

ABALOPARATIDE

Products Affected

- TYMLOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 MONTHS
Other Criteria	OSTEOPOROSIS: HAS NOT RECEIVED A TOTAL OF 24 MONTHS CUMULATIVE TREATMENT WITH ANY PARATHYROID HORMONE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ABATACEPT IV

Products Affected

- ORENCIA (WITH MALTOSE)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	RA, PJIA, PSA: INITIAL: 6 MOS, RENEWAL: 12 MOS. ACUTE GRAFT VERSUS HOST DISEASE (AGVHD): 1 MO.
Other Criteria	INITIAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. RENEWAL: RA, PJIA, PSA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

ABATACEPT SQ

Products Affected

- ORENCIA
- ORENCIA CLICKJECT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, PJIA, PSA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ABEMACICLIB

Products Affected

- VERZENIO

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

ABIRATERONE

Products Affected

- *abiraterone*
- *abirtega*

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

ABIRATERONE SUBMICRONIZED

Products Affected

- *abiraterone, submicronized*
- YONSA

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

ACALABRUTINIB

Products Affected

- CALQUENCE (ACALABRUTINIB MAL)

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

ACORAMIDIS

Products Affected

- ATTRUBY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CARDIOMYOPATHY OF WILD TYPE OR VARIANT TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM): INITIAL: 1) NEW YORK HEART ASSOCIATION (NYHA) CLASS I, II, OR III HEART FAILURE, AND 2) DIAGNOSIS CONFIRMED BY (A) BONE SCAN (SCINTIGRAPHY) STRONGLY POSITIVE FOR MYOCARDIAL UPTAKE OF TC-99M-PYP, OR (B) BIOPSY OF TISSUE OF AFFECTED ORGAN(S) (CARDIAC AND POSSIBLY NON-CARDIAC SITES) TO CONFIRM AMYLOID PRESENCE AND CHEMICAL TYPING TO CONFIRM PRESENCE OF TRANSTHYRETIN (TTR) PROTEIN.
Age Restrictions	
Prescriber Restrictions	ATTR-CM: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST, ATTR SPECIALIST, OR MEDICAL GENETICIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	ATTR-CM: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER ATTR-CM TTR STABILIZERS (E.G., TAFAMIDIS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ADAGRASIB

Products Affected

- KRAZATI

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

ADALIMUMAB

Products Affected

- HUMIRA PEN
- HUMIRA PEN CROHNS-UC-HS START
- HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML
- HUMIRA(CF)
- HUMIRA(CF) PEDI CROHNS STARTER
- HUMIRA(CF) PEN
- HUMIRA(CF) PEN CROHNS-UC-HS
- HUMIRA(CF) PEN PEDIATRIC UC
- HUMIRA(CF) PEN PSOR-UV-ADOL HS

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

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. UVEITIS: DOES NOT HAVE ISOLATED ANTERIOR UVEITIS. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), PSA, AS, PSO, HS, UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ADALIMUMAB-AATY

Products Affected

- *adalimumab-aaty*
- *adalimumab-aaty(cf) ai crohns*
- YUFLYMA(CF)
- YUFLYMA(CF) AI CROHN'S-UC-HS
- YUFLYMA(CF) AUTOINJECTOR

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

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. UVEITIS: DOES NOT HAVE ISOLATED ANTERIOR UVEITIS. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), PSA, AS, PSO, HS, UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ADALIMUMAB-ADBIM

Products Affected

- CYLTEZO(CF)
- CYLTEZO(CF) PEN
- CYLTEZO(CF) PEN CROHN'S-UC-HS
- CYLTEZO(CF) PEN PSORIASIS-UV

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

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. UVEITIS: DOES NOT HAVE ISOLATED ANTERIOR UVEITIS. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), PSA, AS, PSO, HS, UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ADALIMUMAB-BWWD

Products Affected

- HADLIMA
- HADLIMA PUSHTOUCH
- HADLIMA(CF)
- HADLIMA(CF) PUSHTOUCH

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

PA Criteria	Criteria Details
Other Criteria	INITIAL: RHEUMATOID ARTHRITIS (RA): TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. UVEITIS: DOES NOT HAVE ISOLATED ANTERIOR UVEITIS. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), PSA, AS, PSO, HS, UVEITIS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

AFATINIB

Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ALECTINIB

Products Affected

- ALECENSA

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

ALPELISIB-PIQRAY

Products Affected

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

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

AMIKACIN LIPOSOMAL INH

Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE: RENEWAL: 1) NO POSITIVE MAC SPUTUM CULTURE AFTER CONSECUTIVE NEGATIVE CULTURES, AND 2) IMPROVEMENT IN SYMPTOMS. ADDITIONALLY, FOR FIRST RENEWAL, APPROVAL REQUIRES AT LEAST ONE NEGATIVE SPUTUM CULTURE FOR MAC BY SIX MONTHS OF ARIKAYCE TREATMENT. FOR SECOND AND SUBSEQUENT RENEWALS, APPROVAL REQUIRES AT LEAST THREE NEGATIVE SPUTUM CULTURES FOR MAC BY 12 MONTHS OF ARIKAYCE TREATMENT.
Age Restrictions	
Prescriber Restrictions	MAC LUNG DISEASE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	INITIAL/RENEWAL: 6 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

AMIVANTAMAB-HYALURONIDASE-LPUJ

Products Affected

- RYBREVANT FASPRO

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

AMIVANTAMAB-VMJW

Products Affected

- RYBREVANT

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

ANAKINRA

Products Affected

- KINERET

PA Criteria	Criteria Details
Exclusion Criteria	CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS.
Required Medical Information	INITIAL: CRYOPYRIN-ASSOCIATED PERIODIC SYNDROMES (CAPS): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE NLRP3 GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR, SERUM AMYLOID A PROTEIN (SAA) OR S100 PROTEINS), AND 2) TWO OF THE FOLLOWING: URTICARIAL-LIKE RASH (NEUTROPHILIC DERMATITIS), COLD-TRIGGERED EPISODES, SENSORINEURAL HEARING LOSS, MUSCULOSKELETAL SYMPTOMS, CHRONIC ASEPTIC MENINGITIS, SKELETAL ABNORMALITIES. DEFICIENCY OF INTERLEUKIN-1 RECEPTOR ANTAGONIST (DIRA): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE IL1RN GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR), AND 2) ONE OF THE FOLLOWING: PUSTULAR PSORIASIS-LIKE RASHES, OSTEOMYELITIS, ABSENCE OF BACTERIAL OSTEOMYELITIS, ONYCHOMADESIS.
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	RA: INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS. CAPS, DIRA: LIFETIME.

PA Criteria	Criteria Details
Other Criteria	INITIAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. RENEWAL: RA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

APALUTAMIDE

Products Affected

- ERLEADA ORAL TABLET 240 MG, 60 MG

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

APOMORPHINE - ONAPGO

Products Affected

- ONAPGO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PD: RENEWAL: IMPROVEMENT IN MOTOR SYMPTOMS WHILE ON THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

APREMILAST

Products Affected

- OTEZLA
- OTEZLA STARTER
- OTEZLA XR
- OTEZLA XR INITIATION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: MILD PLAQUE PSORIASIS (PSO): 1) PSORIASIS COVERING LESS THAN 3 PERCENT OF BODY SURFACE AREA (BSA), 2) STATIC PHYSICIAN GLOBAL ASSESSMENT (SPGA) SCORE OF 2, OR 3) PSORIASIS AREA AND SEVERITY INDEX (PASI) SCORE OF 2 TO 9. MODERATE TO SEVERE PSO: PSORIASIS COVERING 3 PERCENT OR MORE OF BSA, OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. BEHCETS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: MILD PSO: TRIAL OF OR CONTRAINDICATION TO ONE CONVENTIONAL SYSTEMIC THERAPY (E.G., METHOTREXATE, ACITRETIN, CYCLOSPORINE) OR ONE CONVENTIONAL TOPICAL THERAPY (E.G., TOPICAL CORTICOSTEROIDS). MODERATE TO SEVERE PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. BEHCETS DISEASE: 1) HAS ORAL ULCERS OR A HISTORY OF RECURRENT ORAL ULCERS BASED ON CLINICAL SYMPTOMS, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OR MORE CONSERVATIVE TREATMENTS (E.G., COLCHICINE, TOPICAL CORTICOSTEROID, ORAL CORTICOSTEROID). INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: ALL INDICATIONS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ARIMOCLOMOL

Products Affected

- MIPLYFFA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	NIEMANN-PICK DISEASE TYPE C (NPC): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH NEUROLOGIST OR GENETICIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	NPC: RENEWAL: IMPROVEMENT OR SLOWING OF DISEASE PROGRESSION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ASCIMINIB

Products Affected

- SCEMBLIX ORAL TABLET 100 MG, 20 MG, 40 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED OR T315I MUTATION PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND SCEMBLIX IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ASFOTASE ALFA

Products Affected

- STRENSIQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HYPOPHOSPHATASIA (HPP): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST, GENETICIST, OR METABOLIC SPECIALIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: PERINATAL/INFANTILE-ONSET HPP: 1) 6 MONTHS OF AGE OR YOUNGER AT ONSET OF HPP, AND 2) POSITIVE FOR A TISSUE NON-SPECIFIC ALKALINE PHOSPHATASE (TNSALP) (ALPL) GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR TWO OF THE FOLLOWING: (A) SERUM ALKALINE PHOSPHATASE (ALP) LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE, (B) ELEVATED SERUM PYRIDOXAL-5'-PHOSPHATE (PLP) LEVELS AND NO VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK, (C) URINE PHOSPHOETHANOLAMINE (PEA) LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE, (D) RADIOGRAPHIC EVIDENCE OF HPP, (E) AT LEAST TWO OF THE FOLLOWING: (I) RACHITIC CHEST DEFORMITY, (II) CRANIOSYNOSTOSIS, (III) DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, (IV) HISTORY OF VITAMIN B6 DEPENDENT SEIZURES, (V) NEPHROCALCINOSIS OR HISTORY OF ELEVATED SERUM CALCIUM, (VI) HISTORY OR PRESENCE OF NON-TRAUMATIC POSTNATAL FRACTURE AND DELAYED FRACTURE HEALING. JUVENILE-ONSET HPP: 1) 18 YEARS OF AGE OR YOUNGER AT ONSET OF HPP, AND 2) POSITIVE FOR A TNSALP ALPL GENE MUTATION AS CONFIRMED BY GENETIC TESTING OR TWO OF THE FOLLOWING: (A) SERUM ALP LEVEL BELOW THAT OF NORMAL RANGE FOR PATIENT AGE, (B) ELEVATED SERUM PLP LEVELS AND NO VITAMIN B6 SUPPLEMENTATION IN THE PREVIOUS WEEK, (C) URINE PEA LEVEL ABOVE THAT OF NORMAL RANGE FOR PATIENT AGE, (D) RADIOGRAPHIC EVIDENCE OF HPP, (E) AT LEAST TWO OF THE FOLLOWING: (I) RACHITIC DEFORMITIES, (II) PREMATURE LOSS OF PRIMARY TEETH PRIOR TO 5 YEARS OF AGE, (III) DELAY IN SKELETAL GROWTH RESULTING IN DELAY OF MOTOR DEVELOPMENT, (IV) HISTORY OR PRESENCE OF NON-TRAUMATIC FRACTURES OR DELAYED FRACTURE HEALING. ALL INDICATIONS: 1) NOT CURRENTLY RECEIVING TREATMENT WITH A BISPHOSPHONATE, 2) CALCIUM OR PHOSPHATE LEVELS ARE NOT BELOW THE NORMAL RANGE, 3)
	NOT HAVE A TREATABLE FORM OF RICKETS. RENEWAL: ALL INDICATIONS: 1) IMPROVEMENT IN THE SKELETAL CHARACTERISTICS OF HPP, AND 2) NOT CURRENTLY RECEIVING TREATMENT WITH A BISPHOSPHONATE.
Indications	All FDA-approved Indications.
Off Label Uses	

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

ATOGEPANT

Products Affected

- QULIPTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MIGRAINE PREVENTION: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

AVACOPAN

Products Affected

- TAVNEOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	ANTI-NEUTROPHIL CYTOPLASMIC AUTOANTIBODY (ANCA)-ASSOCIATED VASCULITIS: INITIAL: ANCA SEROPOSITIVE (ANTI-PR3 OR ANTI-MPO).
Age Restrictions	
Prescriber Restrictions	ANCA-ASSOCIATED VASCULITIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 6 MONTHS.
Other Criteria	ANCA-ASSOCIATED VASCULITIS: RENEWAL: CONTINUES TO BENEFIT FROM THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

AVAPRITINIB

Products Affected

- AYVAKIT

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

AVUTOMETINIB-DEFACTINIB

Products Affected

- AVMAPKI
- AVMAPKI-FAKZYNJA
- FAKZYNJA

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

AXATILIMAB-CSFR

Products Affected

- NIKTIMVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CHRONIC GRAFT VS HOST DISEASE (CGVHD): 1) FAILURE OF AT LEAST TWO LINES OF SYSTEMIC THERAPY, ONE OF WHICH MUST BE A TRIAL OF OR CONTRAINDICATION TO JAKAFI, AND 2) NO CONCURRENT USE WITH JAKAFI, REZUROCK, OR IMBRUVICA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

AXITINIB

Products Affected

- INLYTA ORAL TABLET 1 MG, 5 MG

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

AZACITIDINE

Products Affected

- ONUREG

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

AZTREONAM INHALED

Products Affected

- CAYSTON

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	7 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

BEDAQUILINE

Products Affected

- SIRTURO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 WEEKS
Other Criteria	N/A
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

BELIMUMAB

Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: SYSTEMIC LUPUS ERYTHEMATOSUS (SLE): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. LUPUS NEPHRITIS (LN): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: SLE: CURRENTLY TAKING CORTICOSTEROIDS, ANTIMALARIALS, NSAIDS, OR IMMUNOSUPPRESSIVE AGENTS. RENEWAL: SLE: PATIENT HAD CLINICAL IMPROVEMENT. LN: IMPROVEMENT IN RENAL RESPONSE FROM BASELINE LABORATORY VALUES (I.E., EGFR OR PROTEINURIA) AND/OR CLINICAL PARAMETERS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

BELUMOSUDIL

Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CHRONIC GRAFT VS HOST DISEASE (CGVHD): 1) FAILURE OF AT LEAST TWO LINES OF SYSTEMIC THERAPY, ONE OF WHICH MUST BE A TRIAL OF OR CONTRAINDICATION TO JAKAFI, AND 2) NO CONCURRENT USE WITH JAKAFI, NIKTIMVO, OR IMBRUVICA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

BELZUTIFAN

Products Affected

- WELIREG

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

BENDAMUSTINE

Products Affected

- *bendamustine intravenous recon soln*
- BENDAMUSTINE INTRAVENOUS SOLUTION
- BENDEKA
- VIVIMUSTA

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

BENRALIZUMAB

Products Affected

- FASENRA
- FASENRA PEN

PA Criteria	Criteria Details
Exclusion Criteria	PA Criteria: Pending CMS Approval
Required Medical Information	PA Criteria: Pending CMS Approval
Age Restrictions	PA Criteria: Pending CMS Approval
Prescriber Restrictions	PA Criteria: Pending CMS Approval
Coverage Duration	PA Criteria: Pending CMS Approval
Other Criteria	PA Criteria: Pending CMS Approval
Indications	PA Criteria: Pending CMS Approval
Off Label Uses	PA Criteria: Pending CMS Approval
Part B Prerequisite	PA Criteria: Pending CMS Approval
Prerequisite Therapy Required	PA Criteria: Pending CMS Approval

BETAINE

Products Affected

- *betaine*

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

BEVACIZUMAB-BVZR

Products Affected

- ZIRABEV

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

BEXAROTENE

Products Affected

- *bexarotene*

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

BINIMETINIB

Products Affected

- MEKTOVI

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

BORTEZOMIB

Products Affected

- *bortezomib injection*
- BORUZU

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

BOSENTAN

Products Affected

- *bosentan oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PAH: INITIAL: 1) DOES NOT HAVE ELEVATED LIVER ENZYMES (ALT, AST) MORE THAN 3 TIMES UPPER LIMIT OF NORMAL (ULN) OR INCREASE IN BILIRUBIN BY 2 OR MORE TIMES ULN, AND 2) NO CONCURRENT USE WITH CYCLOSPORINE A OR GLYBURIDE. RENEWAL: NO CONCURRENT USE WITH CYCLOSPORINE A OR GLYBURIDE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

BOSUTINIB

Products Affected

- BOSULIF ORAL CAPSULE 100 MG, 50 MG
- BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG
- *bosutinib oral tablet 100 mg, 400 mg, 500 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND BOSULIF IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

BRIGATINIB

Products Affected

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

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

C1 ESTERASE INHIBITOR-HAEGARDA

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HEREDITARY ANGIOEDEMA (HAE): INITIAL: 1) TYPE III HAE, OR 2) TYPE I OR II HAE CONFIRMED BY ONE OF THE FOLLOWING COMPLEMENT TESTS: C1-INH PROTEIN LEVELS, C4 PROTEIN LEVELS, C1-INH FUNCTIONAL LEVELS, C1Q.
Age Restrictions	
Prescriber Restrictions	HAE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, IMMUNOLOGIST, ALLERGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	HAE: INITIAL/RENEWAL: NOT ON CONCURRENT TREATMENT WITH ALTERNATIVE PROPHYLACTIC AGENT FOR HAE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

CABOZANTINIB CAPSULE

Products Affected

- COMETRIQ ORAL CAPSULE 100 MG/DAY(80 MG X1-20 MG X1), 140 MG/DAY(80 MG X1-20 MG X3), 60 MG/DAY (20 MG X 3/DAY)

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

CABOZANTINIB TABLET

Products Affected

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

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

CANNABIDIOL

Products Affected

- EPIDIOLEX

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

CAPIVASERTIB

Products Affected

- TRUQAP

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

CAPMATINIB

Products Affected

- TABRECTA

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

CARGLUMIC ACID

Products Affected

- *carglumic acid*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ACUTE OR CHRONIC HYPERAMMONEMIA (HA) DUE TO N ACETYLGLUTAMATE SYNTHASE (NAGS) DEFICIENCY: NAGS GENE MUTATION IS CONFIRMED BY BIOCHEMICAL OR GENETIC TESTING. ACUTE HA DUE TO PROPIONIC ACIDEMIA (PA): 1) CONFIRMED BY ELEVATED METHYLCITRIC ACID AND NORMAL METHYLMALONIC ACID, OR 2) GENETIC TESTING CONFIRMS MUTATION IN THE PCCA OR PCCB GENE. ACUTE HA DUE TO METHYLMALONIC ACIDEMIA (MMA): 1) CONFIRMED BY ELEVATED METHYLMALONIC ACID, METHYLCITRIC ACID, OR 2) GENETIC TESTING CONFIRMS MUTATION IN THE MMUT, MMA, MMAB OR MMADHC GENES.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ACUTE HA DUE TO NAGS/PA/MMA: 7 DAYS. CHRONIC HA DUE TO NAGS: INITIAL: 6 MOS, RENEWAL: 12 MOS.
Other Criteria	RENEWAL: CHRONIC HA DUE TO NAGS: PATIENT HAS SHOWN CLINICAL IMPROVEMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

CERITINIB

Products Affected

- ZYKADIA

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

CERTOLIZUMAB PEGOL

Products Affected

- CIMZIA
- CIMZIA POWDER FOR RECONST
- CIMZIA STARTER KIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA. NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. PSA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, XELJANZ, RINVOQ, SKYRIZI, TREMFYA, ORENCIA, OTEZLA. AS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ. CD: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, RINVOQ, SKYRIZI, TREMFYA. PSO: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, SKYRIZI, TREMFYA, OTEZLA. NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PJIA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ IR, ORENCIA, RINVOQ. INITIAL FOR RA, PSA, PSO, AS, CD, PJIA: TRIAL OF OR CONTRAINDICATION TO THE STEP AGENTS IS NOT REQUIRED IF THE PATIENT IS PREGNANT, BREASTFEEDING, OR TRYING TO BECOME PREGNANT. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL FOR RA, PSA, AS, PSO, NR-AXSPA, PJIA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

CETUXIMAB

Products Affected

- ERBITUX

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

CLADRIBINE

Products Affected

- *cladribine 10 mg x 10 tab pk outer* (8 pack), 10 mg (9 pack)
- *cladribine(multiple sclerosis) oral tablet 10 mg (10 pack), 10 mg (4 pack), 10 mg (5 pack), 10 mg (6 pack), 10 mg (7 pack), 10 mg*

