AND FAILURE (DEFINED AS AN INADEQUATE RESPONSE) OF TWO OF THE FOLLOWING: ARIPIPRAZOLE, ASENAPINE (SAPHRIS), BREXPIPRAZOLE (REXULTI), CARIPRAZINE (VRAYLAR), CHLOPROMAZINE, HALDOL, HALOPERIDOL, ILOPERIDONE (FANAPT), LUMATEPERONEE (CAPLYTA), LURASIDONE (LATUDA), OLANZAPINE, PALIPERIDONE (INVEGA), PERPHENAZINE, QUETIAPINE, RISPERIDONE, THIORIDAZINE, THIOTHIXENE, TRIFLUOPERAZINE, OR ZIPRASIDONE (AT LEAST A 30-DAY SUPPLY IN THE PRIOR 180 DAYS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZANIDATAMAB HRII

Products Affected

- ZIIHERA

PA Criteria	Criteria Details
Exclusion Criteria	PREGNANCY
Required Medical Information	1) DIAGNOSIS: FOR THE TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE OR METASTATIC HER2-POSITIVE (IHC 3+) BILIARY TRACT CANCER (BTC). 2) DOCUMENT: A) POSITIVE HER2 STATUS, B) DOCUMENTATION OF PRIOR AND/OR CURRENT FIRST-LINE (I.E. CHEMOTHERAPY-BASED REGIMEN) TREATMENTS PATIENT HAS USED FOR BTC AND C) PATIENTS ACTUAL BODY WEIGHT (WEIGHT-BASED DOSING).
Age Restrictions	18 YEARS OF AGE AND OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZANUBRUTINIB

Products Affected

- BRUKINSA ORAL CAPSULE
- BRUKINSA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZENOCUTUZUMAB ZBCO

Products Affected

- BIZENGRI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: A) TREATMENT OF ADULTS WITH ADVANCED, UNRESECTABLE, OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) HARBORING A NEUREGULIN 1 (NRG1) GENE FUSION WITH DISEASE PROGRESSION ON OR AFTER PRIOR SYSTEMIC THERAPY, B) TREATMENT OF ADULTS WITH ADVANCED, UNRESECTABLE, OR METASTATIC PANCREATIC ADENOCARCINOMA HARBORING A NRG1 GENE FUSION WITH DISEASE PROGRESSION ON OR AFTER PRIOR SYSTEMIC THERAPY. 2) DOCUMENT FOR ALL INDICATIONS: A) PRESENCE OF NRG1 GENE FUSION AS DETERMINED BY AN FDA-APPROVED TEST, B) DISEASE PROGRESSION ON OR AFTER PRIOR SYSTEMIC THERAPY
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	1) FDA APPROVED TEST TO DETERMINE PRESENCE OF NRG1 GENE FUSION DETECTED BY DNA-BASED NEXT-GENERATION SEQUENCING (NGS) OR BY RNA-BASED NGS THAT MAY INCLUDE DNA SEQUENCING.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZOLBETUXIMAB CLZB

Products Affected

- VYLOY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) DIAGNOSIS: LOCALLY ADVANCED UNRESECTABLE OR METASTATIC HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA WHOSE TUMORS ARE CLAUDIN (CLDN) 18.2 POSITIVE. 2) DOCUMENTATION OF A) USE IN COMBINATION WITH FLUOROPYRIMIDINE AND PLATINUM-CONTAINING CHEMOTHERAPY. B) EVIDENCE OR PRESENCE OF (HER2)-NEGATIVE OR CLAUDIN (CLDN) 18.2 POSITIVE AS DETERMINED BY AN FDA-APPROVED TEST.
Age Restrictions	18 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZONGERTINIB

