ABALOPARATIDE

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	ONE OF THE FOLLOWING: (1) HIGH RISK FOR FRACTURES DEFINED AS ONE OF THE FOLLOWING: HISTORY OF OSTEOPOROTIC (I.E., FRAGILITY, LOW TRAUMA) FRACTURE(S). 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.). 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) ADEQUATE TRIAL OF, INTOLERANCE TO, OR A CONTRAINDICATION TO BISPHOSPHONATES (E.G., ALENDRONATE, RISEDRONATE, IBANDRONATE).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ABATACEPT IV

Products Affected

- ORENCIA (WITH MALTOSE)

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, JUVENILE IDIOPATHIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: RA: 6 MOS. JIA: 4 MOS. PSA: 12 MOS. RENEWAL: 12 MOS ALL INDICATIONS
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. JUVENILE IDIOPATHIC ARTHRITIS (JIA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX.

ABATACEPT SQ

Products Affected

- ORENCIA
- ORENCIA CLICKJECT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, JUVENILE IDIOPATHIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. JUVENILE IDIOPATHIC ARTHRITIS (JIA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX.

ABEMACICLIB

Products Affected

- VERZENIO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINDICATION TO IBRANCE (PALBOCICLIB) WHEN REQUEST IS FOR COMBINATION THERAPY WITH FULVESTRANT FOR HORMONE RECEPTOR (HR)-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER.

ABIRATERONE

Products Affected

- ZYTIGA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ABIRATERONE SUBMICRONIZED

Products Affected

- YONSA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ACALABRUTINIB

Products Affected

- CALQUENCE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ADALIMUMAB

Products Affected

- HUMIRA
- HUMIRA PEDIATRIC CROHN'S START
- HUMIRA PEN
- HUMIRA PEN CROHN'S-UC-HS START
- HUMIRA PEN PSORIASIS-UVEITIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS: CURRENT WEIGHT. PLAQUE PSORIASIS: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSORIASIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. CROHN'S DISEASE/ULCERATIVE COLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL:RA:6 MO PSA/AS:4 MO PJIA:5 MO PSO/CD/UC/HS:3 MO UVEITIS:6 MO RENEWAL:12 MO ALL INDICATIONS

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PREVIOUS TRIAL OF ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. ANKYLOSING SPONDYLITIS: TRIAL OF FORMULARY AGENTS NOT REQUIRED. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF ONE OF THE FOLLOWING CONVENTIONAL THERAPIES SUCH AS PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF ONE CONVENTIONAL AGENT SUCH AS A CORTICOSTEROID (I.E., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE. ULCERATIVE COLITIS (UC): PREVIOUS TRIAL OF ONE CONVENTIONAL AGENT SUCH AS A CORTICOSTEROID (I.E., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE.

AFATINIB DIMALEATE

Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ALECTINIB

Products Affected

- ALECENSA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ALEMTUZUMAB - LEMTRADA

Products Affected

- LEMTRADA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 1 MONTH. RENEWAL: 12 MONTHS.
Other Criteria	TRIAL WITH TWO OF THE FOLLOWING AGENTS FOR MULTIPLE SCLEROSIS: AUBAGIO, AVONEX, GILENYA, PLEGRIDY, REBIF, TECFIDERA, OR GLATIRAMER. RENEWAL REQUESTS FOR ALEMTUZUMAB REQUIRE THAT AT LEAST 12 MONTHS HAVE ELAPSED SINCE RECEIVING THE FIRST COURSE OF LEMTRADA. PATIENTS ARE LIMITED TO TWO COURSES OF THERAPY WITH LEMTRADA WITHIN A LIFETIME.

ALIROCUMAB

Products Affected

- PRALUENT PEN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE AND OLDER.
Prescriber Restrictions	CARDIOLOGIST, ENDOCRINOLOGIST OR LIPIDOLOGIST
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	MUST HAVE A LDL CHOLESTEROL LEVEL GREATER THAN 100MG/DL WHILE ON MAXIMAL DRUG TREATMENT FOR THE PAST 2 MONTHS AND ONE OF THE FOLLOWING DIAGNOSES: (1) HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH) DETERMINED BY SIMON BROOME DIAGNOSTIC CRITERIA FOR HEFH OR A SCORE OF 6 OR GREATER ON THE DUTCH LIPID NETWORK CRITERIA FOR HEFH OR (2) HISTORY OF ATHEROSCLEROTIC CARDIOVASCULAR DISEASE (ASCVD) AS DOCUMENTED BY PHYSICIAN ATTESTATION. PATIENT MUST NOT HAVE CONCURRENT USE OF REPATHA OR OTHER PCSK9 AGENT. INITIAL THERAPY: FOR STATIN TOLERANT PATIENTS: MUST HAVE TAKEN ATORVASTATIN OR ROSUVASTATIN FOR THE PAST 2 MONTHS. FOR STATIN INTOLERANT PATIENTS: DOCUMENTATION OF STATIN INTOLERANCE BY ONE OF THE FOLLOWING: (1) PHYSICIAN ATTESTATION, (2) PATIENT HAS TRIED ROSUVASTATIN, ATORVASTATIN, OR STATIN THERAPY AT ANY DOSE AND HAS EXPERIENCED SKELETAL-MUSCLE RELATED SYMPTOMS (E.G., MYOPATHY). PATIENTS WITH CONTRAINDICATIONS TO STATINS INCLUDING ACTIVE DECOMPENSATED LIVER DISEASE, NURSING FEMALE, PREGNANCY OR PLANS TO BECOME PREGNANT OR HYPERSENSITIVITY REACTIONS WILL BE APPROVED FOR PRALUENT THERAPY WITHOUT REQUIREMENT OF DOCUMENTATION OF STATIN INTOLERANCE. RENEWAL CRITERIA: RECEIVING PRIOR PRALUENT THERAPY FOR THE PAST 6 MONTHS AND NO CLAIMS FOR REPATHA, JUXTAPID, OR KYNAMRO SINCE PRALUENT APPROVAL.

AMANTADINE

Products Affected

- GOCOVRI ORAL
  CAPSULE,EXTENDED RELEASE
  24HR 137 MG, 68.5 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ANAKINRA

Products Affected

- KINERET

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RHEUMATOID ARTHRITIS (RA) RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: RA: 6 MONTHS NOMID/CAPS: 12 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA.

APALUTAMIDE

Products Affected

- ERLEADA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

APREMILAST

Products Affected

- OTEZLA
- OTEZLA STARTER

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSORIASIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST
Coverage Duration	INITIAL: PSORIATIC ARTHRITIS: 4 MONTHS. PSORIASIS: 5 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA AND ONE CONVENTIONAL THERAPY SUCH AS PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE.

ASFOTASE

Products Affected

- STRENSIQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	NON-SPECIFIC ALKALINE PHOSPHATASE (TNSALP) (ALPL) GENE MUTATION, SERUM ALKALINE PHOSPHATASE (ALP) LEVEL, SERUM PYRIDOXAL-5'-PHOSPHATE (PLP) LEVELS, URINE PHOSPHOETHANOLAMINE (PEA) LEVEL, RADIOGRAPHIC EVIDENCE OF HYPOPHOSPHATASIA (HPP)
Age Restrictions	PERINATAL/INFANTILE-ONSET HYPOPHOSPHATASIA (HPP): 6 MONTHS OF AGE OR YOUNGER AT HYPOPHOSPHATASIA (HPP) ONSET. JUVENILE-ONSET HYPOPHOSPHATASIA (HPP): 18 YEARS OF AGE OR YOUNGER AT HYPOPHOSPHATASIA (HPP) ONSET.
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN ENDOCRINOLOGIST, A GENETICIST, OR A METABOLIC SPECIALIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: FOR PATIENTS WITH PERINATAL/INFANTILE-ONSET HYPOPHOSPHATASIA (HPP), ALL OF THE FOLLOWING CRITERIA MUST BE MET: POSITIVE FOR A TISSUE NON-SPECIFIC ALKALINE PHOSPHATASE (TNSALP) (ALPL) GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR MEETS AT LEAST TWO OF THE FOLLOWING CRITERIA: 1.) SERUM ALKALINE PHOSPHATASE (ALP) LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE 2.) SERUM PYRIDOXAL-5'-PHOSPHATE (PLP) LEVELS ELEVATED AND PATIENT HAS NOT RECEIVED VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK 3.) URINE PHOSPHOETHANOLAMINE (PEA) LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE 4.) RADIOGRAPHIC EVIDENCE OF HYPOPHOSPHATASIA (HPP) (E.G., FLARED AND FRAYED METAPHYSES, OSTEOPENIA, WIDENED GROWTH PLATES, AREAS OF RADIOLUCENCY OR SCLEROSIS) 5.) PRESENCE OF TWO OR MORE OF THE FOLLOWING: RACHITIC CHEST DEFORMITY, CRANIOSYNOSTOSIS (PREMATURE CLOSURE OF SKULL BONES), DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, HISTORY OF VITAMIN B6 DEPENDENT SEIZURES, NEPHROCALCINOSIS, OR HISTORY OF ELEVATED SERUM CALCIUM. HISTORY OR PRESENCE OF NON-TRAUMATIC POSTNATAL FRACTURE AND DELAYED FRACTURE HEALING. FOR PATIENTS WITH JUVENILE-ONSET HYPOPHOSPHATASIA (HPP), ALL OF THE FOLLOWING CRITERIA MUST BE MET: POSITIVE FOR A TISSUE NON-SPECIFIC ALKALINE PHOSPHATASE (TNSALP) (ALPL) GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR MEETS AT LEAST TWO OF THE FOLLOWING CRITERIA: 1.) SERUM ALKALINE PHOSPHATASE (ALP) LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE 2.) SERUM PYRIDOXAL-5'-PHOSPHATE (PLP) LEVELS ELEVATED AND PATIENT HAS NOT RECEIVED VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK 3.)URINE PHOSPHOETHANOLAMINE (PEA) LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE 4.)RADIOGRAPHIC EVIDENCE OF HYPOPHOSPHATASIA (HPP) (E.G., FLARED AND FRAYED METAPHYSES, OSTEOPENIA, OSTEOMALACIA, WIDENED GROWTH PLATES, AREAS OF RADIOLUCENCY OR SCLEROSIS) 5.)PRESENCE OF TWO OR MORE OF THE FOLLOWING:RACHITIC DEFORMITIES (RACHITIC CHEST,

PA Criteria	Criteria Details
	BOWED LEGS, KNOCK-KNEES),PREMATURE LOSS OF PRIMARY TEETH PRIOR TO 5 YEARS OF AGE, DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, OR HISTORY OR PRESENCE OF NON-TRAUMATIC FRACTURES OR DELAYED FRACTURE HEALING. STRENSIQ WILL NOT BE APPROVED FOR THE FOLLOWING PATIENTS: PATIENTS CURRENTLY RECEIVING TREATMENT WITH A BISPHOSPHONATE [E.G., BONIVA (IBANDRONATE), FOSAMAX (ALENDRONATE), ACTONEL (RISEDRONATE)], PATIENTS WITH SERUM CALCIUM OR PHOSPHATE LEVELS BELOW THE NORMAL RANGE, PATIENTS WITH A TREATABLE FORM OF RICKETS. RENEWAL: PATIENT HAS EXPERIENCED AN IMPROVEMENT IN THE SKELETAL CHARACTERISTICS OF HYPOPHOSPHATASIA (HPP) (E.G., IMPROVEMENT OF THE IRREGULARITY OF THE PROVISIONAL ZONE OF CALCIFICATION, PHYSEAL WIDENING, METAPHYSEAL FLARING, RADIOLUCENCIES, PATCHY OSTEOSCLEROSIS, RATIO OF MID-DIAPHYSEAL CORTEX TO BONE THICKNESS, GRACILE BONES, BONE FORMATION AND FRACTURES.

ASPARAGINASE

Products Affected

- ONCASPAR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 MONTHS
Other Criteria	

ATEZOLIZUMAB

Products Affected

- TECENTRIQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

AVATROMBOPAG

Products Affected

- DOPTELET

PA Criteria	Criteria Details
Covered Uses	ALL FDA-APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 MONTHS
Other Criteria	

AVELUMAB

Products Affected

- BAVENCIO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

AXITINIB

Products Affected

- INLYTA ORAL TABLET 1 MG, 5 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF AT LEAST ONE SYSTEMIC THERAPY FOR THE TREATMENT OF RCC SUCH AS NEXAVAR (SORAFENIB), TORISEL (TEMSIROLIMUS), SUTENT (SUNITINIB), VOTRIENT (PAZOPANIB), OR AVASTIN (BEVACIZUMAB) IN COMBINATION WITH INTERFERON.

BARICITINIB

Products Affected

- OLUMIANT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PREVIOUS TRIAL OF OR CONTRAINDICATION TO HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, XELJANZ XR, CIMZIA, OR ACTEMRA.

BECAPLERMIN

Products Affected

- REGRANEX

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	NON-DIABETIC ULCERS, NEOPLASM AT APPLICATION SITE, PRESSURE OR VENOUS STASIS ULCERS AND ULCERS THAT DO NOT EXTEND THROUGH THE DERMIS.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	VASCULAR SURGEON, PODIATRIST, ENDOCRINOLOGIST, PHYSICIAN PRACTICING IN A SPECIALTY WOUND CLINIC OR INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	3 MONTHS
Other Criteria	

BEDAQUILINE FUMARATE

Products Affected

- SIRTURO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE AND OLDER.
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.

BELIMUMAB

Products Affected

- BENLYSTA INTRAVENOUS
- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	AUTOANTIBODY POSITIVE LUPUS TEST.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: MEMBER IS CURRENTLY TAKING CORTICOSTEROIDS, ANTIMALARIALS, NSAIDS, OR IMMUNOSUPPRESSIVE AGENTS. NO APPROVAL FOR DIAGNOSIS OF SEVERE ACTIVE LUPUS NEPHRITIS, SEVERE CENTRAL NERVOUS SYSTEM LUPUS OR CONCURRENT USE OF BIOLOGIC AGENTS OR INTRAVENOUS CYCLOPHOSPHAMIDE. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.