PA Criteria	Criteria Details
Exclusion Criteria	PA Criteria: Pending CMS Approval
Required Medical Information	PA Criteria: Pending CMS Approval
Age Restrictions	PA Criteria: Pending CMS Approval
Prescriber Restrictions	PA Criteria: Pending CMS Approval
Coverage Duration	PA Criteria: Pending CMS Approval
Other Criteria	PA Criteria: Pending CMS Approval
Indications	PA Criteria: Pending CMS Approval
Off Label Uses	PA Criteria: Pending CMS Approval
Part B Prerequisite	PA Criteria: Pending CMS Approval
Prerequisite Therapy Required	PA Criteria: Pending CMS Approval

CLOBAZAM-SYMPAZAN

Products Affected

- SYMPAZAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: LENNOX-GASTAUT SYNDROME (LGS): THERAPY IS PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	LGS: INITIAL: CONTRAINDICATION TO OR UNABLE TO SWALLOW CLOBAZAM TABLETS OR SUSPENSION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

COBIMETINIB

Products Affected

- COTELLIC

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

CORTICOTROPIN

Products Affected

- CORTROPHIN GEL INJECTION

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL: NOT APPROVED FOR DIAGNOSTIC PURPOSES.
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MULTIPLE SCLEROSIS (MS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, ALLERGIST/IMMUNOLOGIST, OPHTHALMOLOGIST, PULMONOLOGIST OR NEPHROLOGIST.
Coverage Duration	INFANTILE SPASMS AND MS: 28 DAYS. ALL OTHER FDA APPROVED INDICATIONS: INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS: TRIAL OF OR CONTRAINDICATION TO INTRAVENOUS (IV) CORTICOSTEROIDS. RENEWAL: ALL FDA APPROVED INDICATIONS EXCEPT INFANTILE SPASMS AND MS: DEMONSTRATED CLINICAL BENEFIT WHILE ON THERAPY AS INDICATED BY SYMPTOM RESOLUTION AND/OR NORMALIZATION OF LABORATORY TESTS. PART B BEFORE PART D STEP THERAPY, APPLIES ONLY TO BENEFICIARIES IN AN MA-PD PLAN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	Yes

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

CRIZOTINIB CAPSULE

Products Affected

- XALKORI ORAL CAPSULE

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

CRIZOTINIB PELLETS

Products Affected

- XALKORI ORAL PELLET 150 MG, 20 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	NON-SMALL CELL LUNG CANCER (NSCLC), ANAPLASTIC LARGE CELL LYMPHOMA (ALCL), INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT): UNABLE TO SWALLOW CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

DABRAFENIB CAPSULES

Products Affected

- TAFINLAR ORAL CAPSULE

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

DABRAFENIB SUSPENSION

Products Affected

- TAFINLAR ORAL TABLET FOR SUSPENSION

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

DACOMITINIB

Products Affected

- VIZIMPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC): NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE-INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

DALFAMPRIDINE

Products Affected

- *dalfampridine*

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

DAROLUTAMIDE

Products Affected

- NUBEQA

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

DASATINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND DASATINIB IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

DATOPOTAMAB DERUXTECAN-DLNK

Products Affected

- DATROWAY

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

DECITABINE/CEDAZURIDINE

Products Affected

- INQOVI

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

DEFERASIROX

Products Affected

- *deferasirox oral granules in packet*
- *deferasirox oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 1000 MCG/L. CHRONIC IRON OVERLOAD IN NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT): 1) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS), AND 2) LIVER IRON CONCENTRATION (LIC) OF 5 MG FE/G OF LIVER DRY WEIGHT OR GREATER. RENEWAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS: SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 500 MCG/L. NTDT: 1) SERUM FERRITIN LEVEL CONSISTENTLY ABOVE 300 MCG/L (AT LEAST TWO LAB VALUES IN THE PREVIOUS THREE MONTHS) OR 2) LIC OF 3 MG FE/G OF LIVER DRY WEIGHT OR GREATER.
Age Restrictions	
Prescriber Restrictions	INITIAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS OR NTDT: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR HEMATOLOGIST/ONCOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: CHRONIC IRON OVERLOAD DUE TO BLOOD TRANSFUSIONS OR NTDT: DEFERASIROX SPRINKLE PACKETS: TRIAL OF OR CONTRAINDICATION TO GENERIC DEFERASIROX ORAL TABLET OR TABLET FOR ORAL SUSPENSION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

DENOSUMAB-BMWO - OSENVELT

Products Affected

- OSENVELT

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

DEUTETRABENAZINE

Products Affected

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

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

DICLOFENAC TOPICAL SOLUTION

Products Affected

- *diclofenac sodium topical solution in metered-dose pump*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	OSTEOARTHRITIS OF THE KNEE: TRIAL OF OR CONTRAINDICATION TO A FORMULARY VERSION OF DICLOFENAC SODIUM 1.5% TOPICAL DROPS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DICLOFENAC-FLECTOR

Products Affected

- *diclofenac epolamine*

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

DIMETHYL FUMARATE

Products Affected

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

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

DIROXIMEL FUMARATE

Products Affected

- VUMERITY

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

DORDAVIPRONE

Products Affected

- MODEYSO

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

DOSTARLIMAB-GXLY

Products Affected

- JEMPERLI

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

DRONABINOL CAPSULE

Products Affected

- *dronabinol oral capsule*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	NAUSEA AND VOMITING ASSOCIATED WITH CANCER CHEMOTHERAPY: TRIAL OF OR CONTRAINDICATION TO ONE ANTIEMETIC THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DROXIDOPA

Products Affected

- *droxidopa*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	NEUROGENIC ORTHOSTATIC HYPOTENSION (NOH): INITIAL: 1) BASELINE BLOOD PRESSURE READINGS WHILE THE PATIENT IS SITTING AND ALSO WITHIN 3 MINUTES OF STANDING FROM A SUPINE POSITION. 2) 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.
Age Restrictions	
Prescriber Restrictions	NOH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR CARDIOLOGIST.
Coverage Duration	INITIAL: 3 MONTHS RENEWAL: 12 MONTHS
Other Criteria	NOH: RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

DUPILUMAB

Products Affected

- DUPIXENT PEN
- DUPIXENT SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: EOSINOPHILIC ASTHMA: BLOOD EOSINOPHIL LEVEL OF 150 TO 1500 CELLS/MCL WITHIN THE PAST 12 MONTHS. EOSINOPHILIC ESOPHAGITIS (EOE): DIAGNOSIS CONFIRMED BY ESOPHAGOGASTRODUODENOSCOPY (EGD) WITH BIOPSY.
Age Restrictions	
Prescriber Restrictions	INITIAL: AD, PN, CSU: PRESCRIBED OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED OR IN CONSULTATION WITH ALLERGIST OR PULMONOLOGIST. CRSWNP: PRESCRIBED OR IN CONSULTATION WITH OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. EOE: PRESCRIBED OR IN CONSULTATION WITH GASTROENTEROLOGIST, ALLERGIST, OR IMMUNOLOGIST. COPD: PRESCRIBED OR IN CONSULTATION WITH PULMONOLOGIST. RENEWAL: CSU: PRESCRIBED OR IN CONSULTATION WITH ALLERGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
Coverage Duration	BP, AFRS: 12 MO. AD/CRSWNP/EOE/PN/CSU: INITIAL/RENEWAL: 6 MO/12 MO. ASTHMA/COPD: INIT/RENEW: 12 MO.

PA Criteria	Criteria Details
Other Criteria	INITIAL: AD: 1) INTRACTABLE PRURITUS OR CRACKING/OOZING/BLEEDING OF AFFECTED SKIN, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE TOPICAL (CORTICOSTEROID, CALCINEURIN INHIBITOR, PDE4 INHIBITOR, OR JAK INHIBITOR). ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY-TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND ONE OTHER MAINTENANCE MEDICATION, AND 2) ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING AT LEAST 3 DAYS WITHIN THE PAST 12 MONTHS, OR ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: (A) DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, (B) ANY NIGHT WAKING DUE TO ASTHMA, (C) SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, (D) ANY ACTIVITY LIMITATION DUE TO ASTHMA. CHRONIC RHINOSINUSITIS WITH NASAL POLYPS (CRSWNP): 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, 2) EVIDENCE OF NASAL POLYPS BY DIRECT EXAMINATION, ENDOSCOPY, OR SINUS CT SCAN, AND 3) INADEQUATELY CONTROLLED DISEASE. PRURIGO NODULARIS (PN): CHRONIC PRURITUS (ITCH MORE THAN 6 WEEKS), MULTIPLE PRURIGINOUS LESIONS, AND HISTORY OR SIGN OF A PROLONGED SCRATCHING BEHAVIOR. EOSINOPHILIC COPD: USED IN COMBINATION WITH A LAMA/LABA/ICS. CHRONIC SPONTANEOUS URTICARIA (CSU): 1) TRIAL OF AND MAINTAINED ON, OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE AND 2) STILL EXPERIENCES HIVES OR ANGIOEDEMA MOST DAYS OF THE WEEK FOR AT LEAST 6 WEEKS. INITIAL/RENEWAL : ALL INDICATIONS EXCEPT BULLOUS PEMPHIGOID (BP): NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4

PA Criteria	Criteria Details
	INHIBITOR) FOR THE SAME INDICATION. RENEWAL: AD, CRSWNP, EOE: IMPROVEMENT WHILE ON THERAPY. ASTHMA: 1) CONTINUED USE OF ICS AND ONE OTHER MAINTENANCE MEDICATION, AND 2) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE, OR (D) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS. PN: IMPROVEMENT OR REDUCTION OF PRURITUS OR PRURIGINOUS LESIONS. EOSINOPHILIC COPD: 1) USED IN COMBINATION WITH A LAMA/LABA/ICS, AND 2) CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: (A) REDUCTION IN COPD EXACERBATIONS FROM BASELINE, OR (B) REDUCTION IN SEVERITY OR FREQUENCY OF COPD-RELATED SYMPTOMS. CSU: MAINTAINED ON OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

DUVELISIB

Products Affected

- COPIKTRA

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

EFLORNITHINE

Products Affected

- IWILFIN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ELACESTRANT

Products Affected

- ORSERDU ORAL TABLET 345 MG, 86 MG

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

ELAGOLIX

Products Affected

- ORILISSA ORAL TABLET 150 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
Age Restrictions	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: 18 YEARS OF AGE OR OLDER.
Prescriber Restrictions	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS
Other Criteria	MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS: INITIAL: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, AND 2) TRIAL OF OR CONTRAINDICATION TO AN NSAID AND A PROGESTIN-CONTAINING PREPARATION. RENEWAL: 1) IMPROVEMENT IN PAIN ASSOCIATED WITH ENDOMETRIOSIS WHILE ON THERAPY, AND 2) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ELAPEGADEMASE-LVLR

Products Affected

- REVCOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	ADENOSINE DEAMINASE SEVERE COMBINED IMMUNE DEFICIENCY (ADA-SCID): INITIAL: ADA-SCID AS MANIFESTED BY: 1) CONFIRMATORY GENETIC TEST, OR 2) SUGGESTIVE LABORATORY FINDINGS (E.G., ELEVATED DEOXYADENOSINE NUCLEOTIDE [DAXP] LEVELS, LYMPHOPENIA) AND HALLMARK SIGNS/SYMPTOMS (E.G., RECURRENT INFECTIONS, FAILURE TO THRIVE, PERSISTENT DIARRHEA).
Age Restrictions	
Prescriber Restrictions	ADA-SCID: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH IMMUNOLOGIST, HEMATOLOGIST/ONCOLOGIST, OR PHYSICIAN SPECIALIZING IN INHERITED METABOLIC DISORDERS.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	ADA-SCID: RENEWAL: 1) IMPROVEMENT OR MAINTENANCE OF IMMUNE FUNCTION FROM BASELINE, AND 2) HAS NOT RECEIVED SUCCESSFUL HCT OR GENE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ELEXACFTOR-TEZACFTOR-IVACFTOR

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CYSTIC FIBROSIS (CF): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME.
Other Criteria	CF: INITIAL: NO CONCURRENT USE WITH ANOTHER CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) MODULATOR. RENEWAL: 1) IMPROVEMENT IN CLINICAL STATUS, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ELRANATAMAB-BCMM

Products Affected

- ELREXFIO 44 MG/1.1 ML VIAL INNER, SUV, P/F
- ELREXFIO SUBCUTANEOUS SOLUTION 40 MG/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	RELAPSED OR REFRACTORY MULTIPLE MYELOMA: RENEWAL: 1) HAS RECEIVED AT LEAST 24 WEEKS OF TREATMENT WITH ELREXFIO, AND 2) HAS RESPONDED TO TREATMENT (PARTIAL RESPONSE OR BETTER), AND HAS MAINTAINED THIS RESPONSE FOR AT LEAST 2 MONTHS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ELTROMBOPAG - ALVAIZ

Products Affected

- ALVAIZ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PERSISTENT OR CHRONIC IMMUNE THROMBOCYTOPENIA (ITP): INITIAL: 1) PLATELET COUNT IS LESS THAN $30 \times 10^9/L$, OR 2) PLATELET COUNT IS LESS THAN $50 \times 10^9/L$ AND HAD A PRIOR BLEEDING EVENT.
Age Restrictions	
Prescriber Restrictions	INITIAL: ITP: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
Coverage Duration	ITP: INITIAL: 6 MO, RENEWAL: 12 MO. HEPATITIS C, SEVERE APLASTIC ANEMIA: 12 MO.
Other Criteria	INITIAL: ITP: 1) TRIAL OF OR CONTRAINDICATION TO ONE CORTICOSTEROID OR IMMUNOGLOBULIN, OR AN INSUFFICIENT RESPONSE TO SPLENECTOMY, AND 2) NO CONCURRENT USE WITH OTHER THROMBOPOIETIN RECEPTOR AGONISTS (TPO-RAS). RENEWAL: ITP: 1) IMPROVEMENT IN PLATELET COUNT FROM BASELINE OR REDUCTION IN BLEEDING EVENTS, AND 2) NO CONCURRENT USE WITH OTHER TPO-RAS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ELTROMBOPAG - PROMACTA

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PERSISTENT OR CHRONIC IMMUNE THROMBOCYTOPENIA (ITP): INITIAL: 1) PLATELET COUNT OF LESS THAN $30 \times 10^9/L$, OR 2) PLATELET COUNT OF LESS THAN $50 \times 10^9/L$ AND A PRIOR BLEEDING EVENT.
Age Restrictions	
Prescriber Restrictions	INITIAL: ITP: PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST OR IMMUNOLOGIST.
Coverage Duration	ITP: INITIAL: 6 MO, RENEWAL: 12 MO. HEPATITIS C, SEVERE APLASTIC ANEMIA: 12 MO.
Other Criteria	INITIAL: ITP: 1) TRIAL OF OR CONTRAINDICATION TO ONE CORTICOSTEROID OR IMMUNOGLOBULIN, OR HAD AN INSUFFICIENT RESPONSE TO SPLENECTOMY, AND 2) NO CONCURRENT USE WITH OTHER THROMBOPOIETIN RECEPTOR AGONISTS (TPO-RAS). ALL INDICATIONS: ELTROMBOPAG ORAL SUSPENSION PACKETS: TRIAL OF A FORMULARY VERSION OF ELTROMBOPAG TABLET OR PATIENT IS UNABLE TO TOLERATE TABLET FORMULATION. RENEWAL: ITP: 1) IMPROVEMENT IN PLATELET COUNTS FROM BASELINE OR REDUCTION IN BLEEDING EVENTS, AND 2) NO CONCURRENT USE WITH OTHER TPO-RAS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ENASIDENIB

Products Affected

- IDHIFA

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

ENCORAFENIB

Products Affected

- BRAFTOVI

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

ENSARTINIB

Products Affected

- ENSACOVE ORAL CAPSULE 100 MG, 25 MG

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

ENTRECTINIB CAPSULES

Products Affected

- ROZLYTREK ORAL CAPSULE 100 MG, 200 MG

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

ENTRECTINIB PELLETS

Products Affected

- ROZLYTREK ORAL PELLETS IN PACKET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), SOLID TUMORS: 1) TRIAL OF OR CONTRAINDICATION TO ROZLYTREK CAPSULES MADE INTO AN ORAL SUSPENSION, AND 2) DIFFICULTY OR UNABLE TO SWALLOW CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ENZALUTAMIDE

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC): HIGH RISK FOR METASTASIS (I.E. PSA DOUBLING TIME OF 9 MONTHS OR LESS). METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC), NON-METASTATIC CRPC (NMCRPC), METASTATIC CSPC (MCSPC), NMCSPC: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

EPCORITAMAB-BYSP

Products Affected

- EPKINLY

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

EPOETIN ALFA-EPBX

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CHRONIC KIDNEY DISEASE (CKD), ANEMIA RELATED TO ZIDOVUDINE, OR CANCER CHEMOTHERAPY: HEMOGLOBIN LEVEL IS LESS THAN 10G/DL. ELECTIVE, NON-CARDIAC, NON-VASCULAR SURGERY: HEMOGLOBIN LEVEL IS 13G/DL OR LESS. RENEWAL: 1) CKD IN ADULTS NOT ON DIALYSIS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS REACHED 10G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 2) CKD IN PEDIATRIC PATIENTS: (A) HEMOGLOBIN LEVEL IS LESS THAN 10G/DL, OR (B) HEMOGLOBIN LEVEL HAS APPROACHED OR EXCEEDS 12G/DL AND THE DOSE IS BEING OR HAS BEEN REDUCED/INTERRUPTED TO DECREASE THE NEED FOR BLOOD TRANSFUSIONS. 3) ANEMIA RELATED TO ZIDOVUDINE: HEMOGLOBIN LEVEL BETWEEN 10G/DL AND 12G/DL. 4) CANCER CHEMOTHERAPY: (A) HEMOGLOBIN LEVEL IS LESS THAN 10 G/DL, OR (B) HEMOGLOBIN LEVEL DOES NOT EXCEED A LEVEL NEEDED TO AVOID RBC TRANSFUSION.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	ANEMIA FROM CHEMO/CKD WITHOUT DIALYSIS/ZIDOVUDINE: INITIAL/RENEWAL: 12 MONTHS. SURGERY: 1 MONTH.
Other Criteria	RENEWAL: CKD: NOT RECEIVING DIALYSIS TREATMENT. THIS DRUG MAY BE EITHER BUNDLED WITH AND COVERED UNDER END STAGE RENAL DISEASE DIALYSIS RELATED SERVICES OR COVERED UNDER MEDICARE D DEPENDING UPON THE CIRCUMSTANCES.