Products Affected

- HERNEXEOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

ZURANOLONE

Products Affected

- ZURZUVAE ORAL CAPSULE 20 MG, 25 MG, 30 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A PSYCHIATRIST OR OBSTETRICIAN/GYNECOLOGIST
Coverage Duration	14 DAYS
Other Criteria	DIAGNOSIS OF POSTPARTUM DEPRESSION (PPD) 1) PHYSICIAN ATTESTATION OF MODERATE TO SEVERE PPD DIAGNOSIS 2) PATIENT IS LESS THAN OR EQUAL TO 12 MONTHS POSTPARTUM, 3) PATIENT HAS THERAPEUTIC FAILURE OR CONTRAINDICATION TO AT LEAST TWO GENERIC SSRI OR SNRI (E.G FLUOXETINE, SERTRALINE, VENLAFAXINE) FOR PPD
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

INDEX

abiraterone 6
abirtega 6
ACTHAR 63
ACTHAR SELFJECT 63
ACTIMMUNE 153
ADEMPAS 245
AIMOVIG AUTOINJECTOR 115
AKEEGA 196
ALECENSA 13
alprazolam oral tablet 0.25 mg, 0.5 mg, 1 mg, 2 mg 146
ALTRENO 300
ALUNBRIG ORAL TABLET 180 MG, 30 MG, 90 MG 47
ALUNBRIG ORAL TABLETS,DOSE PACK 47
alyq 214
ambrisentan 109
amikacin injection solution 1,000 mg/4 ml, 500 mg/2 ml 15
apomorphine 19
APRIZIO PAK II 170
ARIKAYCE 22
ASTAGRAF XL 276
AUGTYRO 238
AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG 85
AUSTEDO XR ORAL TABLET EXTENDED RELEASE 24 HR 12 MG, 18 MG, 24 MG, 30 MG, 36 MG, 42 MG, 48 MG, 6 MG 85
AUSTEDO XR TITRATION KT(WK1-4) ORAL TABLET, EXT REL 24HR DOSE PACK 12-18-24-30 MG 85
AVASTIN INTRAVENOUS 41
AVMAPKI-FAKZYNJA 28
AVONEX INTRAMUSCULAR PEN INJECTOR KIT 154
AVONEX INTRAMUSCULAR SYRINGE KIT 154
AXTLE 220
AYVAKIT 26
BALVERSA ORAL TABLET 3 MG, 4 MG, 5 MG 114
BAXDELA ORAL 82
BENLYSTA INTRAVENOUS 34
BENLYSTA SUBCUTANEOUS 34
benztropine oral 144
BESREMI 252
BETASERON SUBCUTANEOUS KIT 154
bexarotene 44
BIZENGRI 328
BOSULIF ORAL CAPSULE 46
BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG 46
BRAFTOVI 108
BRUKINSA ORAL CAPSULE 327
BRUKINSA ORAL TABLET 327
CABLIVI INJECTION KIT 54
CABOMETYX ORAL TABLET 20 MG, 40 MG, 60 MG 51
CALQUENCE 8
CALQUENCE (ACALABRUTINIB MAL) 8
CAPRELSA ORAL TABLET 100 MG, 300 MG 315
CAYSTON 32
CERDELGA 105
chlordiazepoxide hcl 146
CINRYZE 48
clobazam oral suspension 60
clobazam oral tablet 60
clonazepam oral tablet 0.5 mg, 1 mg, 2 mg 146
clonazepam oral tablet,disintegrating 0.125 mg, 0.25 mg, 0.5 mg, 1 mg, 2 mg 146
clorazepate dipotassium 146
COBENFY 325
COBENFY STARTER PACK 325
COMETRIQ 50
COPIKTRA 97
CORTROPHIN GEL 63
COSENTYX (2 SYRINGES) 256
COSENTYX PEN (2 PENS) 256
COSENTYX SUBCUTANEOUS SYRINGE 75 MG/0.5 ML 256
COSENTYX UNOREADY PEN 256
COTELLIC 62
cyclobenzaprine oral tablet 10 mg, 5 mg ... 144
cyproheptadine oral syrup 144
CYSTARAN 65