BELINOSTAT

Products Affected

- BELEODAQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BENDAMUSTINE

Products Affected

- BENDEKA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BENRALIZUMAB

Products Affected

- FASENRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BEVACIZUMAB

Products Affected

- AVASTIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BEXAROTENE

Products Affected

- *bexarotene*
- TARGRETIN TOPICAL

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BLINATUMOMAB

Products Affected

- BLINCYTO INTRAVENOUS KIT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: RELAPSED OR REFRACTORY B-CELL: 3 MOS. MRD-POSITIVE B-CELL: 2 MOS. RENEWAL: 12 MOS.
Other Criteria	INITIAL: RELAPSED OR REFRACTORY B-CELL PRECURSOR ALL: APPROVAL IS FOR 2 CYCLES, MAY APPROVE FOR 1 ADDITIONAL CYCLE DUE TO TREATMENT INTERRUPTION FOR DOSE MODIFICATION. RENEWAL: FOR DIAGNOSIS OF RELAPSED OR REFRACTORY B-CELL PRECURSOR ACUTE LYMPHOBLASTIC LEUKEMIA (ALL), RENEWAL IS APPROVED FOR PATIENTS WHO HAVE ACHIEVED COMPLETE REMISSION (CR) OR CR WITH PARTIAL HEMATOLOGICAL RECOVERY OF PERIPHERAL BLOOD COUNTS AFTER 2 CYCLES OF TREATMENT. RENEWAL IS NOT APPROVED FOR PATIENTS WHO RECEIVED AN ALLOGENEIC HEMATOPOIETIC STEM-CELL TRANSPLANT. FOR DIAGNOSIS OF MINIMAL RESIDUAL DISEASE (MRD)-POSITIVE B-CELL PRECURSOR ACUTE LYMPHOBLASTIC LEUKEMIA (ALL), RENEWAL IS APPROVED FOR PATIENTS WHO HAVE ACHIEVED UNDETECTABLE MINIMAL RESIDUAL DISEASE (MRD) WITHIN ONE CYCLE OF BLINCYTO TREATMENT AND IS RELAPSE-FREE (I.E., HEMATOLOGICAL OR EXTRAMEDULLARY RELAPSE, OR SECONDARY LEUKEMIA).

BORTEZOMIB

Products Affected

- BORTEZOMIB
- VELCADE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BOSUTINIB

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CML: BCR-ABL MUTATIONAL ANALYSIS CONFIRMING THAT BOTH T315I AND V299L MUTATIONS ARE NOT PRESENT.

BOTULINUM NEUROTOXIN

Products Affected

- BOTOX INJECTION RECON SOLN 100 UNIT, 200 UNIT
- MYOBLOC INTRAMUSCULAR SOLUTION 10,000 UNIT/2 ML, 2,500 UNIT/0.5 ML, 5,000 UNIT/ML

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	COSMETIC DIAGNOSIS: WRINKLES.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MIGRAINE HEADACHE: TRIAL OF TWO OF THE FOLLOWING: BETA BLOCKERS, TRICYCLIC ANTIDEPRESSANTS, OR VALPROIC ACID. OVERACTIVE BLADDER: TRIAL OF OR CONTRAINDICATION TO THE USE OF ONE ANTICHOLINERGIC MEDICATION SUCH AS ORAL OXYBUTYNIN, ORAL OXYBUTYNIN ER, TOLTERODINE, TOLTERODINE ER, TOVIAZ, TROSPIUM, OR TROSPIUM ER. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

BRIGATINIB

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

BRODALUMAB

Products Affected

- SILIQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: COSENTYX OR OTEZLA. PATIENT HAS BEEN COUNSELED ON AND EXPRESSES UNDERSTANDING OF THE RISK OF SUICIDAL IDEATION AND BEHAVIOR. RENEWAL: PATIENT HAS NOT DEVELOPED OR REPORTED WORSENING DEPRESSIVE SYMPTOMS OR SUICIDAL IDEATION AND BEHAVIORS WHILE ON TREATMENT WITH SILIQ.

C1 ESTERASE INHIBITOR

Products Affected

- CINRYZE
- HAEGARDA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HEMATOLOGIST, IMMUNOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	

CABOZANTINIB

Products Affected

- COMETRIQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

CABOZANTINIB S-MALATE - CABOMETYX

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

CANAKINUMAB

Products Affected

- ILARIS (PF)

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	CAPS: 4 YEARS AND OLDER. SJIA: 2 YEARS AND OLDER.
Prescriber Restrictions	PRESCRIBED OR SUPERVISED BY RHEUMATOLOGIST, DERMATOLOGIST, OR AN IMMUNOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	

CANNABINOIDS

Products Affected

- *dronabinol*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 SUCH AS ONDANSETRON, STEROIDS INDICATED FOR EMESIS OR EMEND. NO ADDITIONAL REQUIREMENTS FOR A DIAGNOSIS OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS.

CANNABINOIDS ORAL SOLUTION

Products Affected

- SYNDROS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 FORMULARY DRONABINOL CAPSULES AND ONE CONVENTIONAL ANTIEMETIC THERAPY SUCH AS ONDANSETRON, STEROIDS INDICATED FOR EMESIS OR APREPITANT. PART D COVERAGE CONSIDERATION FOR A DIAGNOSIS OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS REQUIRES A TRIAL OF OR CONTRAINDICATION TO FORMULARY DRONABINOL CAPSULES.

CARFILZOMIB

Products Affected

- KYPROLIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

CERITINIB

Products Affected

- ZYKADIA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	POSITIVE FOR ANAPLASTIC LYMPHOMA KINASE (ALK) FUSION ONCOGENE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

CERTOLIZUMAB PEGOL

Products Affected

- CIMZIA
- CIMZIA POWDER FOR RECONST

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS/ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. CROHN'S DISEASE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: RA: 6 MONTHS. PSA/AS: 4 MONTHS. CD: 12 MONTHS. RENEWAL: 12 MONTHS FOR ALL DIAGNOSES.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. ANKYLOSING SPONDYLITIS: PREVIOUS TRIAL OF HUMIRA. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF HUMIRA AND ONE CONVENTIONAL AGENT SUCH AS A CORTICOSTEROID (I.E., BUDESONIDE, METHYLPREDNISOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE.

CLOBAZAM

Products Affected

- ONFI ORAL SUSPENSION
- ONFI ORAL TABLET 10 MG, 20 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	2 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF LAMOTRIGINE OR TOPIRAMATE. REQUESTS FOR ORAL SUSPENSION APPROVABLE IF PATIENT IS UNABLE TO SWALLOW OR IS UNDER THE AGE OF 5 YEARS.

COBIMETINIB FUMARATE

Products Affected

- COTELLIC

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

COPANLISIB DI-HCL

Products Affected

- ALIQOPA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

CORTICOTROPIN

Products Affected

- ACTHAR H.P.

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INFANTILE SPASMS AND MULTIPLE SCLEROSIS: 28 DAYS. OTHER FDA APPROVED INDICATIONS:12 MONTHS.
Other Criteria	ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS REQUIRE A TRIAL OF OR CONTRAINDICATION TO IV CORTICOSTEROIDS. NOT APPROVED FOR DIAGNOSTIC PURPOSES.

CRIZOTINIB

Products Affected

- XALKORI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DABRAFENIB MESYLATE

Products Affected

- TAFINLAR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DACLATASVIR

Products Affected

- DAKLINZA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
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. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI, OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. NO APPROVALS FOR CONCURRENT USE WITH ANY OF THESE (CONTRAINDICATED OR NOT RECOMMENDED BY THE MANUFACTURER) MEDICATIONS: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, OR RIFAMPIN.

DALFAMPRIDINE

Products Affected

- AMPYRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	WALKING DISABILITY SUCH AS MILD TO MODERATE BILATERAL LOWER EXTREMITY WEAKNESS OR UNILATERAL WEAKNESS PLUS LOWER EXTREMITY OR TRUNCAL ATAXIA.
Age Restrictions	
Prescriber Restrictions	NEUROLOGIST
Coverage Duration	INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT IN WALKING ABILITY.

DARATUMUMAB

Products Affected

- DARZALEX

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DASATINIB

Products Affected

- SPRYCEL ORAL TABLET 100 MG, 140 MG, 20 MG, 50 MG, 70 MG, 80 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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.

DEFERASIROX

Products Affected

- EXJADE
- JADENU
- JADENU SPRINKLE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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. RENEWAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 500 MCG/L. NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) INITIAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L AND LIVER IRON CONCENTRATION (LIC) OF 5 MG FE/G DRY WEIGHT OR GREATER. RENEWAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L OR LIC OF 3 MG FE/G DRY WEIGHT OR GREATER

DEFERIPRONE

Products Affected

- FERRIPROX

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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	INITIAL CRITERIA: REQUIRES TRIAL OF EXJADE, JADENU, OR GENERIC DEFEROXAMINE AND ONE OF THE FOLLOWING CRITERIA 1) PHYSICIAN ATTESTATION THAT PATIENT IS EXPERIENCING INTOLERABLE TOXICITIES, CLINICALLY SIGNIFICANT ADVERSE EFFECTS, OR CONTRAINDICATION TO THESE THERAPIES OR 2) INADEQUATE CHELATION DEFINED BY ONE OF THE FOLLOWING: A) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 2500 MCG/L OR B) EVIDENCE OF CARDIAC IRON ACCUMULATION. RENEWAL: SERUM FERRITIN LEVELS MUST BE CONSISTENTLY ABOVE 500MCG/L

DEFEROXAMINE

Products Affected

- *deferoxamine*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	AT LEAST 3 YEARS OF AGE OR OLDER
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEMATOLOGIST OR HEMATOLOGIST/ONCOLOGIST
Coverage Duration	INITIAL: 6 MONTHS RENEWAL: 12 MONTHS
Other Criteria	INITIAL: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 1000MCG/L RENEWAL: SERUM FERRITIN LEVELS MUST BE CONSISTENTLY ABOVE 500MCG/L

DEFLAZACORT

Products Affected

- EMFLAZA ORAL SUSPENSION MG, 36 MG, 6 MG
- EMFLAZA ORAL TABLET 18 MG, 30

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PHYSICIAN ATTESTATION OF GENETIC TESTING CONFIRMING DMD DIAGNOSIS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL CRITERIA: REQUIRE TRIAL OF PREDNISONE OR PREDNISOLONE FOR AT LEAST 6 MONTHS 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 CRITERIA ARE MET: A) PATIENT IS NOT IN STAGE 1 (PRE-SYMPTOMATIC PHASE) B) STEROID MYOPATHY HAS BEEN RULED OUT C) PHYSICIAN ATTESTATION OF DETERIORATION IN AMBULATION, FUNCTIONAL STATUS, OR PULMONARY FUNCTION CONSISTENT WITH ADVANCING DISEASE. RENEWAL CRITERIA: 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)

DELAFLOXACIN

Products Affected

- BAXDELA ORAL

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ONE MONTH
Other Criteria	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST OR ABSSSI ORGANISM ANTIMICROBIAL SUSCEPTIBILITY TESTING SHOWS SUSCEPTIBILITY TO DELAFLOXACIN AND RESISTANCE TO ONE PREFERRED FORMULARY STANDARD OF CARE AGENT OR IF SENSITIVITY RESULTS ARE UNAVAILABLE: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED FORMULARY AGENTS: A PENICILLIN, A FLUOROQUINOLONE, A CEPHALOSPORIN, OR A GRAM POSITIVE TARGETING ANTIBIOTIC

DENOSUMAB-XGEVA

Products Affected

- XGEVA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DESIRUDIN

Products Affected

- IPRIVASK

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 MONTH
Other Criteria	

DEUTETRABENAZINE

Products Affected

- AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DICHLORPHENAMIDE

Products Affected

- KEVEYIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	HEPATIC INSUFFICIENCY, PULMONARY OBSTRUCTION, OR A HEALTH CONDITION THAT WARRANTS CONCURRENT USE OF HIGH-DOSE ASPIRIN
Required Medical Information	
Age Restrictions	18 YEARS AND OLDER
Prescriber Restrictions	
Coverage Duration	INITIAL: 2 MONTHS RENEWAL: 12 MONTHS
Other Criteria	RENEWAL REQUIRES THE PATIENT EXPERIENCED AT LEAST TWO FEWER ATTACKS PER WEEK FROM THEIR BASELINE

DICLOFENAC EPOLAMINE

Products Affected

- FLECTOR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

DICLOFENAC TOPICAL

Products Affected

- *diclofenac sodium topical gel 3 %* METERED-DOSE PUMP
- PENNSAID TOPICAL SOLUTION IN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DIMETHYL FUMARATE

Products Affected

- TECFIDERA ORAL CAPSULE,DELAYED RELEASE(DR/EC) 120 MG, 120 MG (14)- 240 MG (46), 240 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DINUTUXIMAB

Products Affected

- UNITUXIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

DROXIDOPA

Products Affected

- NORTHERA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 HAD AN INCREASE IN SYSTOLIC BLOOD PRESSURE FROM BASELINE OF AT LEAST 10 MMHG UPON STANDING FROM A SUPINE (LYING FACE UP) POSITION.

DUPILUMAB

Products Affected

- DUPIXENT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PATIENT HAS MINIMUM BODY SURFACE AREA (BSA) INVOLVEMENT OF AT LEAST 10%, ECZEMA AREA AND SEVERITY INDEX (EASI) SCORE OF AT LEAST 16, OR PHYSICIAN GLOBAL ASSESSMENT (PGA) SCORE OF AT LEAST 3. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST OR IMMUNOLOGIST
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL:12 MONTHS
Other Criteria	INITIAL: PREVIOUS TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING: TOPICAL CORTICOSTEROIDS, TOPICAL CALCINEURIN INHIBITORS [E.G., ELIDEL (PIMECROLIMUS), GENERIC TACROLIMUS OINTMENT], OR TOPICAL PDE4 INHIBITOR [E.G., EUCRISA (CRISABOROLE)].

DURVALUMAB

Products Affected

- IMFINZI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

EDARAVONE

Products Affected

- RADICAVA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ELBASVIR/GRAZOPREVIR

Products Affected

- ZEPATIER

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
Exclusion Criteria	MODERATE OR SEVERE LIVER IMPAIRMENT (CHILD PUGH B OR C)
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS. FOR GENOTYPE 1A -TESTING FOR NS5A RESISTANCE-ASSOCIATED POLYMORPHISMS.
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.

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. NO CONCURRENT USE OF SOVALDI AND ANY OF THE FOLLOWING AGENTS: PHENYTOIN, CARBAMAZEPINE, RIFAMPIN, EFAVIRENZ, ATAZANAVIR, DARUNAVIR, LOPINAVIR, SAQUINAVIR, TIPRANAVIR, CYCLOSPORINE, NAFCILLIN, KETOCONAZOLE, MODAFINIL, BOSENTAN, ETRAVIRINE, ELVITEGRAVIR/COBICISTAT/EMTRICITABINE/TENOFOVIR, ATORVASTATIN AT DOSES GREATER THAN 20MG PER DAY OR ROSUVASTATIN AT DOSES GREATER THAN 10MG PER DAY.

ELIGLUSTAT TARTRATE

Products Affected

- CERDELGA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ELOSULFASE ALFA

Products Affected

- VIMIZIM

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	LIFETIME OF MEMBERSHIP IN PLAN.
Other Criteria	

ELOTUZUMAB

Products Affected

- EMPLICITI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ELTROMBOPAG

Products Affected

- PROMACTA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ITP:INITIAL: 2MO.RENEW:AFTER RESPONSE:12MO, INADEQUATE DOSE:2MO.HCV:12MO.SEVERE APLASTIC ANEMIA:12MO
Other Criteria	CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA PURPURA (ITP): INITIAL: TRIAL OF OR CONTRAINDICATION TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR AN INSUFFICIENT RESPONSE TO SPLENECTOMY. ITP: RENEWAL: PATIENT HAS A CLINICAL RESPONSE AS DEFINED BY AN INCREASE IN PLATELET COUNT OF GREATER THAN OR EQUAL TO 50 X10^9/L (GREATER THAN OR EQUAL TO 50,000 PER UL) AT THE MAX DOSE OF 75MG PER DAY FOR 4 WEEKS. HEPATITIS C: CONCURRENT INTERFERON THERAPY.