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

ERDAFITINIB

Products Affected

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

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

ERENUMAB-AOOE

Products Affected

- AIMOVIG AUTOINJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	MIGRAINE PREVENTION: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ERLOTINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ESKETAMINE

Products Affected

- SPRAVATO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: TREATMENT-RESISTANT DEPRESSION (TRD), MAJOR DEPRESSIVE DISORDER (MDD): PRESCRIBED BY OR IN CONSULTATION WITH A PSYCHIATRIST OR OTHER REMS-CERTIFIED PROVIDER.
Coverage Duration	INITIAL: TRD: 3 MONTHS. MDD: 4 WEEKS. RENEWAL: TRD, MDD: 12 MONTHS.
Other Criteria	INITIAL: TRD, MDD: 1) NON-PSYCHOTIC, UNIPOLAR DEPRESSION, AND 2) NO ACTIVE SUBSTANCE ABUSE. RENEWAL: TRD, MDD: DEMONSTRATED CLINICAL BENEFIT (IMPROVEMENT IN DEPRESSION) COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ETANERCEPT

Products Affected

- ENBREL MINI
- ENBREL SUBCUTANEOUS SOLUTION
- ENBREL SUBCUTANEOUS SYRINGE
- ENBREL SURECLICK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA), ANKYLOSING SPONDYLITIS (AS): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: ALL INDICATIONS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

EVEROLIMUS-AFINITOR

Products Affected

- *everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *torpenz oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *yulithira oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*

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

EVEROLIMUS-AFINITOR DISPERZ

Products Affected

- *everolimus (antineoplastic) oral tablet for suspension*

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

FECAL MICROBIOTA CAPSULE

Products Affected

- VOWST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	30 DAYS
Other Criteria	CLOSTRIDIoidES DIFFICILE INFECTION (CDI): 1) HAS NOT PREVIOUSLY RECEIVED VOWST: COMPLETION OF ANTIBIOTIC TREATMENT FOR RECURRENT CDI (AT LEAST 3 CDI EPISODES), OR 2) PREVIOUSLY RECEIVED VOWST: (A) TREATMENT FAILURE (DEFINED AS THE PRESENCE OF CDI DIARRHEA WITHIN 8 WEEKS OF FIRST DOSE OF VOWST AND A POSITIVE STOOL TEST FOR C. DIFFICILE), AND (B) HAS NOT RECEIVED MORE THAN ONE TREATMENT COURSE OF VOWST WHICH WAS AT LEAST 12 DAYS AND NOT MORE THAN 8 WEEKS PRIOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

FEDRATINIB

Products Affected

- INREBIC

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

FENFLURAMINE

Products Affected

- FINTEPLA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: DRAVET SYNDROME, LENNOX-GASTAUT SYNDROME (LGS); PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: LGS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING ANTIEPILEPTIC MEDICATIONS: RUFINAMIDE, FELBAMATE, CLOBAZAM, TOPIRAMATE, LAMOTRIGINE, CLONAZEPAM.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

FENTANYL CITRATE

Products Affected

- *fentanyl citrate buccal lozenge on a handle*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CANCER RELATED PAIN: 1) CURRENTLY ON A MAINTENANCE DOSE OF CONTROLLED-RELEASE OPIOID PAIN MEDICATION, AND 2) TRIAL OF OR CONTRAINDICATION TO AT LEAST ONE IMMEDIATE-RELEASE ORAL OPIOID PAIN AGENT OR PATIENT HAS DIFFICULTY SWALLOWING TABLETS/CAPSULES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

FEZOLINETANT

Products Affected

- VEOZAH

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	MENOPAUSAL VASOMOTOR SYMPTOMS (VMS): INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO HORMONAL THERAPY (E.G., ESTRADIOL TRANSDERMAL PATCH, ORAL CONJUGATED ESTROGENS), 2) LABORATORY TESTING TO ESTABLISH BASELINE HEPATIC FUNCTION AND CONTINUED MONITORING OF THESE VALUES IN ACCORDANCE WITH THE FDA CURRENT LABEL RECOMMENDATION, AND 3) NO CONCURRENT USE WITH ANOTHER HORMONAL (E.G., PREMPRO) OR NON-HORMONAL (E.G., BRISDELLE) AGENT FOR VMS. RENEWAL: 1) CONTINUED NEED FOR VMS TREATMENT (PERSISTENT HOT FLASHES), 2) REDUCTION IN VMS FREQUENCY OR SEVERITY DUE TO VEOZAH TREATMENT, AND 3) NO NEW SYMPTOMS OF LIVER INJURY AND/OR WORSENING LAB VALUES (E.G., ALT, AST, TOTAL BILIRUBIN).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

FILGRASTIM-AAFI

Products Affected

- NIVESTYM

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

FINERENONE

Products Affected

- KERENDIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: HEART FAILURE (HF): 1) NEW YORK HEART ASSOCIATION (NYHA) CLASS II-IV, AND 2) LEFT VENTRICULAR EJECTION FRACTION OF AT LEAST 40 PERCENT NOT DUE TO AN UNDERLYING CAUSE (E.G., INFILTRATIVE CARDIOMYOPATHY, HYPERTROPHIC CARDIOMYOPATHY, VALVULAR DISEASE, PERICARDIAL DISEASE, HIGH-OUTPUT HEART FAILURE).
Age Restrictions	
Prescriber Restrictions	INITIAL: HF: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST.
Coverage Duration	INITIAL/RENEWAL:12 MONTHS
Other Criteria	CHRONIC KIDNEY DISEASE (CKD) ASSOCIATED WITH TYPE 2 DIABETES (T2D): INITIAL: HISTORY OF AND WILL CONTINUE ON, HAS A CONTRAINDICATION, OR INTOLERANCE TO AN ANGIOTENSIN CONVERTING ENZYME INHIBITOR (ACE-I) OR AN ANGIOTENSIN RECEPTOR BLOCKER (ARB). HF: INITIAL/RENEWAL: NO CONCURRENT USE WITH ANOTHER MINERALOCORTICOID (ALDOSTERONE) RECEPTOR ANTAGONIST.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

FINGOLIMOD

Products Affected

- *fingolimod*

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

FOSCARBIDOPA-FOSLEVODOPA

Products Affected

- VYALEV

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PARKINSONS DISEASE (PD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	PD: INITIAL: ONE OF THE FOLLOWING: 1) UNABLE TO SWALLOW EXTENDED-RELEASE (ER) TABLETS OR ADMINISTER ER CAPSULES VIA A FEEDING TUBE, OR 2) FAILURE TO ADHERE OR TOLERATE VIA A FEEDING TUBE AN ORAL CARBIDOPA/LEVODOPA REGIMEN. RENEWAL: IMPROVEMENT IN MOTOR SYMPTOMS WHILE ON THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

FRUQUINTINIB

Products Affected

- FRUZAQLA ORAL CAPSULE 1 MG, 5 MG

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

FUTIBATINIB

Products Affected

- LYTGOBI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INTRAHEPATIC CHOLANGIOCARCINOMA (ICCA): COMPLETE A COMPREHENSIVE OPHTHALMOLOGICAL EXAMINATION, INCLUDING OPTICAL COHERENCE TOMOGRAPHY (OCT), PRIOR TO THE INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

GALCANEZUMAB-GNLM

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: MIGRAINE PREVENTION: 6 MOS. EPISODIC CLUSTER HEADACHE: 3 MOS. RENEWAL (ALL): 12 MOS.
Other Criteria	MIGRAINE PREVENTION: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY. EPISODIC CLUSTER HEADACHE: RENEWAL: IMPROVEMENT IN EPISODIC CLUSTER HEADACHE FREQUENCY AS COMPARED TO BASELINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

GANAXOLONE

Products Affected

- ZTALMY

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

GEFITINIB

Products Affected

- *gefitinib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EGFR MUTATION: NOT ON CONCURRENT THERAPY WITH AN EGFR TYROSINE KINASE INHIBITOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

GEPIRONE

Products Affected

- EXXUA ORAL TABLET EXTENDED RELEASE 24 HR
- EXXUA ORAL TABLET, EXT REL 24HR DOSE PACK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	MAJOR DEPRESSIVE DISORDER: INITIAL: TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENTS: TRINTELLIX AND ONE GENERIC ANTIDEPRESSANT. INITIAL/RENEWAL: NO CONCURRENT USE WITH ANOTHER 5-HT1A RECEPTOR AGONIST (E.G., BUSPIRONE). RENEWAL: RESPONSE TO OR REMISSION OF DEPRESSIVE SYMPTOMS WITH THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

GILTERITINIB

Products Affected

- XOSPATA

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

GLASDEGIB

Products Affected

- DAURISMO ORAL TABLET 100 MG, 25 MG

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

GLATIRAMER

Products Affected

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

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

GLP1-DULAGLUTIDE

Products Affected

- TRULICITY

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

GLP1-SEMAGLUTIDE

Products Affected

- OZEMPIC ORAL
- OZEMPIC SUBCUTANEOUS
- RYBELSUS

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

GLP1-TIRZEPATIDE

Products Affected

- MOUNJARO

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

GOSERELIN

Products Affected

- ZOLADEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	ENDOMETRIOSIS: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
Age Restrictions	
Prescriber Restrictions	ENDOMETRIOSIS: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
Coverage Duration	STAGE B2-C PROSTATIC CARCINOMA: 4 MOS. ENDOMETRIOSIS: 6 MOS PER LIFETIME. ALL OTHERS: 12 MONTHS.
Other Criteria	ENDOMETRIOSIS: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, 2) TRIAL OF OR CONTRAINDICATION TO NSAID AND PROGESTIN-CONTAINING PREPARATION, AND 3) HAS NOT RECEIVED A TOTAL OF 6 MONTHS OF TREATMENT PER LIFETIME.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

GUSELKUMAB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ULCERATIVE COLITIS (UC), CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSO, PSA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

HIGH CONCENTRATION ORAL OPIOID SOLUTIONS

Products Affected

- *morphine concentrate oral solution*

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

HIGH RISK DRUGS IN THE ELDERLY - BUTALBITAL-CONTAINING AGENTS

Products Affected

- *butalbital-acetaminop-caf-cod oral capsule 50-325-40-30 mg*
- *butalbital-acetaminophen-caff oral capsule*
- *butalbital-acetaminophen-caff oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS ARE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY - DIPYRIDAMOLE

Products Affected

- *dipyridamole oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY - ESTRADIOL-NORETHINDRONE

Products Affected

- *abigale*
- *estradiol-norethindrone acet*
- *mimvey*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	VULVAR/VAGINAL ATROPHY, OSTEOPOROSIS, AND VASOMOTOR SYMPTOMS OF MENOPAUSE: PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HYPOESTROGENISM TREATMENT AND HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY - ESTROGEN-MEDROXYPROGESTERONE

Products Affected

- PREMPHASE
- PREMPRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY - GLYBURIDE FORMULATIONS

Products Affected

- *glyburide*
- *glyburide micronized*
- *glyburide-metformin*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	TYPE 2 DIABETES MELLITUS (DM): 1) TRIAL OF OR CONTRAINDICATION TO GLIMEPIRIDE OR GLIPIZIDE, OR 2) PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS ARE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

HIGH RISK DRUGS IN THE ELDERLY - KETOROLAC

Products Affected

- *ketorolac oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	30 DAYS
Other Criteria	PREScriBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY - PHENOBARBITAL

Products Affected

- *phenobarbital*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	EPILEPSY/SEIZURES: PATIENTS WHO ARE NEWLY PRESCRIBED PHENOBARBITAL: 1) HAS NOT RESPONDED TO AT LEAST ONE ANTICONVULSANT, OR 2) PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

HIGH RISK DRUGS IN THE ELDERLY - PROMETHAZINE

Products Affected

- *promethazine injection solution 25 mg/ml* *mg*
- *promethazine oral tablet*
- *promethazine rectal suppository 25 mg*
- *promethegan rectal suppository 12.5 mg, 25*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRURITUS/URTICARIA/SEASONAL/PERENNIAL ALLERGY: 1) TRIAL OF OR CONTRAINDICATION TO A NON-SEDATING ANTIHISTAMINE SUCH AS LEVOCETIRIZINE, OR 2) PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. NAUSEA AND VOMITING: PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH-RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS REQUIRE PHYSICIAN ATTESTATION THAT REQUESTED MEDICATION IS USED TO TREAT A DIAGNOSIS UNRELATED TO THE TERMINAL ILLNESS OR RELATED CONDITION, AND ARE APPROVED WITHOUT TRIAL OF FORMULARY ALTERNATIVES NOR REQUIRING PRESCRIBER ACKNOWLEDGEMENT.
Indications	All FDA-approved Indications.
Off Label Uses	

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

HIGH RISK DRUGS IN THE ELDERLY - SCOPOLAMINE

Products Affected

- *scopolamine base*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PREScriBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS REQUIRE PHYSICIAN ATTESTATION THAT REQUESTED MEDICATION IS USED TO TREAT A DIAGNOSIS UNRELATED TO THE TERMINAL ILLNESS OR RELATED CONDITION, AND ARE APPROVED WITHOUT REQUIRING PRESCRIBER ACKNOWLEDGEMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY- DIPHENOXYLATE-ATROPINE

Products Affected

- *diphenoxylate-atropine oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY- INDOMETHACIN

Products Affected

- *indomethacin oral capsule*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY- MEGESTROL

Products Affected

- *megestrol oral suspension 400 mg/10 ml (40 mg/ml), 625 mg/5 ml (125 mg/ml)*
- *megestrol oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

HIGH RISK DRUGS IN THE ELDERLY- PAROXETINE

Products Affected

- *paroxetine hcl*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	APPROPRIATE HIGH RISK MEDICATION USE: 65 YEARS OF AGE OR OLDER. PA NOT REQUIRED FOR AGE 0-64 YEARS.
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	PRESCRIBER ACKNOWLEDGEMENT/AWARENESS THAT THE DRUG IS CONSIDERED HIGH RISK FOR PATIENTS 65 YEARS AND OLDER. HOSPICE PATIENTS WILL BE APPROVED WITHOUT REQUIRING ADDITIONAL CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

IBRUTINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CHRONIC GRAFT VS HOST DISEASE (CGVHD): NO CONCURRENT USE WITH JAKAFI, NIKTIMVO, OR REZUROCK.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ICATIBANT

Products Affected

- *icatibant*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HEREDITARY ANGIOEDEMA (HAE): INITIAL: 1) TYPE III HAE, OR 2) TYPE I OR II HAE CONFIRMED BY ONE OF THE FOLLOWING COMPLEMENT TESTS: C1-INH PROTEIN LEVELS, C4 PROTEIN LEVELS, C1-INH FUNCTIONAL LEVELS, C1Q.
Age Restrictions	
Prescriber Restrictions	HAE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST, IMMUNOLOGIST, HEMATOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	HAE: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER MEDICATIONS FOR THE TREATMENT OF ACUTE HAE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

IDEALALISIB

Products Affected

- ZYDELIG

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

IMATINIB

Products Affected

- *imatinib oral tablet 100 mg, 400 mg*

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

IMATINIB SOLUTION

Products Affected

- IMKELDI

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

IMETELSTAT

Products Affected

- RYTELO

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

IMLUNESTRANT

Products Affected

- INLURIYO

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

INAVOLISIB

Products Affected

- ITOVEBI ORAL TABLET 3 MG, 9 MG

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

INFLIXIMAB

Products Affected

- *infliximab*

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

PA Criteria	Criteria Details
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. PSA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, XELJANZ, RINVOQ, SKYRIZI, TREMFYA, ORENCIA, OTEZLA. PSO: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, SKYRIZI, TREMFYA, OTEZLA. AS: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: COSENTYX, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ. MODERATE TO SEVERE CD: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS, WHERE AGES ALIGN: SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, RINVOQ, SKYRIZI, TREMFYA. UC: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS, WHERE AGES ALIGN: SELARSDI/YESINTEK/USTEKINUMAB-AAUZ, XELJANZ, HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, RINVOQ, SKYRIZI, TREMFYA. INITIAL/RENEWAL: RA, PSA, AS, PSO, MODERATE TO SEVERE CD, UC: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, PSA, AS, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

INSULIN SUPPLIES PAYMENT DETERMINATION

Products Affected

- 1ST TIER UNIFINE PENTP 5MM 31G
- 1ST TIER UNIFINE PNTIP 4MM 32G
- 1ST TIER UNIFINE PNTIP 6MM 31G
- 1ST TIER UNIFINE PNTIP 8MM 31G STRL,SINGLE-USE,SHRT
- 1ST TIER UNIFINE PNTTP 29GX1/2"
- 1ST TIER UNIFINE PNTTP 31GX3/16
- 1ST TIER UNIFINE PNTTP 32GX5/32
- ADVOCATE INS 0.3 ML 30GX5/16"
- ADVOCATE INS 0.3 ML 31GX5/16"
- ADVOCATE INS 0.5 ML 30GX5/16"
- ADVOCATE INS 0.5 ML 31GX5/16"
- ADVOCATE INS 1 ML 31GX5/16"
- ADVOCATE INS SYR 1 ML 30GX5/16
- ADVOCATE PEN NEEDLE 32G 4MM
- ADVOCATE PEN NEEDLE 4MM 33G
- ADVOCATE PEN NEEDLES 5MM 31G
- ADVOCATE PEN NEEDLES 8MM 31G
- ALCOHOL PADS
- AQINJECT PEN NEEDLE 31G 5MM
- AQINJECT PEN NEEDLE 32G 4MM
- ASSURE ID DUO PRO NDL 31G 5MM
- ASSURE ID DUO-SHIELD 30GX3/16"
- ASSURE ID DUO-SHIELD 30GX5/16"
- ASSURE ID INSULIN SAFETY SYRINGE 1 ML 29 GAUGE X 1/2"
- ASSURE ID PEN NEEDLE 30GX5/16"
- ASSURE ID PRO PEN NDL 30G 5MM
- ASSURE ID SYR 0.5 ML 31GX15/64"
- ASSURE ID SYR 1 ML 31GX15/64"
- BD AUTOSHIELD DUO NDL 5MMX30G
- BD ECLIPSE 30GX1/2" SYRINGE
- BD INS SYR 0.3 ML 8MMX31G(1/2)
- BD INS SYR UF 0.3 ML 12.7MMX30G
- BD INS SYR UF 0.5 ML 12.7MMX30G NOT FOR RETAIL SALE
- BD INSULIN SYR 1 ML 27GX12.7MM
- BD INSULIN SYR 1 ML 27GX5/8" MICRO-FINE
- BD SAFETYGLD INS 0.3 ML 29G 13MM
- BD SAFETYGLD INS 0.3 ML 31G 8MM
- BD SAFETYGLD INS 0.5 ML 30G 8MM
- BD SAFETYGLD INS 1 ML 29G 13MM
- BD SAFETYGLID INS 1 ML 6MMX31G
- BD SAFETYGLIDE SYRINGE 27GX5/8
- BD SAFTYGLD INS 0.3 ML 6MMX31G
- BD SAFTYGLD INS 0.5 ML 29G 13MM
- BD SAFTYGLD INS 0.5 ML 6MMX31G
- BD SINGLE USE SWAB
- BD UF MICRO PEN NEEDLE 6MMX32G
- BD UF MINI PEN NEEDLE 5MMX31G
- BD UF NANO PEN NEEDLE 4MMX32G
- BD UF ORIG PEN NDL 12.7MMX29G
- BD UF SHORT PEN NEEDLE 8MMX31G
- BD VEO INS 0.3 ML 6MMX31G (1/2)
- BD VEO INS SYRING 1 ML 6MMX31G
- BD VEO INS SYRN 0.3 ML 6MMX31G
- BD VEO INS SYRN 0.5 ML 6MMX31G
- BORDERED GAUZE 2"X2"
- CAREFINE PEN NEEDLE 12.7MM 29G HRI
- CAREFINE PEN NEEDLE 4MM 32G
- CAREFINE PEN NEEDLE 5MM 32G
- CAREFINE PEN NEEDLE 6MM 31G HRI
- CAREFINE PEN NEEDLE 8MM 30G HRI
- CAREFINE PEN NEEDLES 6MM 32G
- CAREFINE PEN NEEDLES 8MM 31G
- CARETOUCH ALCOHOL 70% PREP PAD
- CARETOUCH PEN NEEDLE 29G 12MM
- CARETOUCH PEN NEEDLE 31GX1/4"
- CARETOUCH PEN NEEDLE 31GX3/16"
- CARETOUCH PEN NEEDLE 31GX5/16"
- CARETOUCH PEN NEEDLE 32GX3/16"
- CARETOUCH PEN NEEDLE 32GX5/32"
- CARETOUCH SYR 0.3 ML 31GX5/16"
- CARETOUCH SYR 0.5 ML 30GX5/16"
- CARETOUCH SYR 0.5 ML 31GX5/16"
- CARETOUCH SYR 1 ML 28GX5/16"
- CARETOUCH SYR 1 ML 29GX5/16"
- CARETOUCH SYR 1 ML 30GX5/16"
- CARETOUCH SYR 1 ML 31GX5/16"
- CLICKFINE PEN NEEDLE 32GX5/32"