dalfampridine 68
DANZITEN 69
DARZALEX 70
DARZALEX FASPRO 70
dasatinib oral tablet 100 mg, 140 mg, 20 mg, 50 mg, 70 mg, 80 mg 73
DATROWAY 74
DAURISMO ORAL TABLET 100 MG, 25 MG 137
deferasirox 76
deferiprone 78
deflazacort oral suspension 80
deflazacort oral tablet 18 mg, 30 mg, 36 mg, 6 mg 80
DIACOMIT ORAL CAPSULE 250 MG, 500 MG 274
DIACOMIT ORAL POWDER IN PACKET 250 MG, 500 MG 274
diazepam intensol 146
diazepam oral solution 5 mg/5 ml (1 mg/ml) 146
diazepam oral tablet 146
dichlorphenamide 86
diclofenac epolamine 87
diclofenac sodium topical gel 3 % 88
diclofenac sodium topical solution in metered-dose pump 88
dicyclomine oral capsule 144
dicyclomine oral solution 144
dicyclomine oral tablet 144
dimethyl fumarate oral capsule,delayed release(dr/ec) 120 mg, 120 mg (14)- 240 mg (46), 240 mg 89
diphenoxylate-atropine oral tablet 144
disopyramide phosphate oral capsule 144
DOPTELET (10 TAB PACK) 27
DOPTELET (15 TAB PACK) 27
DOPTELET (30 TAB PACK) 27
DOPTELET SPRINKLE 27
dronabinol 92
droxidopa 93
DUPIXENT PEN SUBCUTANEOUS PEN INJECTOR 200 MG/1.14 ML, 300 MG/2 ML 94
DUPIXENT SYRINGE SUBCUTANEOUS SYRINGE 100 MG/0.67 ML, 200 MG/1.14 ML, 300 MG/2 ML 94
EGRIFTA SV 290
EGRIFTA WR 290
eltrombopag olamine oral powder in packet 12.5 mg, 25 mg 106
eltrombopag olamine oral tablet 12.5 mg, 25 mg, 50 mg, 75 mg 106
EMFLAZA ORAL SUSPENSION 80
EMFLAZA ORAL TABLET 18 MG, 30 MG, 36 MG, 6 MG 80
EMGALITY PEN 133
EMGALITY SYRINGE SUBCUTANEOUS SYRINGE 120 MG/ML, 300 MG/3 ML (100 MG/ML X 3) 133
EMPRICAINE-II 170
EMRELIS 285
ENBREL MINI 119
ENBREL SUBCUTANEOUS RECON SOLN 119
ENBREL SUBCUTANEOUS SOLUTION 119
ENBREL SUBCUTANEOUS SYRINGE 25 MG/0.5 ML (0.5), 50 MG/ML (1 ML) .. 119
ENBREL SURECLICK 119
ENDARI 168
ENVARSUS XR 276
EPIDIOLEX 52
ERIVEDGE 322
ERLEADA 18
erlotinib oral tablet 100 mg, 150 mg, 25 mg 116
EULEXIN 129
EVENITY SUBCUTANEOUS SYRINGE 105 MG/1.17 ML, 210MG/2.34ML (105MG/1.17MLX2) 250
everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg 121
everolimus (antineoplastic) oral tablet for suspension 121
EVRYSDI 249
EXKIVITY 188
FASENRA PEN 38