ENASIDENIB

Products Affected

- IDHIFA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ENDOTHELIN RECEPTOR ANTAGONISTS

Products Affected

- LETAIRIS
- OPSUMIT
- TRACLEER ORAL TABLET
- TRACLEER ORAL TABLET FOR SUSPENSION

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	INITIAL: MEAN PULMONARY ARTERY PRESSURE (PAP) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. LETAIRIS: PATIENT DOES NOT HAVE IDIOPATHIC PULMONARY FIBROSIS (IPF) TRACLEER: 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.

ENZALUTAMIDE

Products Affected

- XTANDI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINDICATION TO ZYTIGA (ABIRATERONE ACETATE) IS ALSO REQUIRED IN PATIENTS WHO DO NOT HAVE A CONTRAINDICATION OR INTOLERANCE TO PREDNISONE.

EPOPROSTENOL IV

Products Affected

- *epoprostenol (glycine)*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	COVERED UNDER LOCAL COVERAGE POLICY OF APPLICABLE MEDICARE DMERC.
Required Medical Information	FORMULARY DRUG ADMINISTERED IN A LONG TERM CARE FACILITY TO A PATIENT WHOSE PART A COVERAGE HAS EXPIRED OR FORMULARY DRUG NOT ADMINISTERED VIA AN IMPLANTABLE PUMP OR AN EXTERNAL PUMP OR DRUG ADMINISTERED VIA AN IMPLANTABLE PUMP/AN EXTERNAL PUMP. DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS III-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) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. RENEWAL: PATIENT HAS SHOWN 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.

ERLOTINIB

Products Affected

- TARCEVA ORAL TABLET 100 MG, 150 MG, 25 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ERYTHROPOIESIS STIMULATING AGENTS - EPOETIN ALFA

Products Affected

- EPOGEN 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
- PROCRIT 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
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. ADDITIONAL OFF LABEL ANEMIA IN HEPATITIS C BEING TREATED IN COMBINATION WITH RIBAVIRIN AND INTERFERON ALFA OR PEGINTERFERON ALFA.
Exclusion Criteria	

PA Criteria	Criteria Details
Required Medical Information	INITIAL: CHRONIC RENAL FAILURE (CRF) AND ANEMIA RELATED TO ZIDOVUDINE THERAPY REQUIRES A HEMOGLOBIN LEVEL OF LESS THAN 10G/DL. CANCER CHEMOTHERAPY REQUIRES A HEMOGLOBIN LEVEL OF LESS THAN 10G/DL ANEMIA DUE TO CONCURRENT HEPATITIS C TREATMENT WITH RIBAVIRIN PLUS INTERFERON ALFA/PEGINTERFERON ALFA REQUIRES A HEMOGLOBIN LEVEL LESS THAN 10G/DL AND RIBAVIRIN DOSE REDUCTION (UNLESS CONTRAINDICATED).ELECTIVE NON-CARDIAC OR NON-VASCULAR SURGERY REQUIRES A HEMOGLOBIN LEVEL LESS THAN 13G/DL. RENEWAL: CHRONIC RENAL FAILURE REQUIRES THAT THE PATIENT MEETS ONE OF THE FOLLOWING: IF THE PATIENT IS CURRENTLY RECEIVING DIALYSIS TREATMENT: 1) HEMOGLOBIN LEVEL OF LESS THAN 11G/DL OR 2) HEMOGLOBIN LEVEL THAT HAS REACHED 11G/DL AND DOSE REDUCTION/INTERRUPTION IS REQUIRED TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS. IF THE PATIENT IS NOT RECEIVING DIALYSIS TREATMENT: 1) HEMOGLOBIN LEVEL OF LESS THAN 10G/DL OR 2) HEMOGLOBIN LEVEL THAT HAS REACHED 10G/DL AND DOSE REDUCTION/INTERRUPTION IS REQUIRED TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS. ANEMIA DUE TO CONCURRENT HEPATITIS C TREATMENT WITH RIBAVIRIN PLUS INTERFERON ALFA/PEGINTERFERON ALFA, OR ANEMIA DUE TO ZIDOVUDINE THERAPY REQUIRES HEMOGLOBIN LEVELS BETWEEN 10G/DL AND 12G/DL. ANEMIA DUE TO EFFECT OF CONCOMITANTLY ADMINISTERED CANCER CHEMOTHERAPY REQUIRES A 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 MYELOSUPPRESSIVE CHEMO/CKD WITHOUT DIALYSIS/ZIDOVUDINE:12 MONTHS.SURGERY:1 MO.HCV:6 MOS.

PA Criteria	Criteria Details
Other Criteria	ALL INDICATIONS: TRIAL OF PROCRIT. PART D MEMBER RECEIVING DIALYSIS OR IDENTIFIED AS A PART D END STAGE RENAL DISEASE MEMBER: PAYS UNDER PART B.

ERYTHROPOIESIS STIMULATING AGENTS - MIRCERA

Products Affected

- MIRCERA INJECTION SYRINGE 100 MCG/0.3 ML, 200 MCG/0.3 ML, 50 MCG/0.3 ML, 75 MCG/0.3 ML

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: CHRONIC RENAL FAILURE REQUIRES HEMOGLOBIN LEVELS LESS THAN 10G/DL RENEWAL: CHRONIC RENAL FAILURE REQUIRES THAT THE PATIENT MEETS ONE OF THE FOLLOWING: IF THE PATIENT IS CURRENTLY RECEIVING DIALYSIS TREATMENT: 1) HEMOGLOBIN LEVEL OF LESS THAN 11G/DL OR 2) HEMOGLOBIN LEVEL THAT HAS REACHED 11G/DL AND DOSE REDUCTION/INTERRUPTION IS REQUIRED TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS. IF THE PATIENT IS NOT RECEIVING DIALYSIS TREATMENT: 1) HEMOGLOBIN LEVEL OF LESS THAN 10G/DL OR 2) HEMOGLOBIN LEVEL THAT HAS REACHED 10G/DL AND DOSE REDUCTION/INTERRUPTION IS REQUIRED TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ANEMIA DUE TO CKD WITH OR WITHOUT DIALYSIS: 12 MONTHS.
Other Criteria	TRIAL OF PROCRIT. PART D MEMBER RECEIVING DIALYSIS OR IDENTIFIED AS A PART D END STAGE RENAL DISEASE MEMBER: PAYS UNDER PART B.

ETANERCEPT

Products Affected

- ENBREL
- ENBREL SURECLICK

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING AT LEAST 5% BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS, PSORIATIC ARTHRITIS: 18 YEARS OR OLDER
Prescriber Restrictions	RHEUMATOID ARTHRITIS, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PLAQUE PSORIASIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: RA: 6 MONTHS. PJIA: 3 MONTHS. PSA/AS/PSO: 4 MONTHS. RENEWAL: 12 MONTHS FOR ALL DIAGNOSES

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA.POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA OR ACTEMRA. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA OR COSENTYX. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: COSENTYX OR OTEZLA.

ETEPLIRSEN

Products Affected

- EXONDYS 51

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PHYSICIAN ATTESTATION OF GENETIC TESTING CONFIRMING THAT MUTATION IN DUCHENNE MUSCULAR DYSTROPHY (DMD) GENE IS AMENABLE TO EXON 51 SKIPPING.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 24 WEEKS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL CRITERIA: PATIENT IS AMBULATORY AND IS CURRENTLY RECEIVING TREATMENT WITH OR HAS A CONTRAINDICATION TO CORTICOSTEROIDS. RENEWAL CRITERIA: 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) DURING THE PAST 24 WEEKS. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.

EVEROLIMUS

Products Affected

- AFINITOR DISPERZ MG, 5 MG, 7.5 MG
- AFINITOR ORAL TABLET 10 MG, 2.5

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 SUTENT OR NEXAVAR.

EVOLOCUMAB

Products Affected

- REPATHA PUSHTRONEX
- REPATHA SURECLICK
- REPATHA SYRINGE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CARDIOLOGIST, ENDOCRINOLOGIST OR LIPIDOLOGIST
Coverage Duration	INITIAL: 6 MONTHS RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	FOR HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH): MUST HAVE LDL LEVEL GREATER THAN 100MG/DL ON MAXIMAL DRUG TREATMENT (MDT) FOR AT LEAST 2 MONTHS WITHIN THE PAST 2 MONTHS AND ONE OF THE FOLLOWING: (1) HEFH DETERMINED BY SIMON BROOME DIAGNOSTIC (SBD) CRITERIA OR A SCORE OF 6 OR GREATER ON THE DUTCH LIPID NETWORK (DLN) CRITERIA. HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH): LDL LEVEL GREATER THAN 100MG/DL ON MDT FOR AT LEAST 2 MONTHS WITHIN THE PAST 2 MONTHS AND HOFH DETERMINED BY ONE OF THE FOLLOWING: 1) SBD CRITERIA, 2) A SCORE OF 8 OR GREATER ON THE DLN CRITERIA, OR 3) A 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. NO CONCURRENT USE OF OTHER PCSK9 INHIBITORS. INITIAL THERAPY: FOR STATIN TOLERANT PATIENTS: MUST HAVE TRIED MAXIMALLY TOLERATED DOSE OF HIGH INTENSITY STATIN SUCH AS ATORVASTATIN OR ROSUVASTATIN. FOR STATIN INTOLERANT PATIENTS WITH HEFH: ONE OF THE FOLLOWING MUST BE MET: PHYSICIAN ATTESTATION OF STATIN INTOLERANCE (INCLUDING BUT NOT LIMITED TO MYOPATHY), OR PATIENT HAS TRIED ROSUVASTATIN OR ATORVASTATIN AT ANY DOSE. PATIENTS WITH CONTRAINDICATIONS TO STATINS INCLUDING ACTIVE DECOMPENSATED LIVER DISEASE, NURSING FEMALE, PREGNANCY OR PLANS TO BECOME PREGNANT OR HYPERSENSITIVITY REACTIONS WILL BE APPROVED FOR REPATHA THERAPY WITHOUT DOCUMENTED STATIN INTOLERANCE. FOR STATIN INTOLERANT PATIENTS WITH HOFH: MUST BE ON MAX LIPID-LOWERING THERAPY INCLUDING ONE OF THE FOLLOWING: NIACIN, BILE ACID SEQUESTRANT, LOMITAPIDE OR MIPOMERSEN. QUALIFIERS MUST PROVIDE DOCUMENTATION OF STATIN INTOLERANCE TO ONE OF THE FOLLOWING: A HIGH INTENSITY STATIN (ROSUVASTATIN OR ATORVASTATIN) OR OTHER STATIN THERAPY AT ANY DOSE. STATIN INTOLERANT PATIENTS MUST BE ON MAXIMAL LIPID-LOWERING MEDICATION (NON-STATIN THERAPY) FOR AT LEAST 2 MONTHS WITHIN THE PAST 2 MONTHS WITH DOCUMENTATION OF

PA Criteria	Criteria Details
	STATIN INTOLERANCE TO ATORVASTATIN OR ROSUVASTATIN OR STATIN THERAPY AT ANY DOSE. DOCUMENTATION OF STATIN INTOLERANCE INCLUDES: (1) PHYSICIAN ATTESTATION, OR (2) PATIENT HAS TRIED ROSUVASTATIN OR ATORVASTATIN AND HAS EXPERIENCED SKELETAL MUSCLE RELATED EVENTS (E.G. MYOPATHY). RENEWAL CRITERIA: RECEIVING PRIOR REPATHA THERAPY FOR AT LEAST 6 MONTHS AND NOT ON CONCURRENT THERAPY WITH OTHER PCSK9 INHIBITORS, MIPOMERSEN, OR LOMITAPIDE.

FENTANYL NASAL SPRAY

Products Affected

- LAZANDA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CANCER: CURRENTLY ON A MAINTENANCE DOSE OF CONTROLLED-RELEASE OPIOID PAIN MEDICATION (SUCH AS MORPHINE SULFATE ER, OXYCODONE ER, OR FENTANYL). EITHER A TRIAL OR CONTRAINDICATION TO AT LEAST ONE (1) IMMEDIATE-RELEASE ORAL OPIOID PAIN AGENT (SUCH AS MORPHINE SULFATE IR, OXYCODONE/ASPIRIN, OXYCODONE/ACETAMINOPHEN, CODEINE/ACETAMINOPHEN, HYDROMORPHONE, OR MEPERIDINE) OR MEMBER HAS DIFFICULTY SWALLOWING TABLETS/CAPSULES AND TRIAL OR CONTRAINDICATION TO GENERIC FENTANYL CITRATE LOZENGE. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

FENTANYL TRANSMUCOSAL AGENTS - FENTANYL CITRATE

Products Affected

- *fentanyl citrate*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CANCER: CURRENTLY ON A MAINTENANCE DOSE OF CONTROLLED-RELEASE OPIOID PAIN MEDICATION (SUCH AS MORPHINE SULFATE ER, OXYCODONE ER, OR FENTANYL). EITHER A TRIAL OR CONTRAINDICATION TO AT LEAST ONE (1) IMMEDIATE-RELEASE ORAL OPIOID PAIN AGENT (SUCH AS MORPHINE SULFATE IR, OXYCODONE/ASPIRIN, OXYCODONE/ACETAMINOPHEN, CODEINE/ACETAMINOPHEN, HYDROMORPHONE, OR MEPERIDINE) OR MEMBER HAS DIFFICULTY SWALLOWING TABLETS/CAPSULES. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

FINGOLIMOD

Products Affected

- GILENYA ORAL CAPSULE 0.25 MG, 0.5 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

FOSTAMATINIB DISODIUM

Products Affected

- TAVALISSE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

GEFITINIB

Products Affected

- IRESSA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

GEMTUZUMAB OZOGAMICIN

Products Affected

- MYLOTARG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

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
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

GLECAPREVIR/PIBRENTASVIR

Products Affected

- MAVYRET

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE.
Exclusion Criteria	MODERATE OR SEVERE HEPATIC IMPAIRMENT (CHILD PUGH B OR C)
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS
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.

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS NOT RECOMMENDED OR CONTRAINDICATED BY THE MANUFACTURER: CARBAMAZEPINE, RIFAMPIN, ETHINYL ESTRADIOL-CONTAINING MEDICATION, ATAZANAVIR,DARUNAVIR, LOPINAVIR, RITONAVIR, EFAVIRENZ, ATORVASTATIN, LOVASTATIN, SIMVASTATIN, ROSUVASTATIN AT DOSES GREATER THAN 10MG, OR CYCLOSPORINE AT DOSES GREATER THAN 100MG PER DAY. PATIENT MUST NOT HAVE PRIOR FAILURE OF A DAA REGIMEN WITH NS5A INHIBITOR AND HCV PROTEASE INHIBITOR.

GLYCEROL PHENYLBUTYRATE

Products Affected

- RAVICTI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF OR CONTRAINDICATION TO SODIUM PHENYLBUTYRATE (BUPHENYL).

GOLIMUMAB IV

Products Affected

- SIMPONI ARIA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS: PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA. PSORIATIC ARTHRITIS: PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS: PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA OR COSENTYX.