- 32GX4MM, STERILE
- CLICKFINE UNIVERSAL 31G X 1/4" 6MM, STORE BRAND
- COMFORT EZ 0.3 ML 31G 15/64"
- COMFORT EZ 0.5 ML 31G 15/64"
- COMFORT EZ INS 0.3 ML 30GX1/2"
- COMFORT EZ INS 0.3 ML 30GX5/16"
- COMFORT EZ INS 1 ML 31G 15/64"
- COMFORT EZ INS 1 ML 31GX5/16"
- COMFORT EZ INSULIN SYR 0.3 ML
- COMFORT EZ INSULIN SYR 0.5 ML
- COMFORT EZ PEN NEEDLE 12MM 29G
- COMFORT EZ PEN NEEDLES 4MM 32G SINGLE USE, MICRO
- COMFORT EZ PEN NEEDLES 4MM 33G
- COMFORT EZ PEN NEEDLES 5MM 31G MINI
- COMFORT EZ PEN NEEDLES 5MM 32G SINGLE USE,MINI,HRI
- COMFORT EZ PEN NEEDLES 5MM 33G
- COMFORT EZ PEN NEEDLES 6MM 31G
- COMFORT EZ PEN NEEDLES 6MM 32G
- COMFORT EZ PEN NEEDLES 6MM 33G
- COMFORT EZ PEN NEEDLES 8MM 31G SHORT
- COMFORT EZ PEN NEEDLES 8MM 32G
- COMFORT EZ PEN NEEDLES 8MM 33G
- COMFORT EZ PRO PEN NDL 30G 8MM
- COMFORT EZ PRO PEN NDL 31G 4MM
- COMFORT EZ PRO PEN NDL 31G 5MM
- COMFORT EZ SYR 0.3 ML 29GX1/2"
- COMFORT EZ SYR 0.5 ML 28GX1/2"
- COMFORT EZ SYR 0.5 ML 29GX1/2"
- COMFORT EZ SYR 0.5 ML 30GX1/2"
- COMFORT EZ SYR 1 ML 27G 12.7MM
- COMFORT EZ SYR 1 ML 28GX1/2"
- COMFORT EZ SYR 1 ML 29GX1/2"
- COMFORT EZ SYR 1 ML 30GX1/2"
- COMFORT EZ SYR 1 ML 30GX5/16"
- COMFORT POINT PEN NDL 31GX1/3"
- COMFORT POINT PEN NDL 31GX1/6"
- COMFORT TOUCH PEN NDL 31G 4MM
- COMFORT TOUCH PEN NDL 31G 5MM
- COMFORT TOUCH PEN NDL 31G 6MM
- COMFORT TOUCH PEN NDL 31G 8MM
- COMFORT TOUCH PEN NDL 32G 4MM
- COMFORT TOUCH PEN NDL 32G 5MM
- COMFORT TOUCH PEN NDL 32G 6MM
- COMFORT TOUCH PEN NDL 32G 8MM
- COMFORT TOUCH PEN NDL 33G 4MM
- COMFORT TOUCH PEN NDL 33G 6MM
- COMFORT TOUCH PEN NDL 33GX5MM
- CURITY ALCOHOL PREPS 2 PLY,MEDIUM
- CURITY GAUZE PADS
- CURITY GAUZE SPONGES (12 PLY)-200/BAG
- DERMACEA 2"X2" GAUZE 12 PLY, USP TYPE VII
- DERMACEA GAUZE 2"X2" SPONGE 8 PLY
- DERMACEA NON-WOVEN 2"X2" SPNGE
- DROPLET 0.3 ML 29G 12.7MM(1/2) OUTER
- DROPLET 0.3 ML 30G 12.7MM(1/2) OUTER
- DROPLET 0.5 ML 29GX12.5MM(1/2)
- DROPLET 0.5 ML 30GX12.5MM(1/2)
- DROPLET INS 0.3 ML 29GX12.5MM
- DROPLET INS 0.3 ML 30G 8MM(1/2) OUTER
- DROPLET INS 0.3 ML 30GX12.5MM
- DROPLET INS 0.3 ML 31G 6MM(1/2) OUTER
- DROPLET INS 0.3 ML 31G 8MM(1/2) OUTER
- DROPLET INS 0.5 ML 29G 12.7MM OUTER
- DROPLET INS 0.5 ML 30G 12.7MM OUTER
- DROPLET INS 0.5 ML 30GX6MM(1/2)
- DROPLET INS 0.5 ML 30GX8MM(1/2)
- DROPLET INS 0.5 ML 31GX6MM(1/2)
- DROPLET INS 0.5 ML 31GX8MM(1/2)
- DROPLET INS SYR 0.3 ML 30GX6MM
- DROPLET INS SYR 0.3 ML 30GX8MM
- DROPLET INS SYR 0.3 ML 31GX6MM
- DROPLET INS SYR 0.3 ML 31GX8MM
- DROPLET INS SYR 0.5 ML 30G 8MM OUTER
- DROPLET INS SYR 0.5 ML 31G 6MM OUTER
- DROPLET INS SYR 0.5 ML 31G 8MM

- OUTER
- DROPLET INS SYR 1 ML 29G 12.7MM
- OUTER
- DROPLET INS SYR 1 ML 30G 12.5MM
- DROPLET INS SYR 1 ML 30G 6MM
- DROPLET INS SYR 1 ML 30G 8MM
- OUTER
- DROPLET INS SYR 1 ML 31G 6MM
- OUTER
- DROPLET INS SYR 1 ML 31G 8MM
- DROPLET MICRON 34G X 9/64"
- DROPLET PEN NEEDLE 29G 10MM
- DROPLET PEN NEEDLE 29G 12MM
- DROPLET PEN NEEDLE 30G 8MM
- DROPLET PEN NEEDLE 31G 5MM
- DROPLET PEN NEEDLE 31G 6MM
- DROPLET PEN NEEDLE 31G 8MM
- DROPLET PEN NEEDLE 32G 4MM
- DROPLET PEN NEEDLE 32G 5MM
- DROPLET PEN NEEDLE 32G 6MM
- DROPLET PEN NEEDLE 32G 8MM
- DROPSAFE ALCOHOL 70% PREP PADS
- DROPSAFE INS SYR 0.3 ML 31G 6MM
- DROPSAFE INS SYR 0.3 ML 31G 8MM
- DROPSAFE INS SYR 0.5 ML 31G 6MM
- DROPSAFE INS SYR 0.5 ML 31G 8MM
- DROPSAFE INSUL SYR 1 ML 31G 6MM
- DROPSAFE INSUL SYR 1 ML 31G 8MM
- DROPSAFE INSULN 1 ML 29G 12.5MM
- DROPSAFE PEN NEEDLE 31G 4MM
- DROPSAFE PEN NEEDLE 31G 5MM
- DROPSAFE PEN NEEDLE 31G 8MM
- DROPSAFE PEN NEEDLE 31GX1/4"
- EASY CMFT SFTY PEN NDL 31G 5MM
- EASY CMFT SFTY PEN NDL 31G 6MM
- EASY CMFT SFTY PEN NDL 32G 4MM
- EASY COMFORT 0.3 ML 31G 1/2"
- EASY COMFORT 0.3 ML 31G 5/16"
- EASY COMFORT 0.5 ML 30GX1/2"
- EASY COMFORT 0.5 ML 31GX5/16"
- EASY COMFORT 0.5 ML 32GX5/16"
- EASY COMFORT 0.5 ML SYRINGE
- EASY COMFORT 1 ML 31GX5/16"
- EASY COMFORT 1 ML 32GX5/16"
- EASY COMFORT ALCOHOL 70% PAD
- EASY COMFORT INSULIN 1 ML SYR
- EASY COMFORT PEN NDL 29G 4MM
- EASY COMFORT PEN NDL 29G 5MM
- EASY COMFORT PEN NDL 31GX1/4"
- EASY COMFORT PEN NDL 31GX3/16"
- EASY COMFORT PEN NDL 31GX5/16"
- EASY COMFORT PEN NDL 32GX5/32"
- EASY COMFORT PEN NDL 33G 4MM
- EASY COMFORT PEN NDL 33G 5MM
- EASY COMFORT PEN NDL 33G 6MM
- EASY COMFORT SYR 0.5 ML 29G 8MM
- EASY COMFORT SYR 1 ML 29G 8MM
- EASY COMFORT SYR 1 ML 30GX1/2"
- EASY GLIDE INS 0.3 ML 31GX6MM
- EASY GLIDE INS 0.5 ML 31GX6MM
- EASY GLIDE INS 1 ML 31GX6MM
- EASY GLIDE PEN NEEDLE 4MM 33G
- EASY TOUCH 0.3 ML 30G 8MM(1/2)
- EASY TOUCH 0.3 ML 31G 6MM(1/2)
- EASY TOUCH 0.3 ML 31G 8MM(1/2)
- EASY TOUCH 0.3 ML SYR 30GX1/2"
- EASY TOUCH 0.5 ML SYR 27GX1/2"
- EASY TOUCH 0.5 ML SYR 29GX1/2"
- EASY TOUCH 0.5 ML SYR 30GX1/2"
- EASY TOUCH 0.5 ML SYR 30GX5/16
- EASY TOUCH 1 ML SYR 29GX1/2"
- EASY TOUCH 1 ML SYR 30GX1/2"
- EASY TOUCH ALCOHOL 70% PADS
- GAMMA-STERILIZED
- EASY TOUCH AUTO 0.5 ML 30G 6MM
- EASY TOUCH AUTO 0.5 ML 30G 8MM
- EASY TOUCH AUTORET 1 ML 30G 6MM
- EASY TOUCH AUTORET 1 ML 30G 8MM
- EASY TOUCH FLIPLOK 1 ML 27GX0.5
- EASY TOUCH INS 0.3 ML 30G 8MM
- EASY TOUCH INS 0.3 ML 31G 8MM
- EASY TOUCH INS 0.5 ML 30G 8MM
- EASY TOUCH INS 0.5 ML 31G 6MM
- EASY TOUCH INS 0.5 ML 31G 8MM
- EASY TOUCH INS 1 ML 27G 1/2"
- EASY TOUCH INS 1 ML 28G 12.7MM
- EASY TOUCH INS SYR 1 ML 30G 8MM
- EASY TOUCH INS SYR 1 ML 31G 6MM
- EASY TOUCH INS SYR 1 ML 31G 8MM
- EASY TOUCH INSULIN 1 ML 29GX1/2
- EASY TOUCH INSULIN 1 ML 30GX1/2
- EASY TOUCH INSULIN SYR 1 ML
- EASY TOUCH INSULIN SYR 1 ML
- RETRACTABLE

- EASY TOUCH INSULN 1 ML 29GX1/2"
- EASY TOUCH INSULN 1 ML 30GX1/2"
- EASY TOUCH INSULN 1 ML 30GX5/16
- EASY TOUCH INSULN 1 ML 31GX5/16
- EASY TOUCH LUER LOK INSUL 1 ML
- EASY TOUCH PEN NEEDLE 29GX1/2"
- EASY TOUCH PEN NEEDLE 30GX5/16
- EASY TOUCH PEN NEEDLE 31GX1/4"
- EASY TOUCH PEN NEEDLE 31GX3/16
- EASY TOUCH PEN NEEDLE 31GX5/16
- EASY TOUCH PEN NEEDLE 32GX1/4"
- EASY TOUCH PEN NEEDLE 32GX3/16
- EASY TOUCH PEN NEEDLE 32GX5/32
- EASY TOUCH SAF PEN NDL 29G 5MM
- EASY TOUCH SAF PEN NDL 29G 8MM
- EASY TOUCH SAF PEN NDL 30G 5MM
- EASY TOUCH SAF PEN NDL 30G 8MM
- EASY TOUCH SYR 0.5 ML 28G 12.7MM
- EASY TOUCH SYR 0.5 ML 29G 12.7MM
- EASY TOUCH SYR 1 ML 27G 16MM
- EASY TOUCH UNI-SLIP SYR 1 ML
- EASYLIFE ALCOHOL 70% PADS
- EASYLIFE INS PEN NDL 29G 12MM
- EASYLIFE INS PEN NDL 31G 4MM
- EASYLIFE INS PEN NDL 31G 5MM
- EASYLIFE INS PEN NDL 31G 6MM
- EASYLIFE INS PEN NDL 31G 8MM
- EASYLIFE INS PEN NDL 32G 4MM
- EASYLIFE INS PEN NDL 32G 5MM
- EASYLIFE INS PEN NDL 32G 6MM
- EASYLIFE INS PEN NDL 32G 8MM
- EASYLIFE INS PEN NDL 33G 4MM
- EASYLIFE INS PEN NDL 33G 5MM
- EASYLIFE INS PEN NDL 33G 6MM
- EASYLIFE INS PEN NDL 33G 8MM
- EASYLIFE INS SYR 0.5 ML 30G 8MM
- EASYLIFE INS SYR 1 ML 30G 8MM
- EASYLIFE INS SYR 1 ML 31G 8MM
- EASYLIFE SAFTY PEN NDL 31G 4MM
- EASYLIFE SAFTY PEN NDL 31G 5MM
- EASYLIFE SYR 1 ML 30G 12.7MM
- EASYTOUCH SAF PEN NDL 30G 6MM
- EMBRACE PEN NEEDLE 29G 12MM
- EMBRACE PEN NEEDLE 30G 5MM
- EMBRACE PEN NEEDLE 30G 8MM
- EMBRACE PEN NEEDLE 31G 5MM
- EMBRACE PEN NEEDLE 31G 6MM
- EMBRACE PEN NEEDLE 31G 8MM
- EMBRACE PEN NEEDLE 32G 4MM
- EXEL U100 0.3 ML 29GX1/2"
- EXEL U100 0.3 ML 30GX5/16"
- FT STERILE PADS 2" X 2"
- GAUZE PAD TOPICAL BANDAGE 2 X 2"
- GNP ALCOHOL SWAB STERILE, TWO PLY
- GNP PEN NEEDLE 31G 5MM
- GNP PEN NEEDLE 32G 6MM
- GNP SIMPLI PEN NEEDLE 32G 4MM
- GNP ULT C 0.3 ML 29GX1/2" (1/2) 1/2 UNIT
- GNP ULT CMFRT 0.5 ML 29GX1/2"
- GNP ULTR CMFRT 0.5 ML 30GX5/16
- GNP ULTRA COMFORT 1 ML SYRINGE
- GNP ULTRA COMFORT 3/10 ML SYR
- GS PEN NEEDLE 31G X 8MM
- HEALTHWISE INS 0.3 ML 30GX5/16"
- HEALTHWISE INS 0.3 ML 31GX5/16"
- HEALTHWISE INS 0.5 ML 30GX5/16"
- HEALTHWISE INS 0.5 ML 31GX5/16"
- HEALTHWISE INS 1 ML 30GX5/16"
- HEALTHWISE INS 1 ML 31GX5/16"
- HEALTHWISE PEN NEEDLE 31G 5MM
- HEALTHWISE PEN NEEDLE 31G 8MM
- HEALTHWISE PEN NEEDLE 32G 4MM
- HEALTHY ACCENTS PENTIP 5MM 31G
- HEALTHY ACCENTS PENTIP 6MM 31G
- HEALTHY ACCENTS PENTIP 8MM 31G
- HEALTHY ACCENTS PENTIP 12MM 29G
- HEB INCONTROL ALCOHOL 70% PADS
- INCONTROL PEN NEEDLE 12MM 29G
- INCONTROL PEN NEEDLE 4MM 32G
- INCONTROL PEN NEEDLE 5MM 31G
- INCONTROL PEN NEEDLE 6MM 31G
- INCONTROL PEN NEEDLE 8MM 31G
- INSULIN 1 ML SYRINGE
- INSULIN 1/2 ML SYRINGE
- INSULIN 3/10 ML SYRINGE
- INSULIN SYR 0.3 ML 31GX1/4(1/2)
- INSULIN SYRIN 0.5 ML 28GX1/2" (OTC)
- INSULIN SYRIN 0.5 ML 29GX1/2" (OTC)
- INSULIN SYRIN 0.5 ML 30GX1/2" (RX)
- INSULIN SYRIN 0.5 ML 30GX5/16" SHORT NEEDLE (OTC)

- INSULIN SYRINGE 0.5 ML 27G 1/2" INNER
- INSULIN SYRINGE 0.3 ML
- INSULIN SYRINGE 0.3 ML 31GX1/4
- INSULIN SYRINGE 0.5 ML
- INSULIN SYRINGE 0.5 ML 31GX1/4
- INSULIN SYRINGE 1 ML
- INSULIN SYRINGE 1 ML 27G 1/2" INNER
- INSULIN SYRINGE 1 ML 28GX1/2" (OTC)
- INSULIN SYRINGE 1 ML 30GX1/2" SHORT NEEDLE (OTC)
- INSULIN SYRINGE 1 ML 30GX5/16" SHORT NEEDLE (OTC)
- INSULIN SYRINGE 1 ML 31GX1/4"
- INSULIN SYRINGE 1 ML 31GX5/16" SHORT NEEDLE, THIN II (OTC)
- INSULIN SYRINGE-NEEDLE U-100 SYRINGE 0.3 ML 29 GAUGE, 1 ML 29 GAUGE X 1/2", 1/2 ML 28 GAUGE
- INSULIN U-500 SYRINGE-NEEDLE
- INSUPEN PEN NEEDLE 29GX1/2"
- INSUPEN PEN NEEDLE 31G 8MM
- INSUPEN PEN NEEDLE 31GX3/16"
- INSUPEN PEN NEEDLE 32G 4MM
- INSUPEN PEN NEEDLE 32G 6MM (RX)
- IV ANTISEPTIC WIPES
- KENDALL ALCOHOL 70% PREP PAD
- LISCO SPONGES 100/BAG
- LITE TOUCH 31GX1/4" PEN NEEDLE
- LITE TOUCH INSULIN 0.5 ML SYR
- LITE TOUCH INSULIN 1 ML SYR
- LITE TOUCH INSULIN SYR 1 ML
- LITE TOUCH PEN NEEDLE 29G
- LITE TOUCH PEN NEEDLE 31G
- LITETOUCH INS 0.3 ML 29GX1/2"
- LITETOUCH INS 0.3 ML 30GX5/16"
- LITETOUCH INS 0.3 ML 31GX5/16"
- LITETOUCH INS 0.5 ML 31GX5/16"
- LITETOUCH SYR 0.5 ML 28GX1/2"
- LITETOUCH SYR 0.5 ML 29GX1/2"
- LITETOUCH SYR 0.5 ML 30GX5/16"
- LITETOUCH SYRIN 1 ML 28GX1/2"
- LITETOUCH SYRIN 1 ML 29GX1/2"
- LITETOUCH SYRIN 1 ML 30GX5/16"
- MAGELLAN INSUL SYRINGE 0.3 ML
- MAGELLAN INSUL SYRINGE 0.5 ML
- MAGELLAN INSULIN SYR 0.3 ML
- MAGELLAN INSULIN SYR 0.5 ML
- MAGELLAN INSULIN SYRINGE 1 ML
- MAXI-COMFORT INS 0.5 ML 28G
- MAXI-COMFORT INS 1 ML 28GX1/2"
- MAXICOMFORT II PEN NDL 31GX6MM
- MAXICOMFORT INS 0.5 ML 27GX1/2"
- MAXICOMFORT INS 1 ML 27GX1/2"
- MAXICOMFORT PEN NDL 29G X 5MM
- MAXICOMFORT PEN NDL 29G X 8MM
- MICRODOT PEN NEEDLE 31GX6MM
- MICRODOT PEN NEEDLE 32GX4MM
- MICRODOT PEN NEEDLE 33GX4MM
- MICRODOT READYGARD NDL 31G 5MM OUTER
- MINI PEN NEEDLE 32G 5MM
- MINI PEN NEEDLE 32G 8MM
- MINI PEN NEEDLE 33G 5MM
- MINI PEN NEEDLE 33G 6MM
- MINI ULTRA-THIN II PEN NDL 31G STERILE
- MONOJECT 0.5 ML SYRN 28GX1/2"
- MONOJECT 1 ML SYRN 27X1/2"
- MONOJECT 1 ML SYRN 28GX1/2" (OTC)
- MONOJECT INSUL SYR U100 (OTC)
- MONOJECT INSUL SYR U100 .5ML, 29GX1/2" (OTC)
- MONOJECT INSUL SYR U100 0.5 ML CONVERTS TO 29G (OTC)
- MONOJECT INSUL SYR U100 1 ML
- MONOJECT INSUL SYR U100 1 ML 3'S, 29GX1/2" (OTC)
- MONOJECT INSUL SYR U100 1 ML W/O NEEDLE (OTC)
- MONOJECT INSULIN SYR 0.3 ML
- MONOJECT INSULIN SYR 0.3 ML (OTC)
- MONOJECT INSULIN SYR 0.5 ML
- MONOJECT INSULIN SYR 0.5 ML (OTC)
- MONOJECT INSULIN SYR 1 ML 3'S (OTC)
- MONOJECT INSULIN SYR U-100
- MONOJECT SYRINGE 0.3 ML
- MONOJECT SYRINGE 0.5 ML
- MONOJECT SYRINGE 1 ML
- NANO 2 GEN PEN NEEDLE 32G 4MM
- NANO PEN NEEDLE 32G 4MM