FASENRA SUBCUTANEOUS SYRINGE 30 MG/ML 38
fentanyl citrate buccal lozenge on a handle 125
FERRIPROX ORAL SOLUTION 78
fingolimod 128
FINTEPLA 124
FIRMAGON KIT W DILUENT SYRINGE 81
flurazepam 146
FOTIVDA 297
FRUZAQLA ORAL CAPSULE 1 MG, 5 MG 131
FULPHILA 215
FYARRO 264
GALAFOLD 183
GATTEX 30-VIAL 284
GAVRETO 231
gefitinib 135
GILOTRIF 12
glatiramer subcutaneous syringe 20 mg/ml, 40 mg/ml 138
glatopa subcutaneous syringe 20 mg/ml, 40 mg/ml 138
GLEOSTINE 173
glutamine (sickle cell) 168
GOMEKLI ORAL CAPSULE 1 MG, 2 MG 186
GOMEKLI ORAL TABLET FOR SUSPENSION 186
GRAFAPEX 304
HAEGARDA SUBCUTANEOUS RECON SOLN 2,000 UNIT, 3,000 UNIT 49
HERNEXEOS 330
HETLIOZ LQ 281
HUMIRA PEN 10
HUMIRA PEN CROHNS-UC-HS START . 10
HUMIRA PEN PSOR-UVEITS-ADOL HS 10
HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML 10
HUMIRA(CF) PEDI CROHNS STARTER SUBCUTANEOUS SYRINGE KIT 80 MG/0.8 ML, 80 MG/0.8 ML-40 MG/0.4 ML 10
HUMIRA(CF) PEN CROHNS-UC-HS 10
HUMIRA(CF) PEN PEDIATRIC UC 10
HUMIRA(CF) PEN PSOR-UV-ADOL HS . 10
HUMIRA(CF) PEN SUBCUTANEOUS PEN INJECTOR KIT 40 MG/0.4 ML, 80 MG/0.8 ML 10
HUMIRA(CF) SUBCUTANEOUS SYRINGE KIT 10 MG/0.1 ML, 20 MG/0.2 ML, 40 MG/0.4 ML 10
hydroxyzine hcl oral tablet 144
IBRANCE 210
IBTROZI 280
icatibant 148
ICLUSIG 229
IDHIFA 107
ILUMYA 295
imatinib oral tablet 100 mg, 400 mg 150
IMBRUVICA ORAL CAPSULE 140 MG, 70 MG 147
IMBRUVICA ORAL SUSPENSION 147
IMBRUVICA ORAL TABLET 147
IMJUDO 303
IMKELDI 150
IMPAVIDO 185
INBRIJA INHALATION CAPSULE, W/INHALATION DEVICE 167
INGREZZA 314
INGREZZA SPRINKLE 314
INLYTA ORAL TABLET 1 MG, 5 MG 30
INQOVI 75
INREBIC 123
IRESSA 135
ITOVEBI 151
IWILFIN 101
JAKAFI 255
JAYPIRCA 227
jaythari oral tablet 18 mg, 30 mg, 36 mg, 6 mg 80
JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 5 MG 171
JYNARQUE ORAL TABLET 299
JYNARQUE ORAL TABLETS, SEQUENTIAL 299
KALYDECO 155
KESIMPTA PEN 200
KEVEYIS 86
KEYTRUDA 219
KEYTRUDA QLEX SUBCUTANEOUS SOLUTION 395 MG-4,800 UNIT/2.4 ML, 790 MG-9,600 UNIT/4.8 ML 219