GOLIMUMAB SQ

Products Affected

- SIMPONI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. ULCERATIVE COLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: RA: 6 MONTHS. PSA/AS: 4 MONTHS. UC: 12 MONTHS. RENEWAL: 12 MONTHS FOR ALL DIAGNOSES.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, CIMZIA OR ACTEMRA. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA OR COSENTYX. ULCERATIVE COLITIS (UC): PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING CONVENTIONAL AGENTS SUCH AS A CORTICOSTEROID (I.E., BUDESONIDE, METHYLPREDINSOLONE), AZATHIOPRINE, MERCAPTOPURINE, METHOTREXATE, OR MESALAMINE.

GUSELKUMAB

Products Affected

- TREMFYA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL WITH HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: COSENTYX OR OTEZLA.

HYDROXYPROGESTERONE CAPROATE-DELALUTIN GENERIC

Products Affected

- *hydroxyprogesterone caproate*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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.

IBRUTINIB

Products Affected

- IMBRUVICA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IBUPROFEN-FAMOTIDINE

Products Affected

- DUEXIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF ONE OF THE FOLLOWING GENERIC, FEDERAL LEGEND HISTAMINE H2-RECEPTOR ANTAGONISTS: FAMOTIDINE, CIMETIDINE, NIZATIDINE, OR RANITIDINE, AND TRIAL OF GENERIC, FEDERAL LEGEND IBUPROFEN.

IDELALISIB

Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IMATINIB MESYLATE

Products Affected

- *imatinib oral tablet 100 mg, 400 mg*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ALL DIAGNOSES: 12 MONTHS. ADJUVANT GASTROINTESTINAL STROMAL TUMOR (GIST) TREATMENT: 36 MONTHS.
Other Criteria	PATIENTS WITH PREVIOUSLY-TREATED CML REQUIRE A BCR-ABL MUTATIONAL ANALYSIS CONFIRMING THAT THE PATIENT IS NEGATIVE FOR THE FOLLOWING MUTATIONS: T315I, V299L, F317L/V/I/C, Y253H, E255K/V, F359V/C/I.

IMIQUIMOD - ALDARA

Products Affected

- *imiquimod*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	ACTINIC KERATOSIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. SUPERFICIAL BASAL CELL CARCINOMA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR AN ONCOLOGIST.
Coverage Duration	4 MONTHS
Other Criteria	EXTERNAL GENITAL WARTS: TRIAL OF PODOFILOX (CONDYLOX) 0.5% TOPICAL SOLUTION. ACTINIC KERATOSIS BRAND DRUG REQUEST: TRIAL OF GENERIC IMIQUIMOD 5% CREAM. SUPERFICIAL BASAL CELL CARCINOMA: LESS THAN 2CM IN SIZE AND NOT ON THE FACE.

INFLIXIMAB

Products Affected

- REMICADE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5 % BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSORIASIS PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. CROHN'S DISEASE/ULCERATIVE COLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: CD/UC: 8 MO. RA: 6 MO. PSA/AS/PSO: 4 MO. RENEWAL FOR ALL INDICATIONS: 12 MO.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA. PSORIATRIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY CIMZIA OR COSENTYX. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: COSENTYX OR OTEZLA. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY CIMZIA. ULCERATIVE COLITIS (UC): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: SIMPONI OR ENTYVIO.

INFLIXIMAB-ABDA

Products Affected

- RENFLEXIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSORIASIS PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHN'S DISEASE/ULCERATIVE COLITIS: GASTROENTEROLOGIST.
Coverage Duration	INITIAL: CD/UC: 8 MOS. RA: 6 MOS. PSA/AS/PSO: 4 MOS. RENEWAL FOR ALL INDICATIONS: 12 MOS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA. PSORIATRIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY CIMZIA OR COSENTYX. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: COSENTYX OR OTEZLA. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY CIMZIA. ULCERATIVE COLITIS (UC): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: SIMPONI OR ENTYVIO.

INFLIXIMAB-DYYB

Products Affected

- INFLECTRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS: SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5 PERCENT BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS, ANKYLOSING SPONDYLITIS: PRESCRIBED BY GIVEN OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSORIASIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. CROHN'S DISEASE/ULCERATIVE COLITIS: GASTROENTEROLOGIST.
Coverage Duration	INITIAL: CD/UC: 8 MOS. RA: 6 MOS. PSA/AS/PSO: 4 MOS. RENEWAL FOR ALL INDICATIONS: 12 MOS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: ORENCIA, XELJANZ, CIMZIA, OR ACTEMRA. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: CIMZIA, OTEZLA, OR COSENTYX. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: CIMZIA OR COSENTYX. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: COSENTYX OR OTEZLA. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY CIMZIA. ULCERATIVE COLITIS (UC): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING: SIMPONI OR ENTYVIO.

INOTUZUMAB OZOGAMICIN

Products Affected

- BESPONSA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

INTERFERON ALFA-2B

Products Affected

- INTRON A INJECTION

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. FOR USE TO TREAT HEPATITIS C, CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HEPATITIS C: GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (E.G. HEPATOLOGIST). NO REQUIREMENT FOR OTHER FDA APPROVED INDICATIONS.
Coverage Duration	6 MONTHS
Other Criteria	LIMITED TO 1 YEAR OF THERAPY EXCEPT 18 MONTHS FOR FOLLICULAR LYMPHOMA. HEPATITIS C GENOTYPE 1, 2, 3, 4, 5, OR 6: REQUIRES A TRIAL OF OR CONTRAINDICATION TO PEGINTERFERON ALFA-2A OR PEGINTERFERON ALFA-2B USED IN COMBINATION WITH RIBAVIRIN UNLESS CONTRAINDICATED.

INTERFERONS FOR MS-AVONEX, PLEGRIDY, REBIF

Products Affected

- AVONEX (WITH ALBUMIN)
- AVONEX INTRAMUSCULAR PEN INJECTOR KIT
- AVONEX INTRAMUSCULAR SYRINGE KIT
- PLEGRIDY
- REBIF (WITH ALBUMIN)
- REBIF REBIDOSE
- REBIF TITRATION PACK

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

INTERFERONS FOR MS-BETASERON, EXTAVIA

Products Affected

- BETASERON SUBCUTANEOUS KIT
- EXTAVIA SUBCUTANEOUS KIT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	TRIAL WITH TWO OF THE FOLLOWING AGENTS FOR MULTIPLE SCLEROSIS: AUBAGIO, AVONEX, GILENYA, PLEGRIDY, REBIF, TECFIDERA, AND GLATIRAMER
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IPILIMUMAB

Products Affected

- YERVOY

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: UNRESECTABLE/MET MELANOMA: 4 MO, RCC: 3 MO. CUTANEOUS MELANOMA: INITIAL AND RENEWAL: 6 MO
Other Criteria	RENEWAL FOR ADJUVANT CUTANEOUS MELANOMA: NO EVIDENCE OF DISEASE RECURRENCE (DEFINED AS THE APPEARANCE OF ONE OR MORE NEW MELANOMA LESIONS: LOCAL, REGIONAL OR DISTANT METASTASIS).

IVABRADINE

Products Affected

- CORLANOR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PATIENT MUST HAVE NEW YORK HEART ASSOCIATION (NYHA) CLASS II TO IV HEART FAILURE
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A CARDIOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: APPROVAL REQUIRES THE PATIENT DOES NOT HAVE A DEMAND PACEMAKER SET TO A RATE OF 60 BEATS PER MINUTE OR GREATER. PATIENT IS CURRENTLY RECEIVING TREATMENT WITH OR HAS AN INTOLERANCE TO A FORMULARY BETA BLOCKER SUCH AS METOPROLOL SUCCINATE, BISOPROLOL, OR CARVEDILOL. RENEWAL: APPROVAL REQUIRES DIAGNOSIS OF HEART FAILURE AND PATIENT MUST BE IN SINUS RHYTHM.

IVACAFTOR

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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	6 YEARS OF AGE OR OLDER.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IVACAFTOR - GRANULE PACKETS

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	F508DEL MUTATION IN CFTR GENE.
Required Medical Information	CONFIRMED MUTATION IN CFTR GENE ACCEPTABLE FOR THE TREATMENT OF CYSTIC FIBROSIS.
Age Restrictions	2 YEARS OF AGE TO 5 YEARS OF AGE
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IXAZOMIB

Products Affected

- NINLARO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

IXEKIZUMAB

Products Affected

- TALTZ AUTOINJECTOR
- TALTZ SYRINGE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PLAQUE PSORIASIS (PSO): MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5 PERCENT BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL (FOR ALL INDICATIONS): PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PLAQUE PSORIASIS (PSO): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST.
Coverage Duration	PLAQUE PSORIASIS (PSO) AND PSORIATIC ARTHRITIS (PSA): INITIAL 4 MONTHS, RENEWAL 12 MONTHS.
Other Criteria	INITIAL: PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: COSENTYX OR OTEZLA. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX.

LEDIPASVIR-SOFOSBUVIR

Products Affected

- HARVONI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
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, SIMEPREVIR, SOFOSBUVIR (AS A SINGLE AGENT), STRIBILD (ELVITEGRAVIR/COBICISTAT/EMTRICITABINE /TENOFOVIR), OR TIPRANAVIR/RITONAVIR.

LENALIDOMIDE

Products Affected

- REVLIMID

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

LENVATINIB MESYLATE

Products Affected

- LENVIMA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

LETERMOVIR

Products Affected

- PREVYMIS INTRAVENOUS SOLUTION 240 MG/12 ML, 480 MG/24 ML
- PREVYMIS ORAL

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	4 MONTHS
Other Criteria	

L-GLUTAMINE

Products Affected

- ENDARI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	INITIAL CRITERIA FOR ADULTS (18 YEARS OR OLDER): PHYSICIAN ATTESTATION OF 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 (ACS). INITIAL REQUESTS FOR PATIENTS BETWEEN THE AGES OF 5-17 WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA. RENEWAL FOR ALL PATIENTS: PHYSICIAN ATTESTATION THAT PATIENT HAS MAINTAINED OR EXPERIENCED REDUCTION IN ACUTE COMPLICATIONS OF SICKLE CELL DISEASE.

LIDOCAINE

Products Affected

- *lidocaine topical adhesive patch,medicated*
- *lidocaine topical ointment*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. ADDITIONAL COVERAGE FOR DIABETIC NEUROPATHY WILL BE CONSIDERED FOR REQUESTS FOR LIDOCAINE TOPICAL PATCHES.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	PATCH: 12 MONTHS. OINTMENT: 3 MONTHS.
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

LIFITEGRAST OPHTHALMIC

Products Affected

- XIIDRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN OPHTHALMOLOGIST, OPTOMETRIST OR RHEUMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	PREVIOUS TRIAL OF OR CONTRAINDICATION TO CYCLOSPORINE OPHTHALMIC (RESTASIS).

LOMITAPIDE

Products Affected

- JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 40 MG, 5 MG, 60 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	LDL CHOLESTEROL LEVEL, LDL RECEPTOR STATUS
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST, ENDOCRINOLOGIST OR LIPIDOLOGIST.
Coverage Duration	INITIAL: 7 MONTHS RENEWAL: 6 MONTHS

PA Criteria	Criteria Details
Other Criteria	DIAGNOSIS CONFIRMED BY ONE OF THE FOLLOWING: SIMON BROOME DIAGNOSTIC CRITERIA (DEFINITE), (E.G. GENETIC TESTING CONSISTENT WITH HOFH AND PRETREATMENT BASELINE LDL CHOLESTEROL IS GREATER THAN 190 MG/DL), CASCADE SCREENING, DUTCH LIPID NETWORK CRITERIA WITH A SCORE OF AT LEAST 6, OR HISTORY OF UNTREATED CHOLESTEROL GREATER THAN 500MG/DL (OR TREATED CHOLESTEROL GREATER THAN 300MG/DL) AND CUTANEOUS XANTHOMA BEFORE 10 YEARS OF AGE. LOMITAPIDE WILL NOT BE APPROVED FOR PATIENTS CONCURRENTLY USING ANY OF THE FOLLOWING STRONG OR MODERATE CYP3A4 MEDICATIONS: CLARITHROMYCIN, CONIVAPTAN, INDINAVIR, ITRACONAZOLE, KETOCONAZOLE, LOPINAVIR/RITONAVIR, MIBEFRADIL, NEFAZODONE, NELFINAVIR, POSACONAZOLE, RITONAVIR, SAQUINAVIR, TELITHROMYCIN, TIPRANAVIR/RITONAVIR, VORICONAZOLE, AMPRENAVIR, APREPITANT, ATAZANAVIR, CIPROFLOXACIN, CRIZOTINIB, DARUNAVIR/RITONAVIR, DILTIAZEM, ERYTHROMYCIN, FLUCONAZOLE, FOSAMPRENAVIR, IMATINIB, OR VERAPAMIL. INITIAL: LDL CHOLESTEROL LEVEL OF AT LEAST 160MG/DL WHILE ON LIPID-LOWERING THERAPY PRIOR TO INITIATING LOMITAPIDE. PREVIOUS TRIAL OF A PCSK9 INHIBITOR (E.G. ALIROCUMAB OR EVOLOCUMAB), UNLESS THE PATIENT HAS NON-FUNCTIONING LDL RECEPTORS. PREVIOUS TRIAL OF ROSUVASTATIN OR ATORVASTATIN, UNLESS THE PATIENT HAS AN ABSOLUTE CONTRAINDICATION TO STATIN THERAPY (E.G. ACTIVE, DECOMPENSATED LIVER DISEASE, NURSING FEMALE, PREGNANCY OR PLANS TO BECOME PREGNANT, HYPERSENSITIVITY REACTION). STATIN-TOLERANT PATIENTS MUST BE TAKING ATORVASTATIN OR ROSUVASTATIN FOR THE PAST 2 MONTHS PRIOR TO STARTING LOMITAPIDE. LOMITAPIDE MUST BE USED IN COMBINATION WITH ATORVASTATIN OR ROSUVASTATIN. IF THE PATIENT HAS PREVIOUSLY TRIED ATORVASTATIN OR ROSUVASTATIN, LOMITAPIDE MUST BE USED IN COMBINATION WITH ANOTHER STATIN OR FORMULARY LDL-LOWERING AGENT (E.G. BILE ACID SEQUESTRANT, GEMFIBROZIL OR OTHER FIBRATE, EZETIMIBE, OR NIACIN). STATIN-INTOLERANT PATIENTS REQUIRE EITHER PHYSICIAN ATTESTATION OF

PA Criteria	Criteria Details
	STATIN INTOLERANCE OR HISTORY OF SKELETAL-MUSCLE RELATED SYMPTOMS (E.G., MYOPATHY) DUE TO A PREVIOUS TRIAL OF STATINS (E.G. ROSUVASTATIN OR ATORVASTATIN). FOR STATIN-INTOLERANT PATIENTS, LOMITAPIDE MUST BE USED IN COMBINATION WITH ONE OF THE FOLLOWING FORMULARY LIPID-LOWERING TREATMENTS: EZETIMIBE, FENOFIBRATE, NIACIN, OR A BILE ACID SEQUESTRANT (E.G. CHOLESTYRAMINE, COLESTIPOL, COLESEVELAM). RENEWAL: PATIENT HAS RECEIVED AT LEAST 6 MONTHS OF THERAPY WITH LOMITAPIDE IN COMBINATION WITH ANOTHER AND LIPID-LOWERING AGENT.