- NOVOFINE 32G NEEDLES
- NOVOFINE AUTOCOVER 30G NEEDLE
- NOVOFINE PLUS PEN NDL 32GX1/6"
- PEN NEEDLE 30G 5MM OUTER
- PEN NEEDLE 30G 8MM INNER
- PEN NEEDLE 30G X 5/16"
- PEN NEEDLE 31G X 1/4" HRI
- PEN NEEDLE 33G 4MM
- PEN NEEDLE, DIABETIC NEEDLE 29 GAUGE X 1/2"
- PEN NEEDLES 12MM 29G 29GX12MM,STRL
- PEN NEEDLES 4MM 32G
- PEN NEEDLES 5MM 31G 31GX5MM,STRL,MINI (OTC)
- PEN NEEDLES 8MM 31G 31GX8MM,STRL,SHORT (OTC)
- PENTIPS PEN NEEDLE 29G 1/2"
- PENTIPS PEN NEEDLE 31G 1/4"
- PENTIPS PEN NEEDLE 31GX3/16" MINI, 5MM
- PENTIPS PEN NEEDLE 31GX5/16" SHORT, 8MM
- PENTIPS PEN NEEDLE 32G 1/4"
- PENTIPS PEN NEEDLE 32GX5/32"
- PIP PEN NEEDLE 31G X 5MM
- PIP PEN NEEDLE 32G X 4MM
- PREVENT PEN NEEDLE 31GX1/4"
- PREVENT PEN NEEDLE 31GX5/16"
- PRO COMFORT 0.5 ML 30GX1/2"
- PRO COMFORT 0.5 ML 30GX5/16"
- PRO COMFORT 0.5 ML 31GX5/16"
- PRO COMFORT 1 ML 30GX1/2"
- PRO COMFORT 1 ML 30GX5/16"
- PRO COMFORT 1 ML 31GX5/16"
- PRO COMFORT ALCOHOL 70% PADS
- PRO COMFORT PEN NDL 32G 8MM
- PRO COMFORT PEN NDL 32G X 1/4"
- PRO COMFORT PEN NDL 4MM 32G
- PRO COMFORT PEN NDL 5MM 32G
- PRO-COMFORT ALCOHOL 70% PADS
- PRODIGY INS SYR 1 ML 28GX1/2"
- PRODIGY SYRNG 0.5 ML 31GX5/16"
- PRODIGY SYRNGE 0.3 ML 31GX5/16"
- PURE CMFT SFTY PEN NDL 31G 5MM
- PURE CMFT SFTY PEN NDL 31G 6MM
- PURE CMFT SFTY PEN NDL 32G 4MM
- PURE COMFORT ALCOHOL 70% PADS
- PURE COMFORT PEN NDL 32G 4MM
- PURE COMFORT PEN NDL 32G 5MM
- PURE COMFORT PEN NDL 32G 6MM
- PURE COMFORT PEN NDL 32G 8MM
- RA ISOPROPYL ALCOHOL 70% WIPES
- RA STERILE PADS 2"X 2"
- RAYA SURE PEN NEEDLE 29G 12MM
- RAYA SURE PEN NEEDLE 31G 4MM
- RAYA SURE PEN NEEDLE 31G 5MM
- RAYA SURE PEN NEEDLE 31G 6MM
- RELION INS SYR 0.3 ML 31GX6MM
- RELION INS SYR 0.5 ML 31GX6MM
- RELION INS SYR 1 ML 31GX15/64"
- RELION PEN NEEDLE 32GX5/32"
- RELION SYRING 0.3 ML 31GX5/16" INNER
- RELION SYRING 0.5 ML 31GX5/16" INNER (OTC)
- SAFESNAP INS SYR UNITS-100 0.3 ML 30GX5/16",10X10
- SAFESNAP INS SYR UNITS-100 0.5 ML 29GX1/2",10X10
- SAFESNAP INS SYR UNITS-100 0.5 ML 30GX5/16",10X10
- SAFESNAP INS SYR UNITS-100 1 ML 28GX1/2",10X10
- SAFESNAP INS SYR UNITS-100 1 ML 29GX1/2",10X10
- SAFETY PEN NEEDLE 31G 4MM
- SAFETY PEN NEEDLE 5MM X 31G
- SAFETY SYRINGE 0.5 ML 30G 1/2"
- SECURESAFE PEN NDL 30GX5/16" OUTER
- SECURESAFE PEN NEEDLE 31G 5MM OUTER
- SECURESAFE SYR 0.5 ML 29G 1/2" OUTER
- SECURESAFE SYRNG 1 ML 29G 1/2" OUTER
- SKY SAFETY PEN NEEDLE 30G 5MM
- SKY SAFETY PEN NEEDLE 30G 8MM
- SURE CMFT SFTY PEN NDL 31G 6MM
- SURE CMFT SFTY PEN NDL 32G 4MM
- SURE COMFORT 0.5 ML SYRINGE
- SURE COMFORT 1 ML SYRINGE
- SURE COMFORT 3/10 ML SYRINGE

- SURE COMFORT 3/10 ML SYRINGE INSULIN SYRINGE
- SURE COMFORT 30G PEN NEEDLE
- SURE COMFORT ALCOHOL PREP PADS
- SURE COMFORT INS 0.3 ML 31GX1/4
- SURE COMFORT INS 0.5 ML 31GX1/4
- SURE COMFORT INS 1 ML 31GX1/4"
- SURE COMFORT PEN NDL 29GX1/2" 12.7MM
- SURE COMFORT PEN NDL 31G 5MM
- SURE COMFORT PEN NDL 31G 8MM
- SURE COMFORT PEN NDL 32G 4MM
- SURE COMFORT PEN NDL 32G 6MM
- SURE-FINE PEN NEEDLES 12.7MM
- SURE-FINE PEN NEEDLES 5MM
- SURE-FINE PEN NEEDLES 8MM
- SURE-JECT INSU SYR U100 0.3 ML
- SURE-JECT INSU SYR U100 0.5 ML
- SURE-JECT INSU SYR U100 1 ML
- SURE-JECT INSUL SYR U100 1 ML
- SURE-JECT INSULIN SYRINGE 1 ML
- SURE-PREP ALCOHOL PREP PADS
- TECHLITE 0.3 ML 29GX12MM (1/2)
- TECHLITE 0.3 ML 30GX8MM (1/2)
- TECHLITE 0.3 ML 31GX6MM (1/2)
- TECHLITE 0.3 ML 31GX8MM (1/2)
- TECHLITE 0.5 ML 30GX12MM (1/2)
- TECHLITE 0.5 ML 30GX8MM (1/2)
- TECHLITE 0.5 ML 31GX6MM (1/2)
- TECHLITE 0.5 ML 31GX8MM (1/2)
- TECHLITE INS SYR 1 ML 29GX12MM
- TECHLITE INS SYR 1 ML 30GX12MM
- TECHLITE INS SYR 1 ML 31GX6MM
- TECHLITE INS SYR 1 ML 31GX8MM
- TECHLITE PEN NEEDLE 29GX1/2"
- TECHLITE PEN NEEDLE 29GX3/8"
- TECHLITE PEN NEEDLE 31GX1/4"
- TECHLITE PEN NEEDLE 31GX3/16"
- TECHLITE PEN NEEDLE 31GX5/16"
- TECHLITE PEN NEEDLE 32GX1/4"
- TECHLITE PEN NEEDLE 32GX5/16"
- TECHLITE PEN NEEDLE 32GX5/32"
- TECHLITE PLUS PEN NDL 32G 4MM
- TERUMO INS SYRINGE U100-1 ML
- TERUMO INS SYRINGE U100-1/2 ML
- TERUMO INS SYRINGE U100-1/3 ML
- TERUMO INS SYRNG U100-1/2 ML
- THINPRO INS SYRIN U100-0.3 ML
- THINPRO INS SYRIN U100-0.5 ML
- THINPRO INS SYRIN U100-1 ML
- TRUE CMFRT PRO 0.5 ML 30G 5/16"
- TRUE CMFRT PRO 0.5 ML 31G 5/16"
- TRUE CMFRT PRO 0.5 ML 32G 5/16"
- TRUE CMFT SFTY PEN NDL 31G 5MM
- TRUE CMFT SFTY PEN NDL 31G 6MM
- TRUE CMFT SFTY PEN NDL 32G 4MM
- TRUE COMFORT 0.5 ML 30G 1/2"
- TRUE COMFORT 0.5 ML 30G 5/16"
- TRUE COMFORT 0.5 ML 31G 5/16"
- TRUE COMFORT 0.5 ML 31GX5/16"
- TRUE COMFORT 1 ML 31GX5/16"
- TRUE COMFORT ALCOHOL 70% PADS
- TRUE COMFORT PEN NDL 31G 8MM
- TRUE COMFORT PEN NDL 31GX5MM
- TRUE COMFORT PEN NDL 31GX6MM
- TRUE COMFORT PEN NDL 32G 5MM
- TRUE COMFORT PEN NDL 32G 6MM
- TRUE COMFORT PEN NDL 32GX4MM
- TRUE COMFORT PEN NDL 33G 4MM
- TRUE COMFORT PEN NDL 33G 5MM
- TRUE COMFORT PEN NDL 33G 6MM
- TRUE COMFORT PRO 1 ML 30G 1/2"
- TRUE COMFORT PRO 1 ML 30G 5/16"
- TRUE COMFORT PRO 1 ML 31G 5/16"
- TRUE COMFORT PRO 1 ML 32G 5/16"
- TRUE COMFORT PRO ALCOHOL PADS
- TRUE COMFORT SFTY 1 ML 30G 1/2"
- TRUE COMFRT PRO 0.5 ML 30G 1/2"
- TRUE COMFRT SFTY 1 ML 30G 5/16"
- TRUE COMFRT SFTY 1 ML 31G 5/16"
- TRUE COMFRT SFTY 1 ML 32G 5/16"
- TRUE-CMFRT PRO PEN NDL 31G 5MM
- TRUE-CMFRT PRO PEN NDL 31G 6MM
- TRUE-CMFRT PRO PEN NDL 31G 8MM
- TRUE-CMFRT PRO PEN NDL 32G 4MM
- TRUEPLUS PEN NEEDLE 29GX1/2"
- TRUEPLUS PEN NEEDLE 31G X 1/4"
- TRUEPLUS PEN NEEDLE 31GX3/16"
- TRUEPLUS PEN NEEDLE 31GX5/16"
- TRUEPLUS PEN NEEDLE 32GX5/32"
- TRUEPLUS SYR 0.3 ML 29GX1/2"
- TRUEPLUS SYR 0.3 ML 30GX5/16"
- TRUEPLUS SYR 0.3 ML 31GX5/16"
- TRUEPLUS SYR 0.5 ML 28GX1/2"

- TRUEPLUS SYR 0.5 ML 29GX1/2"
- TRUEPLUS SYR 0.5 ML 30GX5/16"
- TRUEPLUS SYR 0.5 ML 31GX5/16"
- TRUEPLUS SYR 1 ML 28GX1/2"
- TRUEPLUS SYR 1 ML 29GX1/2"
- TRUEPLUS SYR 1 ML 30GX5/16"
- TRUEPLUS SYR 1 ML 31GX5/16"
- ULT CFT 0.3 ML 31GX5/16" (1/2) 1/2 UNIT
- ULTICAR INS 0.3 ML 31GX1/4(1/2)
- ULTICARE INS 0.3 ML 31GX1/4"
- ULTICARE INS 0.5 ML 31GX1/4"
- ULTICARE INS 1 ML 31GX1/4"
- ULTICARE INS SYR 1 ML 30GX1/2"
- ULTICARE PEN NEEDLE 31GX3/16"
- ULTICARE PEN NEEDLE 6MM 31G
- ULTICARE PEN NEEDLE 8MM 31G
- ULTICARE PEN NEEDLES 12MM 29G
- ULTICARE PEN NEEDLES 4MM 32G MICRO, 32GX4MM
- ULTICARE PEN NEEDLES 6MM 32G
- ULTICARE SAFE PEN NDL 30G 8MM
- ULTICARE SAFE PEN NDL 5MM 30G
- ULTICARE SYR 0.3 ML 30GX1/2"
- ULTICARE SYR 0.3 ML 31GX5/16" SHORT NDL
- ULTICARE SYR 0.5 ML 30GX1/2"
- ULTICARE SYR 0.5 ML 31GX5/16" SHORT NDL
- ULTICARE SYR 1 ML 31GX5/16"
- ULTIGUARD SAFE 1 ML 30G 12.7MM
- ULTIGUARD SAFE0.3 ML 30G 12.7MM
- ULTIGUARD SAFE0.5 ML 30G 12.7MM
- ULTIGUARD SAFEPACK 1 ML 31G 8MM
- ULTIGUARD SAFEPACK 29G 12.7MM
- ULTIGUARD SAFEPACK 31G 5MM
- ULTIGUARD SAFEPACK 31G 6MM
- ULTIGUARD SAFEPACK 31G 8MM
- ULTIGUARD SAFEPACK 32G 4MM
- ULTIGUARD SAFEPACK 32G 6MM
- ULTIGUARD SAFEPK 0.3 ML 31G 8MM
- ULTIGUARD SAFEPK 0.5 ML 31G 8MM
- ULTILET ALCOHOL STERL SWAB
- ULTILET INSULIN SYRINGE 0.3 ML
- ULTILET INSULIN SYRINGE 0.5 ML
- ULTILET INSULIN SYRINGE 1 ML
- ULTILET PEN NEEDLE
- ULTILET PEN NEEDLE 4MM 32G
- ULTRA COMFORT 0.3 ML 29GX1/2"
- ULTRA COMFORT 0.3 ML SYRINGE
- ULTRA COMFORT 0.5 ML 28GX1/2" CONVERTS TO 29G
- ULTRA COMFORT 0.5 ML 29GX1/2"
- ULTRA COMFORT 0.5 ML 31GX5/16"
- ULTRA COMFORT 0.5 ML SYRINGE
- ULTRA COMFORT 1 ML 29GX1/2"
- ULTRA COMFORT 1 ML 30GX5/16"
- ULTRA COMFORT 1 ML 31GX5/16"
- ULTRA COMFORT 1 ML SYRINGE
- ULTRA FLO 0.3 ML 30G 1/2" (1/2)
- ULTRA FLO 0.3 ML 30G 5/16"(1/2)
- ULTRA FLO 0.3 ML 31G 5/16"(1/2)
- ULTRA FLO PEN NEEDLE 31G 5MM
- ULTRA FLO PEN NEEDLE 31G 8MM
- ULTRA FLO PEN NEEDLE 32G 4MM
- ULTRA FLO PEN NEEDLE 33G 4MM
- ULTRA FLO PEN NEEDLES 12MM 29G
- ULTRA FLO SYR 0.3 ML 29GX1/2"
- ULTRA FLO SYR 0.3 ML 30G 5/16"
- ULTRA FLO SYR 0.3 ML 31G 5/16"
- ULTRA FLO SYR 0.5 ML 29G 1/2"
- ULTRA THIN PEN NDL 32G X 4MM
- ULTRA-FINE 0.3 ML 30G 12.7MM
- ULTRA-FINE 0.3 ML 31G 6MM (1/2)
- ULTRA-FINE 0.3 ML 31G 8MM (1/2)
- ULTRA-FINE 0.5 ML 30G 12.7MM
- ULTRA-FINE INS SYR 1 ML 31G 6MM
- ULTRA-FINE INS SYR 1 ML 31G 8MM
- ULTRA-FINE PEN NDL 29G 12.7MM
- ULTRA-FINE PEN NEEDLE 31G 5MM
- ULTRA-FINE PEN NEEDLE 31G 8MM
- ULTRA-FINE PEN NEEDLE 32G 6MM
- ULTRA-FINE SYR 0.3 ML 31G 6MM
- ULTRA-FINE SYR 0.3 ML 31G 8MM
- ULTRA-FINE SYR 0.5 ML 31G 6MM
- ULTRA-FINE SYR 0.5 ML 31G 8MM
- ULTRA-FINE SYR 1 ML 30G 12.7MM
- ULTRA-THIN II 1 ML 31GX5/16"
- ULTRA-THIN II INS 0.3 ML 30G
- ULTRA-THIN II INS 0.3 ML 31G
- ULTRA-THIN II INS 0.5 ML 29G
- ULTRA-THIN II INS 0.5 ML 30G
- ULTRA-THIN II INS 0.5 ML 31G
- ULTRA-THIN II INS SYR 1 ML 29G

- ULTRA-THIN II INS SYR 1 ML 30G
- ULTRA-THIN II PEN NDL 29GX1/2"
- ULTRA-THIN II PEN NDL 31GX5/16
- ULTRACARE INS 0.3 ML 30GX5/16"
- ULTRACARE INS 0.3 ML 31GX5/16"
- ULTRACARE INS 0.5 ML 30GX1/2"
- ULTRACARE INS 0.5 ML 30GX5/16"
- ULTRACARE INS 0.5 ML 31GX5/16"
- ULTRACARE INS 1 ML 30G X 5/16"
- ULTRACARE INS 1 ML 30GX1/2"
- ULTRACARE INS 1 ML 31G X 5/16"
- ULTRACARE PEN NEEDLE 31GX1/4"
- ULTRACARE PEN NEEDLE 31GX3/16"
- ULTRACARE PEN NEEDLE 31GX5/16"
- ULTRACARE PEN NEEDLE 32GX1/4"
- ULTRACARE PEN NEEDLE 32GX3/16"
- ULTRACARE PEN NEEDLE 32GX5/32"
- ULTRACARE PEN NEEDLE 33GX5/32"
- UNIFINE OTC PEN NEEDLE 31G 5MM
- UNIFINE OTC PEN NEEDLE 32G 4MM
- UNIFINE PENTIPS 12MM 29G 29GX12MM, STRL
- UNIFINE PENTIPS 31GX3/16" 31GX5MM,STRL,MINI
- UNIFINE PENTIPS 32G 4MM
- UNIFINE PENTIPS 32GX1/4"
- UNIFINE PENTIPS 33GX5/32"
- UNIFINE PENTIPS 6MM 31G
- UNIFINE PENTIPS 8MM 31G 31GX8MM, SHORT, STRL
- UNIFINE PENTIPS MAX 30GX3/16"
- UNIFINE PENTIPS PLUS 29GX1/2" 12MM
- UNIFINE PENTIPS PLUS 30GX3/16"
- UNIFINE PENTIPS PLUS 31G 5MM
- UNIFINE PENTIPS PLUS 31GX1/4" ULTRA SHORT, 6MM
- UNIFINE PENTIPS PLUS 32GX5/32"
- UNIFINE PENTIPS PLUS 33GX5/32"
- UNIFINE PROTECT 30G 5MM
- UNIFINE PROTECT 30G 8MM
- UNIFINE PROTECT 32G 4MM
- UNIFINE SAFECONTROL 30G 5MM
- UNIFINE SAFECONTROL 30G 8MM
- UNIFINE SAFECONTROL 31G 5MM
- UNIFINE SAFECONTROL 31G 6MM
- UNIFINE SAFECONTROL 31G 8MM
- UNIFINE SAFECONTROL 32G 4MM
- UNIFINE ULTRA PEN NDL 31G 5MM
- UNIFINE ULTRA PEN NDL 31G 6MM
- UNIFINE ULTRA PEN NDL 31G 8MM
- UNIFINE ULTRA PEN NDL 32G 4MM
- VANISHPOINT 0.5 ML 30GX1/2" SY OUTER
- VANISHPOINT INS 0.5 ML 30G 8MM OUTER
- VANISHPOINT INS 1 ML 30GX3/16"
- VANISHPOINT U-100 29X1/2 SYR
- VERIFINE INS SYR 1 ML 29G 1/2"
- VERIFINE PEN NEEDLE 29G 12MM
- VERIFINE PEN NEEDLE 31G 5MM
- VERIFINE PEN NEEDLE 31G X 6MM
- VERIFINE PEN NEEDLE 31G X 8MM
- VERIFINE PEN NEEDLE 32G 6MM
- VERIFINE PEN NEEDLE 32G X 4MM
- VERIFINE PEN NEEDLE 32G X 5MM
- VERIFINE PLUS PEN NDL 31G 5MM
- VERIFINE PLUS PEN NDL 31G 8MM
- VERIFINE PLUS PEN NDL 32G 4MM
- VERIFINE PLUS PEN NDL 32G 4MM-SHARPS CONTAINER
- VERIFINE SYRING 0.5 ML 29G 1/2"
- VERIFINE SYRING 1 ML 31G 5/16"
- VERIFINE SYRNG 0.3 ML 31G 5/16"
- VERIFINE SYRNG 0.5 ML 31G 5/16"
- VERSALON ALL PURPOSE SPONGE 25'S,N-STERILE,3PLY
- WEBCOL ALCOHOL PREPS 20'S,LARGE
- WM UNIFINE PENTIP PLUS 8MM 31G W/ SAFETY CLICK

PA Criteria	Criteria Details
Exclusion Criteria	ONLY COVERED UNDER PART D WHEN USED CONCURRENTLY WITH INSULIN.