KINERET 16
KISQALI FEMARA CO-PACK ORAL TABLET 200 MG/DAY(200 MG X 1)-2.5 MG, 400 MG/DAY(200 MG X 2)-2.5 MG, 600 MG/DAY(200 MG X 3)-2.5 MG 242
KISQALI ORAL TABLET 200 MG/DAY (200 MG X 1), 400 MG/DAY (200 MG X 2), 600 MG/DAY (200 MG X 3) 242
KOSELUGO ORAL CAPSULE 10 MG, 25 MG 262
KOSELUGO ORAL CAPSULE, SPRINKLE 262
KRAZATI 9
KRYSTEXXA 216
KYNMOBI SUBLINGUAL FILM 10 MG, 15 MG, 20 MG, 25 MG, 30 MG 20
KYPROLIS 56
lanreotide 159
lapatinib 160
LAZCLUZE ORAL TABLET 240 MG, 80 MG 162
ledipasvir-sofosbuvir 163
lenalidomide 164
LENVIMA 165
lidocaine hcl mucous membrane solution 4 % (40 mg/ml) 169
lidocaine topical adhesive patch,medicated 5 % 169
lidocaine topical ointment 169
lidocaine-prilocaine topical cream 170
LIVTENCITY 178
LONSURF ORAL TABLET 15-6.14 MG, 20-8.19 MG 306
lorazepam oral tablet 0.5 mg, 1 mg, 2 mg 146
LORBRENA ORAL TABLET 100 MG, 25 MG 174
LUMAKRAS 273
LUNSUMIO 190
LYBALVI 201
LYNPARZA 202
LYTGOBI ORAL TABLET 12 MG/DAY (4 MG X 3), 16 MG/DAY (4 MG X 4), 20 MG/DAY (4 MG X 5) 132
MAVENCLAD (10 TABLET PACK) 59
MAVENCLAD (4 TABLET PACK) 59
MAVENCLAD (5 TABLET PACK) 59
MAVENCLAD (6 TABLET PACK) 59
MAVENCLAD (7 TABLET PACK) 59
MAVENCLAD (8 TABLET PACK) 59
MAVENCLAD (9 TABLET PACK) 59
MAVYRET ORAL PELLETS IN PACKET 139
MAVYRET ORAL TABLET 139
MAYZENT ORAL TABLET 0.25 MG, 1 MG, 2 MG 263
MAYZENT STARTER(FOR 1MG MAINT) 263
MAYZENT STARTER(FOR 2MG MAINT) 263
MEKINIST ORAL RECON SOLN 302
MEKINIST ORAL TABLET 0.5 MG, 2 MG 302
MEKTOVI 45
mifepristone oral tablet 300 mg 182
miglustat 184
MODEYSO 91
morphine concentrate oral solution 143
MOUNJARO 140
MULPLETA 177
MVASI 42
NERLYNX 191
NEULASTA 215
NEXLETOL 37
NEXLIZET 37
NIKTIMVO 29
NILOTINIB D-TARTRATE ORAL CAPSULE 150 MG, 200 MG, 50 MG 192
nilotinib hcl oral capsule 150 mg, 200 mg, 50 mg 192
NINLARO 157
nitisinone 198
NITYR 198
NIVESTYM 127
NORDITROPIN FLEXPRO 268
NOXAFIL ORAL SUSPENSION 230
NUBEQA 72
NUCALA SUBCUTANEOUS AUTO-INJECTOR 179
NUCALA SUBCUTANEOUS RECON SOLN 179
NUCALA SUBCUTANEOUS SYRINGE 100 MG/ML, 40 MG/0.4 ML 179
NUPLAZID 225
NURTEC ODT 244