LUMACAFTOR-IVACAFTOR

Products Affected

- ORKAMBI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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 12 MONTHS.
Other Criteria	RENEWAL: MAINTAINED OR IMPROVEMENT IN FEV1 OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS OR IMPROVEMENT IN BODY MASS INDEX (BMI).

MEPOLIZUMAB

Products Affected

- NUCALA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	SEVERE ASTHMA: CONCURRENT USE OF XOLAIR.
Required Medical Information	SEVERE ASTHMA: BLOOD EOSINOPHIL LEVEL GREATER THAN OR EQUAL TO 150 CELLS/MCL WITHIN THE LAST 6 WEEKS OR GREATER THAN OR EQUAL TO 300 CELLS/MCL WITHIN THE LAST 12 MONTHS.
Age Restrictions	
Prescriber Restrictions	SEVERE ASTHMA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN PULMONARY MEDICINE, AN ALLERGIST OR AN IMMUNOLOGIST.
Coverage Duration	INITIAL: SEVERE ASTHMA: 24 WEEKS. EGPA: 12 MONTHS. RENEWAL FOR ALL INDICATIONS: 12 MONTHS.
Other Criteria	INITIAL THERAPY: SEVERE ASTHMA: PATIENT CURRENTLY TREATED WITH A MAXIMALLY TOLERATED DOSE OF INHALED CORTICOSTEROIDS AND AT LEAST ONE OTHER MAINTENANCE MEDICATION WHICH INCLUDES ANY OF THE FOLLOWING: LONG-ACTING INHALED BETA2-AGONIST, LONG-ACTING MUSCARINIC ANTAGONIST, A LEUKOTRIENE RECEPTOR ANTAGONIST, THEOPHYLLINE, OR ORAL CORTICOSTEROID. RENEWAL: SEVERE ASTHMA: REQUIRES DOCUMENTATION THAT THE PATIENT HAS EXPERIENCED IMPROVEMENT IN ASTHMA EXACERBATIONS FROM BASELINE (PHYSICIAN ATTESTATION) AND A REDUCTION IN ORAL CORTICOSTEROID DOSE (IF THE PATIENT WAS ON A MAINTENANCE REGIMEN OF ORAL CORTICOSTEROIDS AT THE INITIATION OF TREATMENT).

METHYLNALTREXONE

Products Affected

- RELISTOR SUBCUTANEOUS SOLUTION
- RELISTOR SUBCUTANEOUS SYRINGE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	ADVANCED ILLNESS: OPIOID-INDUCED CONSTIPATION.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS FOR PATIENTS RECEIVING PALLIATIVE CARE,12 MONTHS FOR PATIENTS WITH CHRONIC,NON-CANCER PAIN.
Other Criteria	ADVANCED ILLNESS: PATIENT IS RECEIVING PALLIATIVE CARE. CHRONIC NON-CANCER PAIN: PATIENT HAS BEEN TAKING OPIOIDS FOR AT LEAST 4 WEEKS AND HAD A PREVIOUS TRIAL OF OR CONTRAINDICATION TO NALOXEGOL (MOVANTIK) OR LUBIPROSTONE (AMITIZA).

METHYLNALTREXONE ORAL

Products Affected

- RELISTOR ORAL

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PATIENT HAS BEEN TAKING OPIOIDS FOR AT LEAST 4 WEEKS AND HAD A PREVIOUS TRIAL OF OR CONTRAINDICATION TO NALOXEGOL (MOVANTIK) OR LUBIPROSTONE (AMITIZA).

MIDOSTAURIN

Products Affected

- RYDAPT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ACUTE MYELOID LEUKEMIA: 6 MONTHS. ADVANCED SYSTEMIC MASTOCYTOSIS: 12 MONTHS
Other Criteria	

MIFEPRISTONE

Products Affected

- KORLYM

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

MILTEFOSINE

Products Affected

- IMPAVIDO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

MIPOMERSEN

Products Affected

- KYNAMRO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	LDL CHOLESTEROL LEVEL, LDL RECEPTOR STATUS
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A CARDIOLOGIST, ENDOCRINOLOGIST OR LIPIDOLOGIST
Coverage Duration	INITIAL: 7 MONTHS RENEWAL 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	DIAGNOSIS OF HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA AS DETERMINED BY ONE OF THE FOLLOWING CRITERIA: SIMON BROOME DIAGNOSTIC CRITERIA (DEFINITE) [EXAMPLE: GENETIC TESTING CONSISTENT WITH HOFH AND PRETREATMENT BASELINE LDL CHOLESTEROL IS GREATER THAN 190 MG/DL], CASCADE SCREENING, DUTCH LIPID NETWORK CRITERIA WITH A SCORE AT LEAST 6, OR HISTORY OF UNTREATED CHOLESTEROL GREATER THAN 500MG/DL (OR TREATED GREATER THAN 300MG/DL) AND CUTANEOUS XANTHOMA BEFORE AGE 10. INITIAL CRITERIA: CURRENT LDL CHOLESTEROL LEVEL IS AT LEAST 160MG/DL. PATIENT DOES NOT HAVE ANY OF THE FOLLOWING CONTRAINDICATIONS TO KYNAMRO (MIPOMERSEN): MODERATE OR SEVERE HEPATIC IMPAIRMENT OR ACTIVE LIVER DISEASE, INCLUDING UNEXPLAINED PERSISTENT ELEVATIONS OF SERUM TRANSAMINASES. PREVIOUS TRIAL OF A PCSK9 INHIBITOR (SUCH AS ALIROCUMAB OR EVOLOCUMAB) UNLESS THE PATIENT HAS NON-FUNCTIONING LDL RECEPTORS. PREVIOUS TRIAL WITH ONE OF THE FOLLOWING STATINS: ROSUVASTATIN OR ATORVASTATIN. PATIENTS WITH ABSOLUTE CONTRAINDICATION TO STATIN THERAPY (ACTIVE, DECOMPENSATED LIVER DISEASE, NURSING FEMALE, PREGNANCY OR PLANS TO BECOME PREGNANT, HYPERSENSITIVITY REACTION) WILL BE APPROVED FOR THERAPY WITHOUT REQUIREMENT OF A TRIAL WITH A STATIN. STATIN-TOLERANT PATIENTS: PRIOR TO (KYNAMRO), PATIENT MUST HAVE BEEN TAKING ONE OF THE FOLLOWING: ATORVASTATIN OR ROSUVASTATIN, FOR AT LEAST 2 MONTHS WITHIN THE PAST 2 MONTHS. FOR STATIN-INTOLERANT PATIENTS: DOCUMENTATION OF STATIN INTOLERANCE WHICH INCLUDES THE FOLLOWING: PHYSICIAN ATTESTATION OR PATIENT HAS TRIED ROSUVASTATIN OR ATORVASTATIN AND HAS EXPERIENCED SKELETAL-MUSCLE RELATED SYMPTOMS (E.G., MYOPATHY). UNLESS CONTRAINDICATED, PATIENT MUST BE ON CONCURRENT THERAPY WITH ONE OF THE FOLLOWING LIPID-LOWERING TREATMENTS (SUCH AS A STATIN [SIMVASTATIN, ATORVASTATIN], EZETIMIBE, FENOFIBRATE, NIACIN, OR BILE ACID SEQUESTRANT [CHOLESTYRAMINE, COLESTIPOL, COLESEVELAM]). RENEWAL CRITERIA:

PA Criteria	Criteria Details
	PATIENT HAS RECEIVED THERAPY FOR AT LEAST 6 MONTHS AND MUST ALSO BE TAKING KYNAMRO IN COMBINATION WITH ANOTHER LIPID-LOWERING AGENT.

NAPROXEN- ESOMEPRAZOLE

Products Affected

- VIMOVO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF ONE OF THE FOLLOWING GENERIC, FEDERAL LEGEND PROTON PUMP INHIBITORS: OMEPRAZOLE, LANSOPRAZOLE, OR PANTOPRAZOLE AND A TRIAL OF GENERIC, FEDERAL LEGEND NAPROXEN.

NATALIZUMAB

Products Affected

- TYSABRI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CROHN'S DISEASE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	MULTIPLE SCLEROSIS: 12 MONTHS. CROHN'S DISEASE: INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	MULTIPLE SCLEROSIS INITIAL CRITERIA: PREVIOUS TRIAL OF TWO OF THE FOLLOWING PREFERRED AGENTS FOR MULTIPLE SCLEROSIS: GLATIRAMER, REBIF, AVONEX, PLEGRIDY, TECFIDERA, GILENYA, OR AUBAGIO. CROHN'S DISEASE INITIAL CRITERIA: PREVIOUS TRIAL OF HUMIRA AND CIMZIA. CROHN'S DISEASE RENEWAL CRITERIA: PATIENT HAS RECEIVED AT LEAST 12 MONTHS OF THERAPY WITH TYSABRI WITH PHYSICIAN ATTESTATION THAT THE PATIENT HAS NOT REQUIRED MORE THAN 3 MONTHS OF CORTICOSTEROID USE WITHIN THE PAST 12 MONTHS TO CONTROL THEIR CROHN'S DISEASE WHILE ON TYSABRI, OR PATIENT HAS ONLY RECEIVED 6 MONTHS OF THERAPY WITH TYSABRI WITH PHYSICIAN ATTESTATION THAT THE PATIENT HAS TAPERED OFF CORTICOSTEROIDS DURING THE FIRST 24 WEEKS OF TYSABRI THERAPY.

NECITUMUMAB

Products Affected

- PORTRAZZA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

NERATINIB MALEATE

Products Affected

- NERLYNX

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	EARLY-STAGE TUMOR (STAGE I-III) AND TUMOR IS HORMONE-RECEPTOR POSITIVE AND THE MEDICATION IS BEING REQUESTED WITHIN 2 YEARS OF COMPLETING THE LAST TRASTUZUMAB DOSE

NILOTINIB

Products Affected

- TASIGNA ORAL CAPSULE 150 MG, 200 MG, 50 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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, AND F359V/C/I.

NINTEDANIB

Products Affected

- OFEV

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	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, AND CANCER. NOT APPROVED IF PATIENT DOES NOT HAVE A PREDICTED FORCED VITAL CAPACITY (FVC) OF AT LEAST 50 PERCENT OR HAS NOT OBTAINED LIVER FUNCTION TESTS
Required Medical Information	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.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PULMONOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	

NIRAPARIB TOSYLATE

Products Affected

- ZEJULA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

NITISINONE

Products Affected

- ORFADIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	DIAGNOSIS OF HEREDITARY TYROSINEMIA TYPE 1 AS CONFIRMED BY ELEVATED URINARY OR PLASMA SUCCINYLACETONE LEVELS OR A MUTATION IN THE FUMARYLACETOACETATE HYDROLASE GENE.
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	ORFADIN SUSPENSION: TRIAL OF ORFADIN CAPSULES. RENEWAL: THE PATIENT'S URINARY OR PLASMA SUCCINYLACETONE LEVELS HAVE DECREASED FROM BASELINE WHILE ON TREATMENT WITH NITISINONE.

NIVOLUMAB

Products Affected

- OPDIVO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MELANOMA: OPDIVO IS NOT APPROVED FOR COMBINATION THERAPY WITH TAFINLAR, MEKINIST (TRAMETINIB), COTELLIC (COBIMETINIB), OR ZELBORAF.

NUEDEXTA

Products Affected

- NUEDEXTA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

OBETICHOLIC ACID

Products Affected

- OCALIVA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	PATIENTS WITH COMPLETE BILIARY OBSTRUCTION.
Required Medical Information	DIAGNOSIS OF PRIMARY BILIARY CHOLANGITIS AS CONFIRMED BY AT LEAST TWO OF THE FOLLOWING CRITERIA: AN ALKALINE PHOSPHATASE LEVEL OF AT LEAST 1.5 TIMES THE UPPER LIMIT OF NORMAL (ULN), THE PRESENCE OF ANTIMITOCHONDRIAL ANTIBODIES AT A TITER OF 1:40 OR HIGHER, HISTOLOGIC EVIDENCE OF NON-SUPPURATIVE DESTRUCTIVE CHOLANGITIS AND DESTRUCTION OF INTERLOBULAR BILE DUCTS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST OR HEPATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	INITIAL: USED IN COMBINATION WITH URSODEOXYCHOLIC ACID (E.G., URSODIOL, URSO 250, URSO FORTE) IN ADULTS WITH AN INADEQUATE RESPONSE TO URSODEOXYCHOLIC ACID AT A DOSAGE OF 13-15 MG/KG/DAY FOR AT LEAST 1 YEAR, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE URSODEOXYCHOLIC ACID. RENEWAL: PATIENT'S ALKALINE PHOSPHATASE LEVELS ARE LESS THAN 1.67-TIMES THE UPPER LIMIT OF NORMAL OR HAVE DECREASED BY AT LEAST 15% FROM BASELINE WHILE ON TREATMENT WITH OBETICHOLIC ACID.

OBINUTUZUMAB

Products Affected

- GAZYVA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	

OCRELIZUMAB

Products Affected

- OCREVUS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RELAPSING FORM OF MULTIPLE SCLEROSIS: TRIAL OF TWO OF THE FOLLOWING AGENTS FOR MULTIPLE SCLEROSIS: AUBAGIO, AVONEX, GILENYA, PLEGRIDY, REBIF, TECFIDERA, OR GLATIRAMER. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.

OLAPARIB

Products Affected

- LYNPARZA ORAL CAPSULE
- LYNPARZA ORAL TABLET

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

OLARATUMAB

Products Affected

- LARTRUVO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THIS DRUG MAY BE COVERED UNDER MEDICARE PART B OR D DEPENDING UPON THE CIRCUMSTANCES. INFORMATION MAY NEED TO BE SUBMITTED DESCRIBING THE USE AND SETTING OF THE DRUG TO MAKE THE DETERMINATION.

OMACETAXINE

Products Affected

- SYNRIBO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INDUCTION: 3 MONTHS. POST INDUCTION OR 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 AN ABSOLUTE NEUTROPHIL COUNT (ANC) GREATER THAN OR EQUAL TO 1.5 X 10^9/L, PLATELETS GREATER THAN OR EQUAL TO 100 X 10^9/L WITHOUT BLOOD BLASTS OR 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.