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

INTERFERON FOR MS-AVONEX

Products Affected

- AVONEX INTRAMUSCULAR PEN INJECTOR KIT
- AVONEX INTRAMUSCULAR SYRINGE KIT

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

INTERFERON FOR MS-BETASERON

Products Affected

- BETASERON SUBCUTANEOUS KIT

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

INTERFERON FOR MS-PLEGRIDY

Products Affected

- PLEGRIDY SUBCUTANEOUS PEN INJECTOR 125 MCG/0.5 ML, 63 MCG/0.5 ML- 94 MCG/0.5 ML
- PLEGRIDY SUBCUTANEOUS SYRINGE

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

INTERFERON GAMMA-1B

Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: CHRONIC GRANULOMATOUS DISEASE (CGD): PRESCRIBED BY OR IN CONSULTATION WITH A HEMATOLOGIST, INFECTIOUS DISEASE SPECIALIST, OR IMMUNOLOGIST. SEVERE MALIGNANT OSTEOPETROSIS (SMO): PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST OR HEMATOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	RENEWAL: CGD, SMO: 1) DEMONSTRATED CLINICAL BENEFIT COMPARED TO BASELINE, AND 2) HAS NOT RECEIVED HEMATOPOIETIC CELL TRANSPLANTATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

IPILIMUMAB

Products Affected

- YERVOY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: UNRESECT/MET MEL: 4MO, RCC/CRC/HCC: 3MO, ALL OTHERS: 12MO. INITIAL/RENEWAL: CUTAN MEL: 6MO
Other Criteria	RENEWAL: ADJUVANT CUTANEOUS MELANOMA: NO EVIDENCE OF DISEASE RECURRENCE (DEFINED AS THE APPEARANCE OF ONE OR MORE NEW MELANOMA LESIONS: LOCAL, REGIONAL OR DISTANT METASTASIS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

ISAVUCONAZONIUM

Products Affected

- CRESEMBA ORAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INVASIVE ASPERGILLOSIS, INVASIVE MUCORMYCOSIS: PRESCRIBED BY OR IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	6 MONTHS
Other Criteria	INVASIVE ASPERGILLOSIS: TRIAL OF OR CONTRAINDICATION TO VORICONAZOLE. CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE REQUIRES NO EXTRA CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

IVACAFTOR

Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CYSTIC FIBROSIS (CF): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT
Coverage Duration	INITIAL: 12 MONTHS. RENEWAL: LIFETIME
Other Criteria	CF: INITIAL: 1) NOT HOMOZYGOUS FOR F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR. RENEWAL: 1) IMPROVEMENT IN CLINICAL STATUS, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

IVOSIDENIB

Products Affected

- TIBSOVO

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

IXAZOMIB

Products Affected

- NINLARO

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

LAMOTRIGINE

Products Affected

- SUBVENITE ORAL SUSPENSION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ALL INDICATIONS: CONTRAINDICATION TO OR UNABLE TO SWALLOW LAMOTRIGINE TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LANREOTIDE

Products Affected

- *lanreotide subcutaneous syringe 120 mg/0.5 ml*
- SOMATULINE DEPOT SUBCUTANEOUS SYRINGE 60 MG/0.2 ML, 90 MG/0.3 ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	ACROMEGALY: INITIAL: THERAPY IS PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	ACROMEGALY: INITIAL/RENEWAL: 12 MOS. GEP-NETS, CARCINOID SYNDROME: 12 MOS.
Other Criteria	ACROMEGALY: RENEWAL: 1) REDUCTION, NORMALIZATION, OR MAINTENANCE OF IGF-1 LEVELS BASED ON AGE AND GENDER, AND 2) IMPROVEMENT OR SUSTAINED REMISSION OF CLINICAL SYMPTOMS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LAPATINIB

Products Affected

- *lapatinib*

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

LAROTRECTINIB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	VITRAKVI ORAL SOLUTION: 1) TRIAL OF VITRAKVI CAPSULES, OR 2) UNABLE TO TAKE CAPSULE FORMULATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

LAZERTINIB

Products Affected

- LAZCLUZE ORAL TABLET 240 MG, 80 MG

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

LEDIPASVIR-SOFOSBUVIR

Products Affected

- HARVONI ORAL PELLETS IN PACKET
33.75-150 MG, 45-200 MG
- HARVONI ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, AND 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING: CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, ROSUVASTATIN, TIPRANAVIR/RITONAVIR, SOFOSBUVIR (AS A SINGLE AGENT), EPCLUSA, ZEPATIER, MAVYRET, OR VOSEVI.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

LENALIDOMIDE

Products Affected

- *lenalidomide*

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

LENVATINIB

Products Affected

- LENVIMA

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

LETERMOVIR

Products Affected

- PREVYMIS ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	HSCT: NOT AT RISK FOR LATE CMV: 4 MOS, AT RISK FOR LATE CMV: 7 MOS. KIDNEY TRANSPLANT: 7 MOS.
Other Criteria	HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT): 1) THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 28 POST TRANSPLANT, AND 2) WILL NOT RECEIVE THE MEDICATION BEYOND 100 DAYS POST TRANSPLANT IF NOT AT RISK FOR LATE CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE, OR BEYOND 200 DAYS POST TRANSPLANT IF AT RISK FOR LATE CMV INFECTION AND DISEASE. KIDNEY TRANSPLANT: 1) THERAPY WILL BE INITIATED BETWEEN DAY 0 AND DAY 7 POST TRANSPLANT, AND 2) WILL NOT RECEIVE THE MEDICATION BEYOND 200 DAYS POST TRANSPLANT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

LEUPROLIDE

Products Affected

- *leuprolide subcutaneous kit*

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

LEUPROLIDE DEPOT

Products Affected

- *leuprolide acetate (3 month)*
- LUTRATE DEPOT (3 MONTH)

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

LEUPROLIDE MESYLATE

Products Affected

- CAMCEVI (6 MONTH)

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

LEUPROLIDE-ELIGARD

Products Affected

- ELIGARD
- ELIGARD (3 MONTH)
- ELIGARD (4 MONTH)
- ELIGARD (6 MONTH)

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

LEUPROLIDE-LUPRON DEPOT

Products Affected

- LUPRON DEPOT
- LUPRON DEPOT (3 MONTH)
- LUPRON DEPOT (4 MONTH)
- LUPRON DEPOT (6 MONTH)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ENDOMETRIOSIS: DIAGNOSIS IS CONFIRMED VIA SURGICAL OR DIRECT VISUALIZATION (E.G., PELVIC ULTRASOUND) OR HISTOPATHOLOGICAL CONFIRMATION (E.G., LAPAROSCOPY OR LAPAROTOMY) IN THE LAST 10 YEARS.
Age Restrictions	
Prescriber Restrictions	INITIAL: ENDOMETRIOSIS: PRESCRIBED BY OR IN CONSULTATION WITH AN OBSTETRICIAN/GYNECOLOGIST.
Coverage Duration	PROSTATE CA: 12 MOS. UTERINE FIBROIDS: 3 MOS. ENDOMETRIOSIS: INITIAL/RENEWAL: 6 MOS.
Other Criteria	INITIAL: ENDOMETRIOSIS: 1) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, 2) TRIAL OF OR CONTRAINDICATION TO NSAID AND PROGESTIN-CONTAINING PREPARATION, AND 3) HAS NOT RECEIVED A TOTAL OF 12 MONTHS OF TREATMENT PER LIFETIME. RENEWAL: ENDOMETRIOSIS: 1) IMPROVEMENT OF PAIN RELATED TO ENDOMETRIOSIS WHILE ON THERAPY, 2) RECEIVING CONCOMITANT ADD-BACK THERAPY (I.E., COMBINATION ESTROGEN-PROGESTIN OR PROGESTIN-ONLY CONTRACEPTIVE PREPARATION), 3) NO CONCURRENT USE WITH ANOTHER GNRH-MODULATING AGENT, AND 4) HAS NOT RECEIVED A TOTAL OF 12 MONTHS OF TREATMENT PER LIFETIME.
Indications	All FDA-approved Indications.
Off Label Uses	

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

LEUPROLIDE-LUPRON DEPOT-PED

Products Affected

- LUPRON DEPOT-PED (3 MONTH)
- LUPRON DEPOT-PED
INTRAMUSCULAR SYRINGE KIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CENTRAL PRECOCIOUS PUBERTY (CPP): INITIAL: FEMALES: ELEVATED LEVEL OF LUTEINIZING HORMONE (LH) LEVEL GREATER THAN 0.2 TO 0.3 MIU/ML AT DIAGNOSIS. MALES: ELEVATED LEVEL OF LH LEVEL GREATER THAN 0.2 TO 0.3 MIU/ML AT DIAGNOSIS.
Age Restrictions	
Prescriber Restrictions	CPP: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	CPP: INITIAL: FEMALES: 1) YOUNGER THAN 8 YEARS OF AGE AT ONSET OF CPP, AND 2) AT TANNER STAGE 2 OR ABOVE FOR BREAST DEVELOPMENT AND PUBIC HAIR GROWTH. MALES: 1) YOUNGER THAN 9 YEARS OF AGE AT ONSET OF CPP, AND 2) AT TANNER STAGE 2 OR ABOVE FOR GENITAL DEVELOPMENT AND PUBIC HAIR GROWTH. RENEWAL: 1) TANNER STAGING AT INITIAL DIAGNOSIS HAS STABILIZED OR REGRESSED DURING THREE SEPARATE MEDICAL VISITS IN THE PREVIOUS YEAR, AND 2) HAS NOT REACHED ACTUAL AGE WHICH CORRESPONDS TO CURRENT PUBERTAL AGE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

L-GLUTAMINE

Products Affected

- *glutamine (sickle cell)*

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

LIDOCAINE OINTMENT

Products Affected

- *lidocaine topical ointment*

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

LIDOCAINE PATCH

Products Affected

- *dermacinrx lidocan 5% patch outer*
- *lidocaine topical adhesive patch,medicated 5 %*
- *lidocan iii*
- *tridacaine iii*
- ZTLIDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	1) PAIN ASSOCIATED WITH POST-HERPETIC NEURALGIA, 2) NEUROPATHY DUE TO DIABETES MELLITUS, 3) CHRONIC BACK PAIN, OR 4) OSTEOARTHRITIS OF THE KNEE OR HIP.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

LIDOCAINE PRILOCAINE

Products Affected

- *lidocaine-prilocaine topical cream*

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

LINVOSELTAMAB-GCPT

Products Affected

- LYNZOZYFIC INTRAVENOUS
SOLUTION 2 MG/ML, 20 MG/ML

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

LONCASTUXIMAB TESIRINE-LPYL

Products Affected

- ZYNLONTA

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

LORLATINIB

Products Affected

- LORBRENA ORAL TABLET 100 MG, 25 MG

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

LOTILANER

Products Affected

- XDEMVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	DEMODEX BLEPHARITIS: 18 YEARS OF AGE OR OLDER
Prescriber Restrictions	
Coverage Duration	6 WEEKS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

LUMACAFTOR-IVACAFTOR

Products Affected

- ORKAMBI ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CYSTIC FIBROSIS (CF): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CF EXPERT.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME.
Other Criteria	CF: INITIAL: NO CONCURRENT USE WITH ANOTHER CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) MODULATOR. RENEWAL: 1) IMPROVEMENT IN CLINICAL STATUS, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

MACITENTAN

Products Affected

- *macitentan*
- OPSUMIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

MARGETUXIMAB-CMKB

Products Affected

- MARGENZA

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

MARIBAVIR

Products Affected

- LIVTENCITY

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

MAVACAMTEN

Products Affected

- CAMZYOS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	OBSTRUCTIVE HYPERTROPHIC CARDIOMYOPATHY(HCM): INITIAL: LEFT VENTRICULAR OUTFLOW TRACK (LVOT) GRADIENT OF 50 MMHG OR HIGHER
Age Restrictions	
Prescriber Restrictions	OBSTRUCTIVE HCM: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	OBSTRUCTIVE HCM: INITIAL: TRIAL OF, CONTRAINDICATION, OR INTOLERANCE TO A BETA-BLOCKER OR A NON-DIHYDROPYRIDINE CALCIUM CHANNEL BLOCKER. RENEWAL: CONTINUED CLINICAL BENEFIT (E.G., REDUCTION OF SYMPTOMS, NYHA CLASSIFICATION IMPROVEMENT).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

MECASERMIN

Products Affected

- INCRELEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	GROWTH FAILURE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST OR NEPHROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	GROWTH FAILURE: INITIAL: OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF WRIST AND HAND. INITIAL/RENEWAL: NO CONCURRENT USE WITH ANOTHER GROWTH HORMONE MEDICATION. RENEWAL: IMPROVEMENT WHILE ON THERAPY (INCREASE IN HEIGHT OR INCREASE IN HEIGHT VELOCITY).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

MECHLORETHAMINE

Products Affected

- VALCHLOR

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

MEPOLIZUMAB

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ASTHMA: BLOOD EOSINOPHIL LEVEL OF AT LEAST 150 CELLS/MCL WITHIN THE PAST 12 MONTHS.
Age Restrictions	
Prescriber Restrictions	INITIAL: ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN PULMONARY OR ALLERGY MEDICINE. CRSWNP: PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. EOSINOPHILIC COPD: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST.
Coverage Duration	INITIAL: CRSWNP: 6 MO. OTHERS: 12 MO. RENEWAL: CRSWNP, ASTHMA, COPD, EGPA, HES: 12 MO.

PA Criteria	Criteria Details
Other Criteria	INITIAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 2) ONE OF THE FOLLOWING: (A) AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING AT LEAST 3 DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA. CHRONIC RHINOSINUSITIS WITH NASAL POLYPS (CRSWNP): 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, 2) EVIDENCE OF NASAL POLYPS BY DIRECT EXAMINATION, ENDOSCOPY, OR SINUS CT SCAN, AND 3) INADEQUATELY CONTROLLED DISEASE. EOSINOPHILIC COPD: USED IN COMBINATION WITH A LAMA/LABA/ICS. RENEWAL: ASTHMA: 1) CONTINUED USE OF ICS AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, 2) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR (D) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE, AND 3) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION. CRSWNP: 1) CLINICAL BENEFIT COMPARED TO BASELINE, AND 2) NO CONCURRENT USE WITH ANOTHER

PA Criteria	Criteria Details
	SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION. EOSINOPHILIC GRANULOMATOSIS WITH POLYANGIITIS (EGPA): 1) REDUCTION IN EGPA SYMPTOMS COMPARED TO BASELINE OR ABILITY TO REDUCE/ELIMINATE CORTICOSTEROID USE, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION. EOSINOPHILIC COPD: 1) USED IN COMBINATION WITH A LAMA/LABA/ICS, 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION, AND 3) CLINICAL RESPONSE AS EVIDENCED BY: (A) REDUCTION IN COPD EXACERBATIONS FROM BASELINE, OR (B) REDUCTION IN SEVERITY OR FREQUENCY OF COPD-RELATED SYMPTOMS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

METYROSIONE

Products Affected

- *metirosine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PHEOCHROMOCYTOMA: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST, ENDOCRINE SURGEON, OR HEMATOLOGIST-ONCOLOGIST.
Coverage Duration	PREOPERATIVE PREPARATION FOR SURGERY: 30 DAYS. MALIGNANT PHEOCHROMOCYTOMA: INITIAL/RENEWAL:12 MOS.
Other Criteria	PHEOCHROMOCYTOMA: INITIAL: HAS NON-METASTATIC PHEOCHROMOCYTOMA. PREOPERATIVE PREPARATION FOR SURGERY: USE IN COMBINATION WITH AN ALPHA-ADRENERGIC RECEPTOR BLOCKER. RENEWAL: MALIGNANT PHEOCHROMOCYTOMA: STABLE OR CLINICAL IMPROVEMENT WHILE ON THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

MIDOSTAURIN

Products Affected

- RYDAPT

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

MIFEPRISTONE

Products Affected

- *mifepristone oral tablet 300 mg*

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

MILTEFOSINE

Products Affected

- IMPAVIDO

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

MIRDAMETINIB

Products Affected

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

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

MIRVETUXIMAB SORAVTANSINE-GYNX

Products Affected

- ELAHERE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER: AN OPHTHALMIC EXAM, INCLUDING VISUAL ACUITY AND SLIT LAMP EXAM, WILL BE COMPLETED PRIOR TO THE INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

MOMELOTINIB

Products Affected

- OJJAARA

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

MOSUNETUZUMAB-AXGB

Products Affected

- LUNSUMIO
- LUNSUMIO VELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA: INITIAL: 6 MONTHS. RENEWAL: 7 MONTHS.
Other Criteria	RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA: RENEWAL: 1) HAS ACHIEVED A PARTIAL RESPONSE TO TREATMENT, AND 2) HAS NOT PREVIOUSLY RECEIVED MORE THAN 17 CYCLES OF TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

NARCOLEPSY AGENTS

Products Affected

- *armodafinil*
- *modafinil oral tablet 100 mg, 200 mg*

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

NAXITAMAB-GQGK

Products Affected

- DANYELZA

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

NERANDOMILAST

Products Affected

- JASCAYD

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: IDIOPATHIC PULMONARY FIBROSIS (IPF): 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. PROGRESSIVE PULMONARY FIBROSIS (PPF): 1) AT LEAST 10% FIBROSIS ON A CHEST HRCT, AND 2) INTERSTITIAL LUNG DISEASE WITH A PROGRESSIVE PHENOTYPE (PF-ILD).
Age Restrictions	
Prescriber Restrictions	INITIAL: IPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST. PPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: IPF: 1) DOES NOT HAVE 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, CANCER), AND 2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: OFEV, PIRFENIDONE. PPF: 1) LUNG FUNCTION AND RESPIRATORY SYMPTOMS OR CHEST IMAGING HAVE WORSENEDED/PROGRESSED DESPITE TREATMENT WITH MEDICATIONS USED IN CLINICAL PRACTICE FOR ILD (NOT ATTRIBUTABLE TO COMORBIDITIES SUCH AS INFECTION, HEART FAILURE), AND 2) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: OFEV (NINTEDANIB). RENEWAL: IPF, PPF: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

NERATINIB

Products Affected

- NERLYNX

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

NILOTINIB - TASIGNA

Products Affected

- *nilotinib hcl oral capsule 150 mg, 200 mg, 50 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND MEDICATION IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

NILOTINIB-DANZITEN

Products Affected

- DANZITEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PREVIOUSLY TREATED PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (Ph+ CML): 1) PERFORMED MUTATIONAL ANALYSIS PRIOR TO INITIATION OF THERAPY, AND 2) THERAPY IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

NINTEDANIB

Products Affected

- *nintedanib*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: IDIOPATHIC PULMONARY FIBROSIS (IPF): 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. SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD): 1) AT LEAST 10% FIBROSIS ON A CHEST HRCT, AND 2) BASELINE FVC AT LEAST 40% OF PREDICTED VALUE. CHRONIC FIBROSING INTERSTITIAL LUNG DISEASE WITH A PROGRESSIVE PHENOTYPE (PF-ILD): AT LEAST 10% FIBROSIS ON A CHEST HRCT.
Age Restrictions	
Prescriber Restrictions	INITIAL: IPF: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST. SSC-ILD, PF-ILD: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR RHEUMATOLOGIST.
Coverage Duration	INITIAL: SSC-ILD: 6 MOS. IPF, PF-ILD: 12 MOS. RENEWAL (ALL INDICATIONS): 12 MOS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: IPF: 1) DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS), AND 2) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: ESBRIET (PIRFENIDONE). SSC-ILD: DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., HEART FAILURE/FLUID OVERLOAD, DRUG-INDUCED LUNG TOXICITY, RECURRENT ASPIRATION). PF-ILD: LUNG FUNCTION AND RESPIRATORY SYMPTOMS OR CHEST IMAGING HAVE WORSENERD/PROGRESSED DESPITE TREATMENT WITH MEDICATIONS USED IN CLINICAL PRACTICE FOR ILD (NOT ATTRIBUTABLE TO COMORBIDITIES SUCH AS INFECTION, HEART FAILURE). RENEWAL: IPF, SSC-ILD, PF-ILD: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