NUVAKAAN-II 170
NYVEPRIA 215
OCREVUS 199
OCREVUS ZUNOVO 199
ODOMZO 271
OFEV 193
OGSIVEO 197
OJEMDA 301
OJJAARA 189
ONUREG 31
OPSUMIT 109
ORENCIA CLICKJECT 3
ORENCIA SUBCUTANEOUS SYRINGE 125 MG/ML, 50 MG/0.4 ML, 87.5 MG/0.7 ML 3
ORFADIN ORAL CAPSULE 20 MG 198
ORFADIN ORAL SUSPENSION 198
ORGOVYX 237
ORILISSA ORAL TABLET 150 MG, 200 MG 103
ORKAMBI ORAL GRANULES IN PACKET 176
ORKAMBI ORAL TABLET 176
ORLADEYO 40
ORSERDU 102
OTEZLA 21
OTEZLA STARTER 21
OTEZLA XR 21
OZEMPIC SUBCUTANEOUS PEN INJECTOR 0.25 MG OR 0.5 MG (2 MG/3 ML), 0.25 MG OR 0.5 MG(2 MG/1.5 ML), 1 MG/DOSE (4 MG/3 ML), 2 MG/DOSE (8 MG/3 ML) 140
PALYNZIQ 217
paroxetine hcl oral tablet 145
pazopanib 212
PEMAZYRE 221
PEMRYDI RTU 220
penicillamine 222
PIQRAY ORAL TABLET 200 MG/DAY (200 MG X 1), 250 MG/DAY (200 MG X1-50 MG X1), 300 MG/DAY (150 MG X 2) 14
pirfenidone oral capsule 226
pirfenidone oral tablet 267 mg, 801 mg 226
PLEGRIDY SUBCUTANEOUS PEN INJECTOR 125 MCG/0.5 ML 154
PLEGRIDY SUBCUTANEOUS SYRINGE 125 MCG/0.5 ML 154
POMALYST 228
posaconazole oral 230
PRALUENT PEN 213
PREVYMIS ORAL PELLETS IN PACKET 120 MG, 20 MG 166
PREVYMIS ORAL TABLET 166
PRIZOPAK II 170
PROMACTA ORAL POWDER IN PACKET 12.5 MG, 25 MG 106
PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG 106
promethazine oral tablet 12.5 mg 144
pyrimethamine 233
PYRUKYND 187
QINLOCK 247
quinine sulfate 234
QULIPTA 24
RAVICTI 141
REPATHA PUSHTRONEX 213
REPATHA SURECLICK 213
REPATHA SYRINGE 213
RETACRIT INJECTION SOLUTION 10,000 UNIT/ML, 2,000 UNIT/ML, 20,000 UNIT/2 ML, 20,000 UNIT/ML, 3,000 UNIT/ML, 4,000 UNIT/ML, 40,000 UNIT/ML 117
RETEVMO ORAL CAPSULE 40 MG, 80 MG 261
RETEVMO ORAL TABLET 261
REVUFORJ 239
REZLIDHIA 204
REZUROCK 35
RIABNI 240
RINVOQ 310
RINVOQ LQ 310
ROLVEDON 100
ROMVIMZA 321
ROZLYTREK ORAL CAPSULE 100 MG, 200 MG 111
ROZLYTREK ORAL PELLETS IN PACKET 111
RUBRACA 254
RYBELSUS 140
RYDAPT 181
RYSTIGGO 253

sajazir 148
SCEMBLIX 23
scopolamine base 144
SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG 270
SIGNIFOR 211
sildenafil (pulm.hypertension) oral tablet 214
SIRTURO 33
SKYRIZI SUBCUTANEOUS PEN INJECTOR 248
SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML, 75 MG/0.83 ML 248
SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR 248
sofosbuvir-velpatasvir 266
SOMAVERT 218
sorafenib 272
STELARA INTRAVENOUS 313
STELARA SUBCUTANEOUS SOLUTION 313
STELARA SUBCUTANEOUS SYRINGE 45 MG/0.5 ML, 90 MG/ML 313
STIVARGA 236
sunitinib malate 275
SUNOSI 267
SYMDEKO 293
SYMLINPEN 120 232
SYMLINPEN 60 232
SYMPAZAN 61
SYNRIBO 205
TABRECTA 55
tadalafil (pulm. hypertension) 214
tadalafil oral tablet 2.5 mg, 5 mg 277
TAFINLAR ORAL CAPSULE 66
TAFINLAR ORAL TABLET FOR SUSPENSION 66
TAGRISSO 208
TAKHZYRO 158
TALZENNA ORAL CAPSULE 0.1 MG, 0.25 MG, 0.35 MG, 0.5 MG, 0.75 MG, 1 MG 279
tasimelteon 281
TAVALISSE 130
TAVNEOS 25
TAZVERIK 282
TECVAYLI 283
temazepam oral capsule 15 mg, 30 mg 146
TEPEZZA 288
TEPMETKO 287
teriflunomide 289
testosterone cypionate 291
testosterone enanthate 291
testosterone transdermal gel in metered-dose pump 12.5 mg/ 1.25 gram (1 %), 20.25 mg/1.25 gram (1.62 %) 291
testosterone transdermal gel in packet 1 % (25 mg/2.5gram), 1 % (50 mg/5 gram) 291
testosterone transdermal solution in metered pump w/app 291
tetrabenazine 292
TEVIMBRA 296
THALOMID 294
THIOLA EC 222
TIBSOVO 156
tiopronin oral tablet,delayed release (dr/ec) 222
tolvaptan (polycys kidney dis) oral tablets, sequential 299
TRACLEER ORAL TABLET 109
TRACLEER ORAL TABLET FOR SUSPENSION 109
TREMFYA INTRAVENOUS 142
TREMFYA ONE-PRESS 142
TREMFYA PEN INDUCTION PK(2PEN) 142
TREMFYA PEN SUBCUTANEOUS PEN INJECTOR 200 MG/2 ML 142
TREMFYA SUBCUTANEOUS SYRINGE 142
tretinoin 300
tridacaine ii 169
trientine 305
trihexyphenidyl 144
TRIKAFTA ORAL GRANULES IN PACKET, SEQUENTIAL 104
TRIKAFTA ORAL TABLETS, SEQUENTIAL 104
TRULICITY 140
TRUQAP 53
TRUSELTIQ 152
TUKYSA ORAL TABLET 150 MG, 50 MG 307
TURALIO 224
TYMLOS 1