OMALIZUMAB

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL CRITERIA FOR ASTHMA: PATIENT MEETS THE CRITERIA OF MODERATE TO SEVERE ASTHMA, POSITIVE SKIN PRICK OR RAST TEST, FEV1 LESS THAN 80%, DEMONSTRATED INADEQUATELY CONTROLLED SYMPTOMS ON INHALED CORTICOSTEROIDS AND SECOND ASTHMA CONTROLLER, BASELINE IGE SERUM LEVEL GREATER THAN OR EQUAL TO 30IU/ML. RENEWAL CRITERIA FOR ASTHMA: PHYSICIAN ATTESTATION OF IMPROVEMENT IN ASTHMA EXACERBATIONS FROM BASELINE OR A REDUCTION IN ORAL OR INHALED CORTICOSTEROID USE.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A SPECIALIST IN ALLERGY, PULMONARY MEDICINE, DERMATOLOGY OR IMMUNOLOGY.
Coverage Duration	INITIAL: ASTHMA: 12 MOS. CHRONIC IDIOPATHIC URTICARIA: 6 MOS. RENEWAL FOR ALL INDICATIONS: 12 MOS.
Other Criteria	FOR CHRONIC IDIOPATHIC URTICARIA: PREVIOUS TRIAL OF OR CONTRAINDICATION TO A MAXIMALLY TOLERATED DOSE OF AN H1 ANTI-HISTAMINE (SUCH AS CLARINEX OR XYZAL) AND PATIENT STILL EXPERIENCES HIVES ON MOST DAYS OF THE WEEK.

OMBITASVIR-PARITAPREVIR-RITONAVIR

Products Affected

- TECHNIVIE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
Exclusion Criteria	DECOMPENSATED CIRRHOSIS, MODERATE OR SEVERE LIVER IMPAIRMENT (CHILD-PUGH B OR C).
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.

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. MUST BE USED CONCURRENTLY WITH RIBAVIRIN. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING (CONTRAINDICATED OR NOT RECOMMENDED BY THE MANUFACTURER): ALFUZOSIN, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, RIFAMPIN, ERGOTAMINE, DIHYDROERGOTAMINE, ERGONOVINE, METHYLERGONOVINE, ETHINYL ESTRADIOL CONTAINING MEDICATIONS (SUCH AS COMBINED ORAL CONTRACEPTIVES, NUVARING, ORTHO EVRA OR XULANE TRANSDERMAL PATCH SYSTEM), LOVASTATIN, SIMVASTATIN, PIMOZIDE, EFAVIRENZ (ATRIPLA, SUSTIVA), REVATIO (SILDENAFIL DOSE OF 20MG AND/OR DOSED THREE TIMES DAILY FOR PAH), TRIAZOLAM, ORAL MIDAZOLAM, LOPINAVIR/RITONAVIR, RILPIVIRINE, SALMETEROL.

OMBITASVIR-PARITAPREVIR-RITONAVIR-DASABUVIR

Products Affected

- VIEKIRA PAK
- VIEKIRA XR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
Exclusion Criteria	DECOMPENSATED CIRRHOSIS, MODERATE OR SEVERE LIVER IMPAIRMENT (CHILD-PUGH B OR C).
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.

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING: ALFUZOSIN, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, GEMFIBROZIL, RIFAMPIN, ERGOTAMINE, DIHYDROERGOTAMINE, ERGONOVINE, METHYLERGONOVINE, ETHINYL ESTRADIOL CONTAINING MEDICATIONS (SUCH AS COMBINED ORAL CONTRACEPTIVES, NUVARING, ORTHO EVRA OR XULANE TRANSDERMAL PATCH SYSTEM), ST. JOHN'S WORT, LOVASTATIN, SIMVASTATIN, PIMOZIDE, EFAVIRENZ, REVATIO, TRIAZOLAM, ORAL MIDAZOLAM, DARUNAVIR/RITONAVIR, LOPINAVIR/RITONAVIR, RILPIVIRINE, SALMETEROL.

OSIMERTINIB

Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	METASTATIC NSCLC WITH EGFR T790M MUTATION: CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE-INHIBITOR.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

OXYMETHOLONE

Products Affected

- ANADROL-50

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	CARCINOMA OF THE PROSTATE OR BREAST IN MALE PATIENTS, CARCINOMA OF THE BREAST IN FEMALES WITH HYPERCALCEMIA, WOMEN WHO ARE OR MAY BECOME PREGNANT, NEPHROSIS OR THE NEPHROTIC PHASE OF NEPHRITIS, HYPERSENSITIVITY TO THE DRUG AND SEVERE HEPATIC DYSFUNCTION.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PALBOCICLIB

Products Affected

- IBRANCE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PALIVIZUMAB

Products Affected

- SYNAGIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	GESTATIONAL AGE
Age Restrictions	LESS THAN 24 MONTHS OF AGE.
Prescriber Restrictions	
Coverage Duration	1 MONTH TO 5 MONTHS. SEE OTHER CRITERIA FOR MORE INFORMATION.
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT RECOMMENDATIONS FROM THE AMERICAN ACADEMY OF PEDIATRICS FOR PALIVIZUMAB PROPHYLAXIS FOR RESPIRATORY SYNCYTIAL VIRUS INFECTIONS. INITIAL: APPROVAL WILL BE FOR AT LEAST 1 MONTH AND NO GREATER THAN 5 MONTHS DEPENDENT UPON REMAINING LENGTH OF RESPIRATORY SYNCYTIAL VIRUS (RSV) SEASON. RENEWAL: ADDITIONAL 1 MONTH OF TREATMENT FOR CARDIOPULMONARY BYPASS SURGERY DURING RSV PROPHYLAXIS SEASON.

PANITUMUMAB

Products Affected

- VECTIBIX

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PANOBINOSTAT

Products Affected

- FARYDAK

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RENEWAL: PATIENT HAS TOLERATED THE FIRST 8 CYCLES OF THERAPY WITHOUT UNRESOLVED SEVERE OR MEDICALLY SIGNIFICANT TOXICITY.

PARATHYROID HORMONE

Products Affected

- NATPARA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PAZOPANIB

Products Affected

- VOTRIENT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PDE5 INHIBITORS FOR PULMONARY ARTERIAL HYPERTENSION

Products Affected

- ADCIRCA
- *sildenafil (antihypertensive) oral*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	PATIENT CANNOT CONCURRENTLY OR INTERMITTENTLY BE TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA), ANY ORGANIC NITRATES IN ANY FORM, OR GUANYLATE CYCLASE (GC) STIMULATORS (ADEMPAS).
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) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. REQUEST FOR ADCIRCA REQUIRE TRIAL OR CONTRAINDICATION TO REVATIO. RENEWAL: PATIENT SHOWS 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. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

PDE5 INHIBITORS FOR PULMONARY ARTERIAL HYPERTENSION - IV

Products Affected

- *sildenafil (antihypertensive) intravenous*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	PATIENT CANNOT CONCURRENTLY OR INTERMITTENTLY BE TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA), ANY ORGANIC NITRATES IN ANY FORM, OR GUANYLATE CYCLASE (GC) STIMULATORS (ADEMPAS).
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) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. RENEWAL: PATIENT SHOWS 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.

PEG-INTERFERON ALFA-2B-SYLATRON

Products Affected

- SYLATRON

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	OVERALL DURATION OF THERAPY LIMITED TO 5 YEARS.

PEGVALIASE-PQPZ

Products Affected

- PALYNZIQ

PA Criteria	Criteria Details
Covered Uses	ALL FDA-APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	

PEMBROLIZUMAB

Products Affected

- KEYTRUDA INTRAVENOUS RECON SOLN
- KEYTRUDA INTRAVENOUS SOLUTION

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PENICILLAMINE

Products Affected

- CUPRIMINE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	RHEUMATOID ARTHRITIS: HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	WILSON'S DISEASE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEPATOLOGIST. CYSTINURIA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEPHROLOGIST. RHEUMATOID ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	WILSON'S DISEASE: GENETIC TESTING FOR ATP7B MUTATIONS. CYSTINURIA: DIAGNOSIS REQUIRES THE PRESENCE OF NEPHROLITHIASIS AND 1 OR MORE OF THE FOLLOWING: STONE ANALYSIS SHOWING PRESENCE OF CYSTEINE, IDENTIFICATION OF PATHOGNOMONIC HEXAGONAL CYSTINE CRYSTALS ON URINALYSIS, POSITIVE FAMILY HISTORY OF CYSTINURIA WITH POSITIVE CYANIDE-NITROPRUSSIDE SCREEN. REQUESTS FOR CUPRIMINE FOR THE TREATMENT OF WILSONS DISEASE, CYSTINURIA, AND RHEUMATOID ARTHRITIS REQUIRE A PREVIOUS TRIAL OF OR CONTRAINDICATION TO DEPEN.

PENICILLAMINE-DEPEN

Products Affected

- DEPEN TITRATABS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	RHEUMATOID ARTHRITIS: HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	WILSON'S DISEASE: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEPATOLOGIST. CYSTINURIA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEPHROLOGIST. RHEUMATOID ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	WILSON'S DISEASE: GENETIC TESTING FOR ATP7B MUTATIONS. CYSTINURIA: DIAGNOSIS REQUIRES THE PRESENCE OF NEPHROLITHIASIS AND 1 OR MORE OF THE FOLLOWING: STONE ANALYSIS SHOWING PRESENCE OF CYSTEINE, IDENTIFICATION OF PATHOGNOMONIC HEXAGONAL CYSTINE CRYSTALS ON URINALYSIS, POSITIVE FAMILY HISTORY OF CYSTINURIA WITH POSITIVE CYANIDE-NITROPRUSSIDE SCREEN.

PERTUZUMAB

Products Affected

- PERJETA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 5 MONTHS RENEWAL: 12 MONTHS
Other Criteria	

PIMAVANSERIN

Products Affected

- NUPLAZID

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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.

PIRFENIDONE

Products Affected

- ESBRIET ORAL CAPSULE MG
- ESBRIET ORAL TABLET 267 MG, 801

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	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, AND CANCER). NOT APPROVED IF THE PATIENT HAS NOT OBTAINED LIVER FUNCTION TESTS.
Required Medical Information	PATIENT WITH USUAL INTERSTITIAL PNEUMONIA (UIP) PATTERN AS EVIDENCED BY HIGH-RESOLUTION COMPUTED TOMOGRAPHY (HRCT) ALONE OR VIA A COMBINATION OF SURGICAL LUNG BIOPSY AND HRCT
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PULMONOLOGIST
Coverage Duration	12 MONTHS
Other Criteria	PATIENT HAS A PREDICTED FORCED VITAL CAPACITY (FVC) OF AT LEAST 50%.

POMALIDOMIDE

Products Affected

- POMALYST

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PONATINIB

Products Affected

- ICLUSIG ORAL TABLET 15 MG, 45 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

PRAMLINTIDE

Products Affected

- SYMLINPEN 120
- SYMLINPEN 60

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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	

PYRIMETHAMINE

Products Affected

- DARAPRIM

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D, ADDITIONAL CONSIDERATION FOR CHRONIC MAINTENANCE THERAPY FOR TOXOPLASMOSIS AND TOXOPLASMOSIS PROPHYLAXIS.
Exclusion Criteria	
Required Medical Information	MALARIA: PLASMODIA SUSCEPTIBLE TESTING. TOXOPLASMOSIS:CD4 LEVEL
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ACUTE MALARIA AND CHEMOPROPHYLAXIS: INITIAL: 3 MONTHS. RENEWAL: 12 MONTHS. SEE OTHER CRITERIA FIELD

PA Criteria	Criteria Details
Other Criteria	INITIAL: ACUTE MALARIA TREATMENT AND MALARIA CHEMOPROPHYLAXIS REQUIRES THAT THE PATIENT HAS MALARIA SUSCEPTIBLE TO PYRIMETHAMINE AND A PREVIOUS TRIAL OF PLAQUENIL (HYDROXYCHLOROQUINE SULFATE) AND MALARONE (ATOVAQUONE/PROGUANIL) (UNLESS THESE REGIMENS ARE RESISTANT IN THE SPECIFIC REGION AS INDICATED BY REGIONAL PLASMODIA SUSCEPTIBILITY). PRIMARY PROPHYLAXIS OF TOXOPLASMOSIS IN PATIENTS WITH HIV REQUIRES PREVIOUS TRIAL OF OR CONTRAINDICATION TO BACTRIM (SMX/TMP). RENEWAL: CONTINUATION OF TREATMENT FOLLOWING ACUTE MALARIA REQUIRES PREVIOUS INFECTION WITH MALARIA SUSCEPTIBLE TO PYRIMETHAMINE WITH SUBSEQUENT CLINICAL CURE (ELIMINATION OF MALARIA SYMPTOMS DEFINED AS CHILLS, FEVER, SWEATS, GENERAL MALAISE) FOLLOWED BY SYMPTOMS OF RELAPSE. CONTINUATION OF MALARIA CHEMOPROPHYLAXIS REQUIRES THE PATIENT WILL BE TRAVELING TO OR RESIDING IN AN AREA WHERE PLASMODIA SUSCEPTIBLE TO PYRIMETHAMINE EXISTS (MALARIA MUST BE SENSITIVE TO PYRIMETHAMINE).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. CONTINUATION OF PRIMARY PROPHYLAXIS FOR TOXOPLASMOSIS WITH HIV REQUIRES CD4 COUNT LESS THAN 200 CELLS/MM3 AND CURRENT ANTI RETROVIRAL THERAPY. TOXOPLASMOSIS: INITIAL: 8 WEEKS. RENEWAL: 6 MONTHS. PRIMARY PROPHYLAXIS OF TOXOPLASMOSIS: INITIAL AND RENEWAL IS 12 MONTHS.

QUININE SULFATE

Products Affected

- *quinine sulfate*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

RAMUCIRUMAB

Products Affected

- CYRAMZA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

REGORAFENIB

Products Affected

- STIVARGA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	FOR COLORECTAL CANCER: TRIAL OF OR CONTRAINDICATION TO AN ANTI-VEGF THERAPY SUCH AS AVASTIN OR ZALTRAP AND A FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY SUCH AS FOLFOX, FOLFOXIRI, FOLFIRI,CAPEOX, INFUSIONAL 5-FU/LV OR CAPECITABINE. IF APPLICABLE, A TRIAL OF OR CONTRAINDICATION TO AN ANTI-EGFR THERAPY SUCH AS ERBITUX OR VECTIBIX IS ALSO REQUIRED FOR KRAS WILD TYPE COLORECTAL CANCER. FOR GIST, A TRIAL OF OR CONTRAINDICATION TO GLEEVEC AND SUTENT IS REQUIRED.

RESLIZUMAB

Products Affected

- CINQAIR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	CONCURRENT USE OF XOLAIR
Required Medical Information	BLOOD EOSINOPHIL LEVEL GREATER THAN OR EQUAL TO 400 CELLS/MCL WITHIN THE LAST 6 MONTHS
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN PULMONARY MEDICINE
Coverage Duration	INITIAL 24 WEEKS. RENEWAL 12 MONTHS
Other Criteria	INITIALTHERAPY: PATIENT CURRENTLY TREATED WITH A MAXIMALLY TOLERATED DOSE OF INHALED CORTICOSTEROIDS. RENEWAL REQUIRES DOCUMENTATION THAT THE PATIENT HAS EXPERIENCED AT LEAST A 25 PERCENT REDUCTION IN ASTHMA EXACERBATIONS (FOR EXAMPLE: HOSPITALIZATIONS, URGENT OR EMERGENT CARE VISITS, USE OF RESCUE MEDICATIONS, ETC.) FROM BASELINE.