NIRAPARIB

Products Affected

- ZEJULA ORAL CAPSULE
- ZEJULA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER: 1) ZEJULA WILL BE USED AS MONOTHERAPY, AND 2) ZEJULA IS STARTED NO LATER THAN 8 WEEKS AFTER THE MOST RECENT PLATINUM-CONTAINING REGIMEN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

NIRAPARIB-ABIRATERONE

Products Affected

- AKEEGA

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

NIROGACESTAT

Products Affected

- OGSIVEO ORAL TABLET 100 MG, 150 MG, 50 MG

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

NITISINONE

Products Affected

- *nitisinone*
- ORFADIN ORAL SUSPENSION

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

NIVOLUMAB

Products Affected

- OPDIVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA: NO CONCURRENT USE WITH TARGETED THERAPY (I.E., BRAF INHIBITORS, MEK INHIBITORS, AND NTRK INHIBITORS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

NIVOLUMAB-HYALURONIDASE-NVHY

Products Affected

- OPDIVO QVANTIG

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

NIVOLUMAB-RELATLIMAB-RMBW

Products Affected

- OPDUALAG

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

NOGAPENDEKIN ALFA

Products Affected

- ANKTIVA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	40 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

OFATUMUMAB SQ

Products Affected

- KESIMPTA PEN

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

OLAPARIB

Products Affected

- LYNPARZA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER: MEDICATION WILL BE USED AS MONOTHERAPY. METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG. ALL OTHER FDA APPROVED INDICATIONS ARE COVERED WITHOUT ADDITIONAL CRITERIA, EXCEPT THOSE CRITERIA IN THE FDA APPROVED LABEL.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

OLUTASIDENIB

Products Affected

- REZLIDHIA

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

OMACETAXINE

Products Affected

- SYNRIBO

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

OMALIZUMAB

Products Affected

- XOLAIR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: ASTHMA: POSITIVE SKIN PRICK OR BLOOD TEST (E.G., ELISA, FEIA) TO A PERENNIAL AEROALLERGEN AND A BASELINE IGE SERUM LEVEL OF AT LEAST 30 IU/ML. FOOD ALLERGY: 1) IGE SERUM LEVEL OF AT LEAST 30 IU/ML, AND 2) POSITIVE SKIN PRICK TEST TO AT LEAST ONE FOOD, OR POSITIVE MEDICALLY MONITORED FOOD CHALLENGE TO AT LEAST ONE FOOD.
Age Restrictions	
Prescriber Restrictions	INITIAL/RENEWAL: CHRONIC SPONTANEOUS URTICARIA (CSU): PRESCRIBED BY OR IN CONSULTATION WITH AN ALLERGIST, DERMATOLOGIST, OR IMMUNOLOGIST. INITIAL: CHRONIC RHINOSINUSITIS WITH NASAL POLYPS (CRSWNP): PRESCRIBED BY OR IN CONSULTATION WITH AN OTOLARYNGOLOGIST, ALLERGIST OR IMMUNOLOGIST. ASTHMA: PRESCRIBED BY OR IN CONSULTATION WITH A PHYSICIAN SPECIALIZING IN ALLERGY OR PULMONARY MEDICINE. FOOD ALLERGY: PRESCRIBED BY OR IN CONSULTATION WITH ALLERGIST OR IMMUNOLOGIST.
Coverage Duration	INITIAL/RENEWAL: ASTHMA 12 MO/12 MO, CSU 6 MO/12 MO, CRSWNP 6 MO/12 MO, FOOD ALLERGY 12 MO/24 MO

PA Criteria	Criteria Details
Other Criteria	INITIAL: CSU: 1) TRIAL OF AND MAINTAINED ON, OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE, 2) STILL EXPERIENCES HIVES OR ANGIOEDEMA ON MOST DAYS OF THE WEEK FOR AT LEAST 6 WEEKS, AND 3) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: DUPIXENT. CRSWNP: 1) A 56 DAY TRIAL OF ONE TOPICAL NASAL CORTICOSTEROID, 2) TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: NUCALA, DUPIXENT, 3) EVIDENCE OF NASAL POLYPS BY DIRECT EXAMINATION, ENDOSCOPY, OR SINUS CT SCAN, AND 4) INADEQUATELY CONTROLLED DISEASE. ASTHMA: 1) CONCURRENT THERAPY WITH A MEDIUM, HIGH-DOSE OR MAXIMALLY TOLERATED DOSE OF AN INHALED CORTICOSTEROID (ICS) AND AT LEAST ONE OTHER MAINTENANCE MEDICATION, AND 2) ONE OF THE FOLLOWING: (A) AT LEAST ONE ASTHMA EXACERBATION REQUIRING SYSTEMIC CORTICOSTEROID BURST LASTING AT LEAST 3 DAYS WITHIN THE PAST 12 MONTHS OR AT LEAST ONE SERIOUS EXACERBATION REQUIRING HOSPITALIZATION OR ER VISIT WITHIN THE PAST 12 MONTHS, OR (B) POOR SYMPTOM CONTROL DESPITE CURRENT THERAPY AS EVIDENCED BY AT LEAST THREE OF THE FOLLOWING WITHIN THE PAST 4 WEEKS: DAYTIME ASTHMA SYMPTOMS MORE THAN TWICE/WEEK, ANY NIGHT WAKING DUE TO ASTHMA, SABA RELIEVER FOR SYMPTOMS MORE THAN TWICE/WEEK, ANY ACTIVITY LIMITATION DUE TO ASTHMA. FOOD ALLERGY: CONCURRENT USE WITH AN ACTIVE PRESCRIPTION FOR EPINEPHRINE AUTO-INJECTOR/INJECTION . INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: CSU: MAINTAINED ON OR CONTRAINDICATION TO A SECOND GENERATION H1 ANTI-HISTAMINE. CRSWNP: CLINICAL BENEFIT COMPARED TO BASELINE. ASTHMA: 1) CONTINUED USE OF ICS AND AT LEAST

PA Criteria	Criteria Details
	ONE OTHER MAINTENANCE MEDICATION, AND 2) CLINICAL RESPONSE AS EVIDENCED BY ONE OF THE FOLLOWING: (A) REDUCTION IN ASTHMA EXACERBATIONS FROM BASELINE, (B) DECREASED UTILIZATION OF RESCUE MEDICATIONS, (C) REDUCTION IN SEVERITY OR FREQUENCY OF ASTHMA-RELATED SYMPTOMS, OR (D) INCREASE IN PERCENT PREDICTED FEV1 FROM PRETREATMENT BASELINE. FOOD ALLERGY: 1) PERSISTENT IGE-MEDIATED FOOD ALLERGY, AND 2) CONCURRENT USE WITH AN ACTIVE PRESCRIPTION FOR EPINEPHRINE AUTO-INJECTOR/INJECTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

OSIMERTINIB

Products Affected

- TAGRISSO

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

OXANDROLONE

Products Affected

- *oxandrolone*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 MONTHS
Other Criteria	PROTEIN CATABOLISM, BONE PAIN: 1) MONITORED FOR PELIOSIS HEPATIS, LIVER CELL TUMORS, AND BLOOD LIPID CHANGES, 2) DOES NOT HAVE KNOWN OR SUSPECTED: CARCINOMA OF THE PROSTATE OR BREAST IN MALE PATIENTS, CARCINOMA OF THE BREAST IN FEMALES WITH HYPERCALCEMIA, NEPHROSIS (THE NEPHROTIC PHASE OF NEPHRITIS), OR HYPERCALCEMIA, AND 3) DOES NOT HAVE SEVERE HEPATIC DYSFUNCTION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PACRITINIB

Products Affected

- VONJO

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

PALBOCICLIB

Products Affected

- IBRANCE

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

PASIREOTIDE DIASPARTATE

Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CUSHINGS DISEASE (CD): INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	CD: RENEWAL: 1) CONTINUED IMPROVEMENT OF CUSHINGS DISEASE, AND 2) MAINTAINED TOLERABILITY TO SIGNIFOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PAZOPANIB

Products Affected

- *pazopanib oral tablet 200 mg, 400 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED SOFT TISSUE SARCOMA (STS): NOT USED FOR ADIPOCYTIC STS OR GASTROINTESTINAL STROMAL TUMORS (GIST)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PEGFILGRASTIM - APGF

Products Affected

- NYVEPRIA

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

PEGFILGRASTIM - CBQV

Products Affected

- UDENYCA ONBODY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	PREScribed BY OR IN CONSULTATION WITH A HEMATOLOGIST OR ONCOLOGIST.
Coverage Duration	12 MONTHS
Other Criteria	NON MYELOID MALIGNANCY: UDENYCA: TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT NYVEPRIA. UDENYCA ONBODY: 1) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT NYVEPRIA, OR 2) BARRIER TO ACCESS (E.G., TRAVEL BARRIERS, UNABLE TO RETURN TO CLINIC FOR INJECTIONS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

PEGINTERFERON ALFA-2A

Products Affected

- PEGASYS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HEPATITIS B: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, INFECTIOUS DISEASE SPECIALIST, OR PHYSICIAN SPECIALIZING IN THE TREATMENT OF HEPATITIS (E.G., HEPATOLOGIST).
Coverage Duration	HEP B/HEP C: 48 WEEKS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PEGVISOMANT

Products Affected

- SOMAVERT

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

PEMBROLIZUMAB

Products Affected

- KEYTRUDA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA: NO CONCURRENT USE WITH TARGETED THERAPY (I.E., BRAF INHIBITORS, MEK INHIBITORS, AND NTRK INHIBITORS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PEMBROLIZUMAB-BERAHYALURONIDASE ALFA-PMPH

Products Affected

- KEYTRUDA QLEX

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

PEMIGATINIB

Products Affected

- PEMAZYRE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CHOLANGIOCARCINOMA, MYELOID/LYMPHOID NEOPLASMS: COMPREHENSIVE OPHTHALMOLOGICAL EXAMINATION, INCLUDING OPTICAL COHERENCE TOMOGRAPHY (OCT), WILL BE COMPLETED PRIOR TO INITIATION OF THERAPY AND AT THE RECOMMENDED SCHEDULED INTERVALS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PENICILLAMINE TABLET

Products Affected

- *penicillamine oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: CYSTINURIA: HAS NEPHROLITHIASIS AND ONE OF THE FOLLOWING: 1) STONE ANALYSIS SHOWING PRESENCE OF CYSTINE, 2) PRESENCE OF PATHOGNOMONIC HEXAGONAL CYSTINE CRYSTALS ON URINALYSIS, OR 3) FAMILY HISTORY OF CYSTINURIA AND POSITIVE CYANIDE-NITROPRUSSIDE SCREENING.
Age Restrictions	
Prescriber Restrictions	INITIAL: WILSONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST OR GASTROENTEROLOGIST. CYSTINURIA: PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST. RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: 12 MONTHS, RENEWAL: LIFETIME.
Other Criteria	INITIAL: WILSONS DISEASE: 1) LEIPZIG SCORE OF 4 OR GREATER. RA: 1) NO HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY, AND 2) TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE GREATER THAN OR EQUAL TO 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. RENEWAL: RA: 1) NO HISTORY OR OTHER EVIDENCE OF RENAL INSUFFICIENCY, AND 2) EXPERIENCED OR MAINTAINED IMPROVEMENT IN TENDER JOINT COUNT OR SWOLLEN JOINT COUNT COMPARED TO BASELINE. WILSONS DISEASE, CYSTINURIA: CONTINUES TO BENEFIT FROM THE MEDICATION.

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

PEXIDARTINIB

Products Affected

- TURALIO

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

PIMAVANSERIN

Products Affected

- NUPLAZID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	PSYCHOSIS IN PARKINSONS DISEASE (PD): INITIAL: 18 YEARS OR OLDER
Prescriber Restrictions	PSYCHOSIS IN PD: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, GERIATRICIAN, OR A BEHAVIORAL HEALTH SPECIALIST (E.G., PSYCHIATRIST).
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PSYCHOSIS IN PD: RENEWAL: IMPROVEMENT IN PSYCHOSIS SYMPTOMS FROM BASELINE AND DEMONSTRATES A CONTINUED NEED FOR TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PIRFENIDONE

Products Affected

- *pirfenidone oral capsule*
- *pirfenidone oral tablet 267 mg, 534 mg, 801 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	IDIOPATHIC PULMONARY FIBROSIS (IPF): INITIAL: 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	IPF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	IPF: INITIAL: 1) DOES NOT HAVE OTHER KNOWN CAUSES OF INTERSTITIAL LUNG DISEASE (E.G., CONNECTIVE TISSUE DISEASE, DRUG TOXICITY, ASBESTOS OR BERYLLIUM EXPOSURE, HYPERSENSITIVITY PNEUMONITIS, SYSTEMIC SCLEROSIS, RHEUMATOID ARTHRITIS, RADIATION, SARCOIDOSIS, BRONCHIOLITIS OBLITERANS ORGANIZING PNEUMONIA, HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION, VIRAL HEPATITIS, OR CANCER). RENEWAL: CLINICAL MEANINGFUL IMPROVEMENT OR MAINTENANCE IN ANNUAL RATE OF DECLINE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

PIRTOBRUTINIB

Products Affected

- JAYPIRCA ORAL TABLET 100 MG, 50 MG

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

POMALIDOMIDE

Products Affected

- *pomalidomide*
- POMALYST

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

PONATINIB

Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CHRONIC MYELOID LEUKEMIA (CML): MUTATIONAL ANALYSIS PRIOR TO INITIATION AND ICLUSIG IS APPROPRIATE PER NCCN GUIDELINE TABLE FOR TREATMENT RECOMMENDATIONS BASED ON BCR-ABL1 MUTATION PROFILE.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

POSACONAZOLE TABLET

Products Affected

- *posaconazole oral tablet, delayed release (dr/ec)*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CONTINUATION OF THERAPY AFTER HOSPITAL DISCHARGE, PROPHYLAXIS: 6 MONTHS. TREATMENT: 12 WEEKS.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

PRALSETINIB

Products Affected

- GAVRETO

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

PYRIMETHAMINE

Products Affected

- *pyrimethamine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	TOXOPLASMOSIS: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH AN INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	TOXOPLASMOSIS: INITIAL: 8 WEEKS, RENEWAL: 6 MOS.
Other Criteria	TOXOPLASMOSIS: RENEWAL: 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 CURRENTLY TAKING AN ANTI-RETROVIRAL THERAPY IF HIV POSITIVE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

QUININE

Products Affected

- *quinine sulfate*

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

QUIZARTINIB

Products Affected

- VANFLYTA

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

REGORAFENIB

Products Affected

- STIVARGA

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

RELUGOLIX

Products Affected

- ORGOVYX

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

REPOTRECTINIB

Products Affected

- AUGTYRO ORAL CAPSULE 160 MG, 40 MG

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

RESMETIROM

Products Affected

- REZDIFFRA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	NONALCOHOLIC STEATOHEPATITIS (NASH): INITIAL: DIAGNOSIS CONFIRMED BY BIOPSY OR NONINVASIVE TESTING, OBTAINED IN THE PAST 12 MONTHS, DEMONSTRATING: 1) LIVER FIBROSIS STAGE 2 OR 3, OR 2) NONALCOHOLIC FATTY LIVER DISEASE (NAFLD) ACTIVITY SCORE OF 4 OR MORE.
Age Restrictions	
Prescriber Restrictions	NASH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST, GASTROENTEROLOGIST, OR ENDOCRINOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	NASH: RENEWAL: CONTINUES TO HAVE NONCIRRHOTIC NASH WITH MODERATE TO ADVANCED LIVER FIBROSIS (CONSISTENT WITH STAGES F2 TO F3 FIBROSIS).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

RETIFANLIMAB-DLWR

Products Affected

- ZYNYZ

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

REVUMENIB

Products Affected

- REVUFORJ ORAL TABLET 110 MG, 160 MG, 25 MG

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

RIBOCICLIB

Products Affected

- KISQALI 200 MG DAILY DOSE
- KISQALI ORAL TABLET 200 MG/DAY (200 MG X 1), 400 MG/DAY (200 MG X 2), 600 MG/DAY (200 MG X 3)

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

RIBOCICLIB-LETROZOLE

Products Affected

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

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

RIFAXIMIN

Products Affected

- XIFAXAN ORAL TABLET 200 MG, 550 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	TRAVELERS DIARRHEA, HEPATIC ENCEPHALOPATHY (HE): 12 MOS. IBS-D: 8 WKS.
Other Criteria	HE: TRIAL OF OR CONTRAINDICATION TO LACTULOSE OR CONCURRENT LACTULOSE THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RILONACEPT

Products Affected

- ARCALYST

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CRYOPYRIN-ASSOCIATED PERIODIC SYNDROMES (CAPS): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE NLRP3 GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR, SERUM AMYLOID A PROTEIN (SAA) OR S100 PROTEINS), AND 2) TWO OF THE FOLLOWING: URTICARIAL-LIKE RASH (NEUTROPHILIC DERMATITIS), COLD-TRIGGERED EPISODES, SENSORINEURAL HEARING LOSS, MUSCULOSKELETAL SYMPTOMS, CHRONIC ASEPTIC MENINGITIS, SKELETAL ABNORMALITIES. DEFICIENCY OF INTERLEUKIN-1 RECEPTOR ANTAGONIST (DIRA): 1) ONE OF THE FOLLOWING: (A) GENETIC TEST FOR GAIN-OF-FUNCTION MUTATIONS IN THE IL1RN GENE, OR (B) HAS INFLAMMATORY MARKERS (I.E., ELEVATED CRP, ESR), AND 2) ONE OF THE FOLLOWING: PUSTULAR PSORIASIS-LIKE RASHES, OSTEOMYELITIS, ABSENCE OF BACTERIAL OSTEOMYELITIS, ONYCHOMADESIS. RECURRENT PERICARDITIS (RP): TWO OF THE FOLLOWING: CHEST PAIN CONSISTENT WITH PERICARDITIS, PERICARDIAL FRICTION RUB, ECG SHOWING DIFFUSE ST-SEGMENT ELEVATION OR PR-SEGMENT DEPRESSION, NEW OR WORSENING PERICARDIAL EFFUSION.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CAPS, DIRA: LIFETIME. RP: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	CAPS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR CAPS. DIRA: 1) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR DIRA, AND 2) TRIAL OF THE PREFERRED AGENT: KINERET. RP: 1) HAD AN EPISODE OF ACUTE PERICARDITIS, 2) SYMPTOM-FREE FOR 4 TO 6 WEEKS, AND 3) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR RP.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

RIMEGEPANT

Products Affected

- NURTEC ODT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	ACUTE MIGRAINE TREATMENT: INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE TRIPTAN (E.G., SUMATRIPTAN, RIZATRIPTAN). INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT. RENEWAL: 1) IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE, OR 2) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS. EPISODIC MIGRAINE PREVENTION: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR MIGRAINE PREVENTION. RENEWAL: REDUCTION IN MIGRAINE OR HEADACHE FREQUENCY, MIGRAINE SEVERITY, OR MIGRAINE DURATION WITH THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

RIOCIGUAT

Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PULMONARY ARTERIAL HYPERTENSION (PAH): DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS. PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH) (WHO GROUP 4): WHO FUNCTIONAL CLASS II-IV SYMPTOMS.
Age Restrictions	
Prescriber Restrictions	INITIAL: PAH, CTEPH: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PAH: NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PHOSPHODIESTERASE (PDE) INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS. CTEPH: 1) NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PDE INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS, AND 2) NOT A CANDIDATE FOR SURGERY OR HAS INOPERABLE CTEPH OR HAS PERSISTENT OR RECURRENT DISEASE AFTER SURGICAL TREATMENT. RENEWAL: PAH, CTEPH: NO CONCURRENT USE WITH NITRATES, NITRIC OXIDE DONORS, PDE INHIBITORS, OR NON-SPECIFIC PDE INHIBITORS.
Indications	All FDA-approved Indications.
Off Label Uses	

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

RIPRETINIB

Products Affected

- QINLOCK

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

RISANKIZUMAB-RZAA

Products Affected

- SKYRIZI INTRAVENOUS 150 MG/ML
- SKYRIZI SUBCUTANEOUS PEN INJECTOR • SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR
- SKYRIZI SUBCUTANEOUS SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PLAQUE PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, FACE, SCALP, OR GENITAL AREA.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSO, PSA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