UBRELVY 308
UDENYCA 215
UDENYCA AUTOINJECTOR 215
UDENYCA ONBODY 215
ULTOMIRIS 309
UPTRAVI ORAL TABLET 1,000 MCG, 1,200 MCG, 1,400 MCG, 1,600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG 258
UPTRAVI ORAL TABLETS,DOSE PACK 258
VANFLYTA 235
VENCLEXTA ORAL TABLET 10 MG, 100 MG, 50 MG 317
VENCLEXTA STARTING PACK 317
VEOZAH 126
VERQUVO 318
VERZENIO 5
vigabatrin 320
vigadrone oral powder in packet 320
VIGAFYDE 320
vigpoder 320
VIJOICE ORAL GRANULES IN PACKET 14
VITRAKVI ORAL CAPSULE 100 MG, 25 MG 161
VITRAKVI ORAL SOLUTION 161
VIZIMPRO 67
VONJO 209
VORANIGO ORAL TABLET 10 MG, 40 MG 323
voriconazole oral suspension for reconstitution 324
VOTRIENT 212
VOWST 122
VUMERITY 90
VYLOY 329
VYNDAMAX 278
VYNDAQEL 278
VYVGART 98
VYVGART HYTRULO 98
WELIREG 36
XALKORI ORAL CAPSULE 64
XALKORI ORAL PELLET 64
XDEMVY 175
XELJANZ ORAL SOLUTION 298
XELJANZ ORAL TABLET 298
XELJANZ XR 298
XERMELO 286
XGEVA 84
XIFAXAN ORAL TABLET 200 MG, 550 MG 243
XOLAIR 206
XOSPATA 136
XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40 MG/WEEK (10 MG X 4), 40 MG/WEEK (40 MG X 1), 40MG TWICE WEEK (40 MG X 2), 60 MG/WEEK (60 MG X 1), 60MG TWICE WEEK (120 MG/WEEK), 80 MG/WEEK (40 MG X 2), 80MG TWICE WEEK (160 MG/WEEK) 260
XTANDI ORAL CAPSULE 112
XTANDI ORAL TABLET 40 MG, 80 MG 112
XURIDEN 312
XYOSTED 291
XYREM 265
YONSA 7
ZARXIO 127
ZEJULA ORAL CAPSULE 195
ZEJULA ORAL TABLET 195
ZELBORAF 316
ZIIHERA 326
ZIRABEV 43
ZTALMY 134
ZTLIDO 169
ZURZUVAE ORAL CAPSULE 20 MG, 25 MG, 30 MG 331
ZYDELIG 149
ZYKADIA 58