RIBOCICLIB

Products Affected

- KISQALI
- 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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

RIFAXIMIN

Products Affected

- XIFAXAN ORAL TABLET 200 MG, 550 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	TRAVELERS' DIARRHEA/HEPATIC ENCEPHALOPATHY: 12 MOS. IBS-D: 12 WKS.
Other Criteria	FOR RIFAXIMIN 550 MG TABLETS ONLY: HEPATIC ENCEPHALOPATHY (HE): PREVIOUS TRIAL OF OR CONTRAINDICATION TO LACTULOSE OR CONCURRENT LACTULOSE THERAPY.

RIOCIGUAT

Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	INITIAL FOR PAH: PATIENT IS NOT CONCURRENTLY TAKING NITRATES OR NITRIC OXIDE DONORS (E.G. AMYL NITRATE), PHOSPHODIESTERASE INHIBITORS (E.G. SILDENAFIL, TADALAFIL, OR VARDENAFIL), OR NON-SPECIFIC PDE INHIBITORS (E.G. DIPYRIDAMOLE, THEOPHYLLINE). INITIAL FOR CTEPH: PATIENT IS NOT A CANDIDATE FOR SURGERY OR HAS INOPERABLE CTEPH. PERSISTENT OR RECURRENT DISEASE AFTER SURGICAL TREATMENT. PATIENT IS NOT CONCURRENTLY OR INTERMITTENTLY TAKING NITRATES, NITRIC OXIDE DONORS OR ANY PDE INHIBITORS (E.G.VIAGRA, CIALIS, DIPYRIDAMOLE).
Required Medical Information	DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS II-IV SYMPTOMS. DIAGNOSIS OF PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH) WHO GROUP 4. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS II-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	INITIAL FOR PAH: MEAN PULMONARY ARTERY PRESSURE (PAP) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. PREVIOUS TRIAL OF OR CONTRAINDICATION TO A PHOSPHODIESTERASE-5 (PDE-5) INHIBITOR, SUCH AS REVATIO OR ADCIRCA. RENEWAL FOR PAH AND CTEPH: 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.

RITUXIMAB

Products Affected

- RITUXAN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: RA: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. NHL, CLL: ONCOLOGIST.
Coverage Duration	RA: INITIAL: 6 MO. RENEWAL: 12 MONTHS. NHL, PV: 12 MONTHS. CLL: 6 MO. WG, MPA: 3 MONTHS.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS: PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: ORENCIA, XELJANZ, CIMZIA OR ACTEMRA.

RITUXIMAB SQ

Products Affected

- RITUXAN HYCELA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	THE PATIENT HAS RECEIVED OR WILL RECEIVE AT LEAST ONE FULL DOSE OF A RITUXIMAB PRODUCT BY INTRAVENOUS INFUSION PRIOR TO INITIATION OF RITUXIMAB AND HYALURONIDASE.

ROMIPLOSTIM

Products Affected

- NPLATE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	INITIAL: ADEQUATE RESPONSE TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR SPLENECTOMY. RENEWAL: NO CLINICAL RESPONSE AS DEFINED BY AN INCREASE IN PLATELET COUNT OF GREATER THAN OR EQUAL TO 50 X10^9/L AT THE MAX DOSE OF 10 MCG/KG PER DAY FOR 4 WEEKS.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITL: 2 MO., RENEW: IF NO RESPONSE AFTER INITIAL APPROVAL: 1 MO. AT MAX DOSE. IF RESPONSE: 12 MO.
Other Criteria	

RUCAPARIB

Products Affected

- RUBRACA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

RUXOLITINIB

Products Affected

- JAKAFI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: IMPROVEMENT OR MAINTENANCE OF SYMPTOM IMPROVEMENT SUCH AS A 50% OR GREATER REDUCTION IN TOTAL SYMPTOM SCORE ON THE MODIFIED MYELOFIBROSIS SYMPTOM ASSESSMENT FORM (MFSAF) V2.0 OR 50% OR GREATER REDUCTION IN PALPABLE SPLEEN LENGTH, OR REDUCTION OF 35% OR GREATER FROM BASELINE SPLEEN VOLUME AFTER 6 MONTHS OF THERAPY.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	

SAFINAMIDE MESYLATE

Products Affected

- XADAGO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

SARILUMAB

Products Affected

- KEVZARA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PREVIOUS TRIAL OF HUMIRA AND ONE OF THE FOLLOWING PREFERRED AGENTS: ACTEMRA, CIMZIA, ORENCIA, OR XELJANZ.

SEBELIPASE ALFA

Products Affected

- KANUMA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	BLOOD TEST OR DRIED BLOOD SPOT TEST INDICATING LOW OR ABSENT LYSOSOMAL ACID LIPASE DEFICIENCY (LAL) ENZYME ACTIVITY, OR A GENETIC TEST INDICATING THE PRESENCE OF ALTERED LIPA GENE(S).
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH AN ENDOCRINOLOGIST, HEPATOLOGIST, GASTROENTEROLOGIST, MEDICAL GENETICIST, LIPIDOLOGIST, OR A METABOLIC SPECIALIST.
Coverage Duration	LAL INITIAL 6 OR 12 MONTHS, SEE OTHER CRITERIA. RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	INITIAL: DIAGNOSIS OF LYSOSOMAL ACID LIPASE (LAL) DEFICIENCY, AS CONFIRMED BY THE PRESENCE OF CLINICAL FEATURES (E.G., HEPATOMEGALY, ELEVATED SERUM TRANSAMINASES, DYSLIPIDEMIA, SPLENOMEGALY) PLUS ANY OF THE FOLLOWING: A BLOOD TEST INDICATING LOW OR ABSENT LEVELS OF LAL ENZYME ACTIVITY, A DRIED BLOOD SPOT TEST INDICATING LOW OR ABSENT LAL ENZYME ACTIVITY, OR A GENETIC TEST INDICATING THE BI-ALLELIC PRESENCE OF ALTERED LIPA GENE(S). RENEWAL:DIAGNOSIS OF LYSOSOMAL ACID LIPASE (LAL) DEFICIENCY PRESENTING AFTER THE FIRST 6 MONTHS OF LIFE AND NOT CONSIDERED RAPIDLY PROGRESSIVE REQUIRES DOCUMENTED IMPROVEMENT IN ANY ONE OF THE FOLLOWING CLINICAL PARAMETERS ASSOCIATED WITH LYSOSOMAL ACID LIPASE (LAL) DEFICIENCY DURING THE PAST 6 MONTHS: A RELATIVE REDUCTION FROM BASELINE IN ANY ONE OF THE FOLLOWING LIPID LEVELS (LDL-C, NON-HDL-C, OR TRIGLYCERIDES), NORMALIZATION OF ASPARTATE AMINOTRANSFERASE (AST) BASED ON AGE- AND GENDER-SPECIFIC NORMAL RANGES, A DECREASE IN LIVER FAT CONTENT COMPARED TO BASELINE ASSESSED BY ABDOMINAL IMAGING (E.G., MULTI-ECHO GRADIENT ECHO [MEGE] MRI). DIAGNOSIS OF RAPIDLY PROGRESSIVE LYSOSOMAL ACID LIPASE (LAL) DEFICIENCY PRESENTING WITHIN THE FIRST 6 MONTHS OF LIFE: 12 MONTHS. A DIAGNOSIS OF LYSOSOMAL ACID LIPASE (LAL) DEFICIENCY PRESENTING AFTER THE FIRST 6 MONTHS OF LIFE AND NOT CONSIDERED RAPIDLY PROGRESSIVE: INITIAL: 6 MONTHS

SECUKINUMAB

Products Affected

- COSENTYX (2 SYRINGES)
- COSENTYX PEN (2 PENS)

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PLAQUE PSORIASIS (PSO): MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5 PERCENT BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PLAQUE PSORIASIS (PSO): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST OR A DERMATOLOGIST. ANKYLOSING SPONDYLITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 4 MONTHS. RENEWAL: 12 MONTHS FOR ALL DIAGNOSES
Other Criteria	INITIAL: PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA AND ONE CONVENTIONAL THERAPY SUCH AS PUVA (PHOTOTHERAPY ULTRAVIOLET LIGHT A), UVB (ULTRAVIOLET LIGHT B), TOPICAL CORTICOSTEROIDS, CALCIPOTRIENE, ACITRETIN, METHOTREXATE, OR CYCLOSPORINE. PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. ANKYLOSING SPONDYLITIS (AS): PREVIOUS TRIAL OF HUMIRA.

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
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. RENEWAL: PATIENT SHOWS 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.

SILTUXIMAB

Products Affected

- SYLVANT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

SIMEPREVIR

Products Affected

- OLYSIO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
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 (FOR EXAMPLE 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.

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. PATIENT MUST NOT BE TAKING ANY OF THE FOLLOWING INTERACTING MEDICATIONS: CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, ERYTHROMYCIN (DOES NOT INCLUDE TOPICAL FORMULATIONS), CLARITHROMYCIN, TELITHROMYCIN, ITRACONAZOLE, KETOCONAZOLE, POSACONAZOLE, FLUCONAZOLE (DOES NOT INCLUDE TOPICAL FORMULATIONS), VORICONAZOLE, DEXAMETHASONE, CISAPRIDE, CYCLOSPORINE, ROSUVASTATIN DOSE ABOVE 10MG, ATORVASTATIN DOSE ABOVE 40MG, OR ANY OF THE FOLLOWING HIV MEDICATIONS: COBICISTAT-CONTAINING MEDS (E.G., STRIBILD), ANY HIV PROTEASE INHIBITOR (ATAZANAVIR, FOSAMPRENAVIR, LOPINAVIR, INDINAVIR, NELFINAVIR, SAQUINAVIR, OR TIPRANAVIR) RITONAVIR, DARUNAVIR/RITONAVIR, DELAVIRDINE, ETRAVIRINE, NEVIRAPINE, EFAVIRENZ). PATIENT MUST ALSO NOT BE TAKING AMIODARONE IF ON COMBINATION REGIMEN OF SOVALDI AND OLYSIO.

SOFOSBUVIR

Products Affected

- SOVALDI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE AND ADDITIONAL CONSIDERATION FOR COVERAGE CONSISTENT WITH FDA LABELING.
Exclusion Criteria	PATIENT WITH END STAGE RENAL DISEASE OR REQUIRES DIALYSIS.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A 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

PA Criteria	Criteria Details
Other Criteria	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE. TRIAL OF A PREFERRED FORMULARY ALTERNATIVE INCLUDING HARVONI OR EPCLUSA WHEN THESE AGENTS ARE CONSIDERED ACCEPTABLE FOR TREATMENT OF THE SPECIFIC GENOTYPE PER AASLD/IDSA GUIDANCE. FOR PATIENTS ON SOVALDI PLUS DAKLINZA REGIMENS THERE WILL BE NO APPROVALS FOR CONCURRENT USE OF ANY OF THESE (CONTRAINDICATED OR NOT RECOMMENDED BY THE MANUFACTURER) MEDICATIONS: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, OR RIFAMPIN. REQUESTS FOR SOVALDI IN COMBINATION WITH DAKLINZA OR OLYSIO WILL REQUIRE THAT THE PATIENT ALSO MEETS ALL CRITERIA FOR THE RESPECTIVE AGENT USED (DAKLINZA OR OLYSIO).

SOFOSBUVIR/VELPATASVIR

Products Affected

- EPCLUSA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE.
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL.
Age Restrictions	18 YEARS OF AGE AND OLDER.
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. PATIENT MUST NOT HAVE SEVERE RENAL IMPAIRMENT, ESRD OR ON HEMODIALYSIS. RIBAVIRIN USE REQUIRED FOR PATIENTS WITH DECOMPENSATED CIRRHOSIS.

SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR

Products Affected

- VOSEVI

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD-IDSA GUIDANCE.
Exclusion Criteria	SEVERE RENAL IMPAIRMENT, ESRD OR ON HEMODIALYSIS. MODERATE OR SEVERE HEPATIC IMPAIRMENT (CHILD-PUGH B OR C).
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS
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. PATIENT IS NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS NOT RECOMMENDED BY THE MANUFACTURER: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, CYCLOSPORINE, PITAVASTATIN, PRAVASTATIN (DOSES ABOVE 40MG), ROSUVASTATIN, METHOTREXATE, MITOXANTRONE, IMATINIB, IRINOTECAN, LAPATINIB, SULFASALAZINE, TOPOTECAN, OR HIV REGIMEN THAT CONTAINS EFAVIRENZ, ATAZANAVIR, LOPINAVIR OR TIPRANAVIR/RITONAVIR.

SOMATROPIN - GROWTH HORMONE

Products Affected

- HUMATROPE
- OMNITROPE
- SAIZEN
- SAIZEN 8.8 MG SAIZENPREP CART
- SAIZEN CLICK.EASY
- ZOMACTON

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES, GROWTH FAILURE WITH CLOSED EPIPHYSES.
Required Medical Information	INDUCTION - PATIENT'S HEIGHT AT LEAST 2 STANDARD DEVIATIONS (SD) BELOW THE MEAN HEIGHT FOR NORMAL CHILDREN OF THE SAME AGE AND GENDER. RENEWAL: GROWTH VELOCITY AND/OR TARGET HEIGHT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: ENDOCRINOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	INITIAL: PREVIOUS TRIAL OF PREFERRED FORMULARY ALTERNATIVES NORDITROPIN AND GENOTROPIN PER FDA INDICATION. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

SOMATROPIN - SEROSTIM

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES
Required Medical Information	HIV/WASTING: MEETS CRITERIA OF 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 20 KG PER METER SQUARED.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST, NUTRITIONAL SUPPORT SPECIALIST (SBS), OR INFECTIOUS DISEASE SPECIALIST
Coverage Duration	3 MONTHS
Other Criteria	HIV/WASTING: CURRENTLY ON ANTIRETROVIRAL THERAPY. IF CURRENTLY ON GROWTH HORMONE, PATIENT HAS SHOWN CLINICAL BENEFIT IN MUSCLE MASS AND WEIGHT OR IF NOT ON GROWTH HORMONE, PATIENT HAS HAD INADEQUATE RESPONSE TO PREVIOUS THERAPY. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

SOMATROPIN - ZORBTIVE

Products Affected

- ZORBTIVE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST
Coverage Duration	SHORT BOWEL: 4 WEEKS ONCE
Other Criteria	THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

SOMATROPIN-NORDITROPIN AND GENOTROPIN

Products Affected

- GENOTROPIN
- GENOTROPIN MINIQUICK
- NORDITROPIN FLEXPRO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES, GROWTH FAILURE WITH CLOSED EPIPHYSES.
Required Medical Information	INDUCTION - PATIENT'S HEIGHT AT LEAST 2 STANDARD DEVIATIONS (SD) BELOW THE MEAN HEIGHT FOR NORMAL CHILDREN OF THE SAME AGE AND GENDER. RENEWAL: GROWTH VELOCITY AND/OR TARGET HEIGHT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: ENDOCRINOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT (I.E. INCREASED HEIGHT OR INCREASED GROWTH VELOCITY). THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

SOMATROPIN-NUTROPIN AND NUTROPIN AQ

Products Affected

- NUTROPIN AQ NUSPIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES, GROWTH FAILURE DUE TO CKD IF PATIENT HAS HAD A RENAL TRANSPLANT, OR GROWTH FAILURE WITH CLOSED EPIPHYSES.
Required Medical Information	INDUCTION - PATIENT'S HEIGHT AT LEAST 2 STANDARD DEVIATIONS (SD) BELOW THE MEAN HEIGHT FOR NORMAL CHILDREN OF THE SAME AGE AND GENDER. RENEWAL: GROWTH VELOCITY AND/OR TARGET HEIGHT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: ENDOCRINOLOGIST. FOR GROWTH HORMONE FAILURE DUE TO CRI: NEPHROLOGIST.
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	ALL DIAGNOSES EXCEPT FOR CHRONIC KIDNEY DISEASE (CKD): INITIAL: PREVIOUS TRIAL OF PREFERRED FORMULARY ALTERNATIVES NORDITROPIN AND GENOTROPIN PER FDA INDICATION. RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT (I.E. INCREASED HEIGHT OR INCREASED GROWTH VELOCITY). FOR GROWTH FAILURE SECONDARY TO CKD: PATIENT HAS NOT RECEIVED A RENAL TRANSPLANT. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

SONIDEGIB

Products Affected

- ODOMZO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	BASELINE SERUM CREATINE KINASE (CK) AND SERUM CREATININE LEVELS
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

SORAFENIB TOSYLATE

Products Affected

- NEXAVAR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

SUNITINIB MALATE

Products Affected

- SUTENT

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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.

TADALAFIL

Products Affected

- CIALIS ORAL TABLET 2.5 MG, 5 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	ERECTILE DYSFUNCTION WITHOUT DIAGNOSIS OF BENIGN PROSTATIC HYPERPLASIA.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TRIAL OF ONE FORMULARY ALPHA BLOCKER SUCH AS DOXAZOSIN, TERAZOSIN, TAMSULOSIN OR ALFUZOSIN) AND ONE FORMULARY 5-ALPHA-REDUCTASE (SUCH AS FINASTERIDE OR DUTASTERIDE). APPLIES TO 2.5MG AND 5MG STRENGTHS ONLY. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION.

TALIMOGENE

Products Affected

- IMLYGIC INJECTION SUSPENSION 10EXP6 (1 MILLION) PFU/ML, 10EXP8 (100 MILLION) PFU/ML

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	HISTORY OF PRIMARY OR ACQUIRED IMMUNODEFICIENT STATES, LEUKEMIA, LYMPHOMA, OR AIDS. PATIENT IS NOT CURRENTLY RECEIVING IMMUNOSUPPRESSIVE THERAPY.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	IMLYGIC TO BE INJECTED INTO CUTANEOUS, SUBCUTANEOUS, AND OR NODAL LESIONS THAT ARE VISIBLE, PALPABLE, OR DETECTABLE BY ULTRASOUND GUIDANCE. NO CONCURRENT USE WITH PEMBROLIZUMAB, NIVOLUMAB, IPILIMUMAB, DABRAFENIB, TRAMETINIB, VEMURAFENIB, INTERLEUKIN-2, INTERFERON, DACARBAZINE, TEMOZOLOMIDE, PACLITAXEL, CARBOPLATIN, IMATINIB, MELPHALAN, IMIQUIMOD, OR RADIATION THERAPY.

TASIMELTEON

Products Affected

- HETLIOZ

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TEDUGLUTIDE

Products Affected

- GATTEX 30-VIAL

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	18 YEARS OF AGE AND OLDER
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.

TELOTRISTAT

Products Affected

- XERMELO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TEMOZOLOMIDE

Products Affected

- TEMODAR INTRAVENOUS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TERIFLUNOMIDE

Products Affected

- AUBAGIO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TERIPARATIDE

Products Affected

- FORTEO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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) ADEQUATE TRIAL OF, INTOLERANCE TO, OR A CONTRAINDICATION TO BISPHOSPHONATES (E.G., ALENDRONATE, RISEDRONATE, IBANDRONATE).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TESTOSTERONE

Products Affected

- ANDRODERM
- ANDROGEL TRANSDERMAL GEL IN METERED-DOSE PUMP 20.25 MG/1.25 GRAM (1.62 %)
- ANDROGEL TRANSDERMAL GEL IN PACKET 1.62 % (20.25 MG/1.25 GRAM), 1.62 % (40.5 MG/2.5 GRAM)
- *testosterone cypionate*
- *testosterone enanthate*
- *testosterone transdermal gel in packet*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D. ADDITIONAL CONSIDERATION FOR GENDER DYSPHORIA.
Exclusion Criteria	
Required Medical Information	MALE HYPOGONADISM CONFIRMED BY EITHER: 1) LAB CONFIRMED TOTAL SERUM TESTOSTERONE LEVEL OF LESS THAN 300 NG/DL OR 2) A LOW TOTAL SERUM TESTOSTERONE LEVEL AS INDICATED BY A LAB RESULT WITH A REFERENCE RANGE OBTAINED WITHIN 90 DAYS, OR 3) A FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 PG/ML.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	LIFETIME OF MEMBERSHIP IN PLAN
Other Criteria	

TETRABENAZINE

Products Affected

- *tetrabenazine*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	NEUROLOGIST
Coverage Duration	12 MONTHS
Other Criteria	

TEZACAFTOR/IVACAFTOR

Products Affected

- SYMDEKO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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: 12 MONTHS
Other Criteria	RENEWAL: MAINTAINED OR IMPROVEMENT IN FEV1 OR BODY MASS INDEX (BMI), OR REDUCTION IN NUMBER OF PULMONARY EXACERBATIONS.

THALIDOMIDE

Products Affected

- THALOMID

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TOCILIZUMAB IV

Products Affected

- ACTEMRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL FOR RA, PJIA, OR SJIA: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	MODERATE TO SEVERE RHEUMATOID ARTHRITIS (RA)/POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA)/ SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST
Coverage Duration	INITIAL: RA: 7 MONTHS. PJIA: 5 MOS. SJIA: 12 MOS. CRS: 1 MO. RENEWAL: 12 MOS FOR RA, PJIA, OR SJIA
Other Criteria	INITIAL: MODERATE TO SEVERE RA AND PJIA: PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) AGENT SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE. INITIAL SJIA: PREVIOUS TRIAL WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) AGENT SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE.

TOCILIZUMAB SQ

Products Affected

- ACTEMRA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RA AND PJIA RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RA AND PJIA: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: RA: 7 MONTHS. PJIA: 5 MONTHS. GCA: 12 MONTHS. ALL RENEWAL: 12 MONTHS
Other Criteria	RA AND PJIA INITIAL: PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) AGENT SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE.

TOFACITINIB

Products Affected

- XELJANZ ORAL TABLET 10 MG, 5 MG
- XELJANZ XR

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	RENEWAL: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	RHEUMATOID ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS: PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST.
Coverage Duration	INITIAL: RA: 6 MOS. PSA: 4 MOS. RENEWAL (ALL INDICATIONS): 12 MOS.
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA) AND PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA AND ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) SUCH AS METHOTREXATE, LEFLUNOMIDE, HYDROXYCHLOROQUINE, OR SULFASALAZINE.

TOLVAPTAN

Products Affected

- JYNARQUE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TOPICAL TRETINOIN

Products Affected

- *tretinoin topical cream*
- *tretinoin topical gel 0.01 %, 0.025 %*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	WRINKLES, PHOTOAGING, MELASMA.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TRABECTEDIN

Products Affected

- YONDELIS

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TRAMETINIB DIMETHYL SULFOXIDE

Products Affected

- MEKINIST ORAL TABLET 0.5 MG, 2 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

TRASTUZUMAB

Products Affected

- HERCEPTIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	BREAST CANCER, METASTATIC BREAST CANCER, GASTRIC CANCER: HER2 POSITIVE
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	B VS D COVERAGE CONSIDERATION.

TREPROSTINIL DIOLAMINE

Products Affected

- ORENITRAM

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	PATIENT DOES NOT HAVE SEVERE HEPATIC IMPAIRMENT.
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) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. PREVIOUS OR CURRENT TREATMENT WITH ONE OF THE FOLLOWING AGENTS: A FORMULARY PHOSPHODIESTERASE-5 INHIBITOR (E.G., SILDENAFIL [GENERIC FOR REVATIO] OR ADCIRCA [TADALAFIL]) OR AN ENDOTHELIN RECEPTOR ANTAGONIST (E.G., TRACLEER [BOSENTAN], LETAIRIS [AMBRISENTAN], OR OPSUMIT [MACITENTAN]). TRIAL OF A FORMULARY PHOSPHODIESTERASE-5 INHIBITOR OR ENDOTHELIN RECEPTOR ANTAGONIST IS NOT REQUIRED IF THE PATIENT WAS PREVIOUSLY STABLE ON ORENITRAM. RENEWAL: PATIENT SHOWS 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.

TREPROSTINIL INHALED

Products Affected

- TYVASO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION. PATIENT HAS NYHA-WHO FUNCTIONAL CLASS III-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS
Other Criteria	THIS DRUG MAYBE COVERED UNDER MEDICARE PART B OR D DEPENDING UPON THE CIRCUMSTANCES. INFORMATION MAY NEED TO BE SUBMITTED DESCRIBING THE USE AND SETTING OF THE DRUG TO MAKE THE DETERMINATION. NEBULIZER THERAPY IS COVERED UNDER PART B FOR PATIENTS WHO ARE USING THE MEDICATION VIA A NEBULIZER IN THEIR OWN HOME. THOSE WHO ARE NOT USING IT IN THEIR HOME WILL BE COVERED UNDER PART D. INITIAL: MEAN PULMONARY ARTERY PRESSURE (PAP) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. 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.

TREPROSTINIL SODIUM INJECTABLE

Products Affected

- REMODULIN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	COVERED UNDER LOCAL COVERAGE POLICY OF APPLICABLE MEDICARE DMERC.
Required Medical Information	FORMULARY DRUG ADMINISTERED IN A LONG TERM CARE FACILITY TO A PATIENT WHOSE PART A COVERAGE HAS EXPIRED OR FORMULARY DRUG NOT ADMINISTERED VIA AN IMPLANTABLE PUMP OR AN EXTERNAL PUMP OR DRUG ADMINISTERED VIA AN IMPLANTABLE PUMP/AN EXTERNAL PUMP. DOCUMENTED CONFIRMATORY PULMONARY ARTERIAL HYPERTENSION (PAH) DIAGNOSIS BASED ON RIGHT HEART CATHETERIZATION.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST
Coverage Duration	INITIAL AND RENEWAL: 12 MONTHS

PA Criteria	Criteria Details
Other Criteria	INITIAL: MEAN PULMONARY ARTERY PRESSURE (PAP) OF AT LEAST 25 MMHG OR GREATER, PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 3 WOOD UNITS. CONTINUATION OF CURRENT REMODULIN THERAPY: PATIENT MUST HAVE NYHA/WHO FC II-IV SYMPTOMS. NEW REQUESTS FOR REMODULIN THERAPY: PATIENT MUST HAVE NYHA/WHO FC III-IV SYMPTOMS. NEW REQUESTS FOR REMODULIN THERAPY FOR PATIENTS WITH NYHA/WHO FC II SYMPTOMS REQUIRES A TRIAL OF OR CONTRAINDICATION TO A PHOSPHODIESTERASE-5 INHIBITOR (PDE-5) (E.G., REVATIO, ADCIRCA) OR AN ENDOTHELIN RECEPTOR ANTAGONIST (ERA) (E.G., LETAIRIS, OPSUMIT, TRACLEER). 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.

TRIENTINE

Products Affected

- *trientine*

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	KNOWN FAMILY HISTORY OF WILSON'S DISEASE OR PHYSICAL EXAMINATION CONSISTENT WITH WILSON'S DISEASE. PLASMA COPPER-PROTEIN CERULOPLASMIN LESS THAN 20 MG/DL. LIVER BIOPSY POSITIVE FOR AN ABNORMALLY HIGH CONCENTRATION OF COPPER (GREATER THAN 250 MCG/G DRY WEIGHT) OR THE PRESENCE OF KAYSER-FLEISCHER RINGS.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A HEPATOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	PREVIOUS TRIAL OF OR CONTRAINDICATION TO PENICILLAMINE (DEPEN).

TRIFLURIDINE/TIPIRACIL

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

URIDINE TRIACETATE

Products Affected

- XURIDEN

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	

USTEKINUMAB

Products Affected

- STELARA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS: MODERATE TO SEVERE PLAQUE PSORIASIS INVOLVING GREATER THAN OR EQUAL TO 5% BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, OR GENITAL AREA. RENEWAL FOR PSORIATIC ARTHRITIS OR PLAQUE PSORIASIS: PHYSICIAN ATTESTATION OF IMPROVEMENT.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH: PSORIATIC ARTHRITIS: DERMATOLOGIST OR RHEUMATOLOGIST. PLAQUE PSORIASIS: DERMATOLOGIST. CROHN'S DISEASE: GASTROENTEROLOGIST.
Coverage Duration	INITIAL: PSA, PSO, CD: 4 MONTHS. CD WITH PREVIOUS DOSE IV: 2 MONTHS. RENEW ALL: 12 MO
Other Criteria	INITIAL: PSORIATIC ARTHRITIS (PSA): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: CIMZIA, OTEZLA, OR COSENTYX. PLAQUE PSORIASIS (PSO): PREVIOUS TRIAL OF HUMIRA FOLLOWED BY ONE OF THE FOLLOWING PREFERRED AGENTS: COSENTYX OR OTEZLA. CROHN'S DISEASE (CD): PREVIOUS TRIAL OF OR CONTRAINDICATION TO PREFERRED TNF INHIBITORS: HUMIRA FOLLOWED BY CIMZIA.

USTEKINUMAB IV

Products Affected

- STELARA

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	2 MONTHS
Other Criteria	PREVIOUS TRIAL OF OR CONTRAINDICATION TO THE PREFERRED TNF INHIBITORS: HUMIRA FOLLOWED BY CIMZIA. THIS DRUG ALSO REQUIRES PAYMENT DETERMINATION AND MAY BE COVERED UNDER MEDICARE PART B OR D.

VALBENAZINE TOSYLATE

Products Affected

- INGREZZA ORAL CAPSULE 40 MG, 80 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	PATIENT HAS A PRIOR HISTORY OF USING ANTIPSYCHOTIC MEDICATIONS OR METOCLOPRAMIDE PER PHYSICIAN ATTESTATION.
Age Restrictions	
Prescriber Restrictions	PRESCRIBED BY OR GIVEN IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR MOVEMENT DISORDER SPECIALIST.
Coverage Duration	12 MONTHS
Other Criteria	

VANDETANIB

Products Affected

- CAPRELSA ORAL TABLET 100 MG, 300 MG

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

VEMURAFENIB

Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

VENETOCLAX

Products Affected

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

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

VINCRISTINE SULFATE LIPOSOMAL

Products Affected

- MARQIBO

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
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.

VISMODEGIB

Products Affected

- ERIVEDGE

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

ZIV-AFLIBERCEPT

Products Affected

- ZALTRAP

PA Criteria	Criteria Details
Covered Uses	ALL FDA APPROVED INDICATIONS NOT OTHERWISE EXCLUDED FROM PART D.
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	

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