RITUXIMAB AND HYALURONIDASE HUMAN-SQ

Products Affected

- RITUXAN HYCELA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	FOLLICULAR LYMPHOMA (FL), DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): 1) HAS RECEIVED OR WILL RECEIVE AT LEAST ONE FULL DOSE OF A RITUXIMAB PRODUCT BY INTRAVENOUS INFUSION PRIOR TO INITIATION OF RITUXIMAB AND HYALURONIDASE, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

RITUXIMAB-ABBS

Products Affected

- TRUXIMA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. NON-HODGKINS LYMPHOMA (NHL), CHRONIC LYMPHOCYTIC LEUKEMIA (CLL): PRESCRIBED BY OR IN CONSULTATION WITH AN ONCOLOGIST.
Coverage Duration	RA: INITIAL: 6 MO, RENEWAL: 12 MO. NHL, GPA, MPA, PV: 12 MO. CLL: 6 MO.
Other Criteria	INITIAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. RENEWAL: RA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

ROPEGINTERFERON ALFA-2B-NJFT

Products Affected

- BESREMI

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

RUCAPARIB

Products Affected

- RUBRACA

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

RUXOLITINIB

Products Affected

- JAKAFI
- JAKAFI XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	MYELOFIBROSIS: INITIAL: 6 MONTHS, RENEWAL: 12 MONTHS. POLYCYTHEMIA VERA, GVHD: 12 MONTHS
Other Criteria	INITIAL: CHRONIC GRAFT VS HOST DISEASE (CGVHD): NO CONCURRENT USE WITH REZUROCK, NIKTIMVO, OR IMBRUVICA. RENEWAL: MYELOFIBROSIS: CONTINUES TO BENEFIT FROM THE MEDICATION. CGVHD: NO CONCURRENT USE WITH REZUROCK, NIKTIMVO, OR IMBRUVICA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

SAPROPTERIN

Products Affected

- *javygtor oral tablet, soluble*
- *sapropterin oral tablet, soluble*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 2 MONTHS, RENEWAL 12 MONTHS.
Other Criteria	HYPERPHENYLALANINEMIA (HPA): INITIAL: NO CONCURRENT USE WITH PALYNZIQ. RENEWAL: 1) CONTINUES TO BENEFIT FROM TREATMENT, AND 2) NO CONCURRENT USE WITH PALYNZIQ.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

SECUKINUMAB SQ

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP, OR FACE. NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN (CRP) LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
Age Restrictions	
Prescriber Restrictions	INITIAL: PSO, HIDRADENITIS SUPPURATIVA (HS): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR A DERMATOLOGIST. ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, ENTHESITIS-RELATED ARTHRITIS (ERA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST.
Coverage Duration	INITIAL: HS: 12 MONTHS, ALL OTHER INDICATIONS: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. AS, NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID. ERA: TRIAL OF OR CONTRAINDICATION TO ONE NSAID, SULFASALAZINE, OR METHOTREXATE. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: ALL INDICATIONS: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SELEXIPAG

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	PAH: INITIAL: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING AGENTS: 1) FORMULARY VERSION OF AN ORAL ENDOTHELIN RECEPTOR ANTAGONIST, 2) FORMULARY VERSION OF AN ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR FOR PAH, 3) FORMULARY VERSION OF AN ORAL CGMP STIMULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

SELINEXOR

Products Affected

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

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

SELPERCATINIB

Products Affected

- RETEVMO ORAL CAPSULE 40 MG, 80 MG
- RETEVMO ORAL TABLET 120 MG, 160 MG, 40 MG, 80 MG

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

SELUMETINIB

Products Affected

- KOSELUGO ORAL CAPSULE 10 MG, 25 MG
- KOSELUGO ORAL CAPSULE, SPRINKLE 5 MG, 7.5 MG

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

SEVABERTINIB

Products Affected

- HYRNUO

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

SILDENAFIL TABLET

Products Affected

- *sildenafil (pulm.hypertension) oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: AGES 18 YEARS OR OLDER: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS. AGES 1 TO 17 YEARS: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PAP GREATER THAN 20 MMHG, 2) PCWP OF 15 MMHG OR LESS, AND 3) PVR OF 3 WOOD UNITS OR GREATER.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PAH: INITIAL/RENEWAL: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA) OR ANY ORGANIC NITRATES IN ANY FORM AND 2) NO CONCURRENT USE WITH GUANYLATE CYCLASE STIMULATORS.
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

SIPONIMOD

Products Affected

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

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

SIROLIMUS PROTEIN-BOUND

Products Affected

- FYARRO

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

SODIUM OXYBATE-XYREM

Products Affected

- *sodium oxybate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: CATAPLEXY IN NARCOLEPSY, EXCESSIVE DAYTIME SLEEPINESS (EDS) IN NARCOLEPSY: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST, PSYCHIATRIST, OR SPECIALIST IN SLEEP MEDICINE
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: EDS IN NARCOLEPSY: 1) NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT, AND 2) AGES 18 YEARS OR OLDER: TRIAL, FAILURE OF, OR CONTRAINDICATION TO A FORMULARY VERSION OF MODAFINIL, ARMODAFINIL, OR SUNOSI AND ONE GENERIC STIMULANT INDICATED FOR EDS IN NARCOLEPSY. CATAPLEXY IN NARCOLEPSY: NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT. RENEWAL: CATAPLEXY IN NARCOLEPSY, EDS IN NARCOLEPSY: 1) SUSTAINED IMPROVEMENT OF SYMPTOMS COMPARED TO BASELINE, AND 2) NO CONCURRENT USE WITH A SEDATIVE HYPNOTIC AGENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

SOFOSBUVIR/VELPATASVIR

Products Affected

- EPCLUSA ORAL PELLETS IN PACKET
150-37.5 MG, 200-50 MG
- EPCLUSA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS: AMIODARONE, CARBAMAZEPINE, PHENYTOIN, PHENOBARBITAL, OXCARBAZEPINE, RIFAMPIN, RIFABUTIN, RIFAPENTINE, HIV REGIMEN THAT CONTAINS EFAVIRENZ, ROSUVASTATIN AT DOSES ABOVE 10MG, TIPRANAVIR/RITONAVIR, TOPOTECAN, SOVALDI (AS A SINGLE AGENT), HARVONI, ZEPATIER, MAVYRET, OR VOSEVI, AND 3) PATIENTS WITH DECOMPENSATED CIRRHOSIS REQUIRE CONCURRENT RIBAVIRIN UNLESS RIBAVIRIN INELIGIBLE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR

Products Affected

- VOSEVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	HCV RNA LEVEL WITHIN PAST 6 MONTHS
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE.
Other Criteria	1) CRITERIA WILL BE APPLIED CONSISTENT WITH CURRENT AASLD/IDSA GUIDANCE, 2) NOT CONCURRENTLY TAKING ANY OF THE FOLLOWING MEDICATIONS: 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, TIPRANAVIR/RITONAVIR, SOVALDI (AS A SINGLE AGENT), EPCLUSA, HARVONI, ZEPATIER, OR MAVYRET, AND 3) DOES NOT HAVE MODERATE OR SEVERE HEPATIC IMPAIRMENT (CHILD-PUGH B OR C).
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	No

SOMATROPIN - NORDITROPIN

Products Affected

- NORDITROPIN FLEXPRO

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL/RENEWAL: ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES.
Required Medical Information	INITIAL: PEDIATRIC GROWTH HORMONE DEFICIENCY (GHD), IDIOPATHIC SHORT STATURE (ISS), SMALL FOR GESTATIONAL AGE (SGA), NOONAN SYNDROME: HEIGHT AT LEAST 2 STANDARD DEVIATIONS BELOW THE MEAN HEIGHT FOR CHILDREN OF THE SAME AGE AND GENDER. TURNER SYNDROME (TS): CONFIRMED BY CHROMOSOMAL ANALYSIS (KARYOTYPING). PRADER WILLI SYNDROME (PWS): CONFIRMED GENETIC DIAGNOSIS OF PWS. ADULT GHD: 1) HAS A CONGENITAL, GENETIC, OR ORGANIC DISEASE (E.G., CRANIOPHARYNGIOMA, PITUITARY HYPOPLASIA, ECTOPIC POSTERIOR PITUITARY, PREVIOUS CRANIAL IRRADIATION), OR 2) GHD CONFIRMED BY ONE OF THE FOLLOWING GROWTH HORMONE (GH) STIMULATION TESTS: (A) INSULIN TOLERANCE TEST (PEAK GH OF 5 NG/ML OR LESS), (B) GLUCAGON-STIMULATION TEST (ONE OF THE FOLLOWING: (I) PEAK RESPONSE OF 3 NG/ML OR LESS AND BMI LESS THAN 25 KG/M2, (II) PEAK RESPONSE OF 3 NG/ML OR LESS AND BMI IS BETWEEN 25 - 30 KG/M2 WITH A PRE-TEST PROBABILITY, (III) PEAK RESPONSE OF 1 NG/ML OR LESS AND BMI IS BETWEEN 25 - 30 KG/M2 WITH LOW TEST PROBABILITY, OR (IV) PEAK RESPONSE OF 1 NG/ML OR LESS AND BMI IS GREATER THAN 30 KG/M2), OR (C) MACIMORELIN TEST (PEAK GH OF 2.8 NG/ML OR LESS).
Age Restrictions	SGA: 2 YEARS OF AGE OR OLDER.
Prescriber Restrictions	INITIAL/RENEWAL: ALL INDICATIONS: PRESCRIBED BY OR IN CONSULTATION WITH AN ENDOCRINOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: PEDIATRIC GHD, ISS, SGA, TS, NOONAN SYNDROME: OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF THE WRIST AND HAND. INITIAL/RENEWAL: ADULT GHD, PEDIATRIC GHD, SGA, TS, PWS, NOONAN SYNDROME: NO CONCURRENT USE WITH INCRELEX. RENEWAL: ISS: 1) IMPROVEMENT WHILE ON THERAPY (INCREASED HEIGHT OR INCREASED GROWTH VELOCITY), AND 2) OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF THE WRIST AND HAND. PEDIATRIC GHD, SGA, TS, NOONAN SYNDROME: 1) IMPROVEMENT WHILE ON THERAPY (INCREASED HEIGHT OR INCREASED GROWTH VELOCITY), AND 2) OPEN EPIPHYSES AS CONFIRMED BY RADIOGRAPH OF THE WRIST AND HAND OR HAS NOT COMPLETED PREPUBERTAL GROWTH. PWS: IMPROVEMENT IN BODY COMPOSITION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

SOMATROPIN - SEROSTIM

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	INITIAL/RENEWAL: ATHLETIC ENHANCEMENT, ANTI-AGING PURPOSES
Required Medical Information	INITIAL: HIV/WASTING: ONE OF THE FOLLOWING FOR WEIGHT LOSS: 1) 10% UNINTENTIONAL WEIGHT LOSS OVER 12 MONTHS OR 5% WEIGHT LOSS OVER 6 MONTHS, 2) 5% BODY CELL MASS (BCM) LOSS WITHIN 6 MONTHS, 3) BCM LESS THAN 35% (MEN) OF TOTAL BODY WEIGHT AND BODY MASS INDEX (BMI) LESS THAN 27 KG PER METER SQUARED, 4) BCM LESS THAN 23% (WOMEN) OF TOTAL BODY WEIGHT AND BMI LESS THAN 27 KG PER METER SQUARED, OR 5) BMI LESS THAN 20 KG PER METER SQUARED.
Age Restrictions	
Prescriber Restrictions	HIV/WASTING: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST, NUTRITIONAL SUPPORT SPECIALIST, OR INFECTIOUS DISEASE SPECIALIST.
Coverage Duration	INITIAL/RENEWAL: 9 MONTHS.
Other Criteria	HIV/WASTING: RENEWAL: 1) CLINICAL BENEFIT IN MUSCLE MASS AND WEIGHT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

SONIDEGIB

Products Affected

- ODOMZO

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

SORAFENIB

Products Affected

- *sorafenib*

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

SOTATERCEPT-CSRK

Products Affected

- WINREVAIR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PAH: INITIAL: 1) ON BACKGROUND PAH THERAPY (FOR AT LEAST 3 MONTHS) WITH AT LEAST TWO OF THE FOLLOWING AGENTS FROM DIFFERENT DRUG CLASSES: A) ORAL ENDOTHELIN RECEPTOR ANTAGONIST, B) ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR FOR PAH, C) ORAL CGMP STIMULATOR, D) IV/SQ PROSTACYCLIN, OR 2) ON ONE AGENT FROM ONE OF THE ABOVE DRUG CLASSES, AND HAS A CONTRAINDICATION OR INTOLERANCE TO ALL OF THE OTHER DRUG CLASSES.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

SOTORASIB

Products Affected

- LUMAKRAS ORAL TABLET 120 MG, 240 MG, 320 MG

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

STIRIPENTOL

Products Affected

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

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

SUNITINIB

Products Affected

- *sunitinib malate*

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

TADALAFIL - ADCIRCA, ALYQ

Products Affected

- *alyq*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	PULMONARY ARTERIAL HYPERTENSION (PAH): INITIAL: DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	PAH: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	PAH: INITIAL/RENEWAL: 1) NOT CONCURRENTLY OR INTERMITTENTLY TAKING ORAL ERECTILE DYSFUNCTION AGENTS (E.G. CIALIS, VIAGRA) OR ANY ORGANIC NITRATES IN ANY FORM, AND 2) NO CONCURRENT USE WITH GUANYLATE CYCLASE STIMULATORS.
Indications	All Medically-accepted Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TADALAFIL-CIALIS

Products Affected

- *tadalafil oral tablet 2.5 mg, 5 mg*

PA Criteria	Criteria Details
Exclusion Criteria	ERECTILE DYSFUNCTION WITHOUT DIAGNOSIS OF BENIGN PROSTATIC HYPERPLASIA (BPH).
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	BPH: 1) TRIAL OF ONE ALPHA BLOCKER (E.G., DOXAZOSIN, TERAZOSIN, TAMSULOSIN, ALFUZOSIN), AND 2) TRIAL OF ONE 5-ALPHA-REDUCTASE INHIBITOR (E.G., FINASTERIDE, DUTASTERIDE). APPLIES TO 2.5MG AND 5MG STRENGTHS ONLY
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TAFAMIDIS

Products Affected

- VYNDAMAX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	CARDIOMYOPATHY ASSOCIATED WITH WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM): INITIAL: 1) NEW YORK HEART ASSOCIATION (NYHA) CLASS I, II, OR III HEART FAILURE, AND 2) DIAGNOSIS CONFIRMED BY (A) BONE SCAN (SCINTIGRAPHY) STRONGLY POSITIVE FOR MYOCARDIAL UPTAKE OF TC-99M-PYP, OR (B) BIOPSY OF TISSUE OF AFFECTED ORGAN(S) (CARDIAC AND POSSIBLY NON-CARDIAC SITES) TO CONFIRM AMYLOID PRESENCE AND CHEMICAL TYPING TO CONFIRM PRESENCE OF TRANSTHYRETIN (TTR) PROTEIN.
Age Restrictions	
Prescriber Restrictions	ATTR-CM: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST, ATTR SPECIALIST, OR MEDICAL GENETICIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	ATTR-CM: INITIAL/RENEWAL: NO CONCURRENT USE WITH OTHER ATTR-CM TTR STABILIZERS (E.G., ACORAMIDIS)
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TALAZOPARIB

Products Affected

- TALZENNA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADVANCED OR METASTATIC BREAST CANCER: 1) HAS BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING, AND 2) IF HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER, RECEIVED PRIOR TREATMENT WITH ENDOCRINE THERAPY OR IS CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY. METASTATIC CASTRATION-RESISTANT PROSTATE CANCER: 1) RECEIVED A BILATERAL ORCHIECTOMY, 2) CASTRATE LEVEL OF TESTOSTERONE (I.E., LESS THAN 50 NG/DL), OR 3) CONCURRENT USE WITH A GONADOTROPIN RELEASING HORMONE (GNRH) ANALOG.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No

PA Criteria	Criteria Details
Prerequisite Therapy Required	Yes

TALETRECTINIB

Products Affected

- IBTROZI

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

TALQUETAMAB-TGVS

Products Affected

- TALVEY

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

TARLATAMAB-DLLE

Products Affected

- IMDELLTRA

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

TEBENTAFUSP-TEBN

Products Affected

- KIMMTRAK

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

TECLISTAMAB-CQYV

Products Affected

- TECVAYLI

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

TELISOTUZUMAB VEDOTIN-TLLV

Products Affected

- EMRELIS

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

TELOTRISTAT

Products Affected

- XERMELO

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

TEPOTINIB

Products Affected

- TEPMETKO

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

TERIPARATIDE

Products Affected

- *teriparatide subcutaneous pen injector 20 mcg/dose (560mcg/2.24ml)*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	24 MONTHS
Other Criteria	OSTEOPOROSIS: HAS NOT RECEIVED A TOTAL OF 24 MONTHS CUMULATIVE TREATMENT WITH ANY PARATHYROID HORMONE THERAPY, UNLESS REMAINS AT OR HAS RETURNED TO HAVING A HIGH RISK FOR FRACTURE.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TESTOSTERONE

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	MALE HYPOGONADISM: INITIAL: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TESTOSTERONE CYPIONATE - DEPO

Products Affected

- *testosterone cypionate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	MALE HYPOGONADISM: INITIAL: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TESTOSTERONE ENANTHATE

Products Affected

- *testosterone enanthate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	MALE HYPOGONADISM: INITIAL: CONFIRMED BY: 1) AT LEAST TWO TOTAL SERUM TESTOSTERONE LEVELS OF LESS THAN 300 NG/DL TAKEN ON SEPARATE OCCASIONS, OR 2) FREE SERUM TESTOSTERONE LEVEL OF LESS THAN 5 NG/DL.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL: MALE DELAYED PUBERTY: 6MO, MALE HYPOGONADISM: 12 MO. OTHER INDICATIONS: 12 MO.
Other Criteria	INITIAL: MALE HYPOGONADISM: 1) 40 YEARS OR OLDER: PROSTATE SPECIFIC ANTIGEN (PSA) HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING. RENEWAL: MALE HYPOGONADISM: 1) 40 YEARS OR OLDER: PSA HAS BEEN EVALUATED FOR PROSTATE CANCER SCREENING, AND 2) IMPROVED SYMPTOMS COMPARED TO BASELINE AND TOLERANCE TO TREATMENT. MALE DELAYED PUBERTY: HAS NOT RECEIVED MORE THAN TWO 6-MONTH COURSES OF TESTOSTERONE REPLACEMENT THERAPY
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TETRABENAZINE

Products Affected

- *tetrabenazine*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	HUNTINGTONS DISEASE: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST OR MOVEMENT DISORDER SPECIALIST
Coverage Duration	12 MONTHS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

THALIDOMIDE

Products Affected

- THALOMID ORAL CAPSULE 100 MG, 150 MG, 200 MG, 50 MG

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

TISLELIZUMAB-JSGR

Products Affected

- TEVIMBRA

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

TISOTUMAB VEDOTIN-TFTV

Products Affected

- TIVDAK

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

TIVOZANIB

Products Affected

- FOTIVDA

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

TOCILIZUMAB-AAZG IV

Products Affected

- TYENNE

PA Criteria	Criteria Details
Exclusion Criteria	CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
Coverage Duration	INITIAL: RA, PJIA, SJIA, GCA: 6 MONTHS. CRS: 1 MONTH. RENEWAL: RA, PJIA, SJIA, GCA: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. GIANT CELL ARTERITIS (GCA): 1) HAS COMPLETED, STARTED, OR WILL SOON START A TAPERING COURSE OF GLUCOCORTICOID, AND 2) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: RINVOQ. PJIA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ IR, ORENCIA, RINVOQ. RENEWAL: RA, PJIA, SJIA: 1) CONTINUES TO BENEFIT FROM THE MEDICATION, AND 2) NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION. GCA: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES FOR THE SAME INDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOCILIZUMAB-AAZG SQ

Products Affected

- TYENNE
- TYENNE AUTOINJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST, DERMATOLOGIST, OR IMMUNOLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ, RINVOQ, ORENCIA. GIANT CELL ARTERITIS (GCA): 1) HAS COMPLETED, STARTED, OR WILL SOON START A TAPERING COURSE OF GLUCOCORTICIDS, AND 2) TRIAL OF OR CONTRAINDICATION TO THE PREFERRED AGENT: RINVOQ. PJIA: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING PREFERRED AGENTS: HUMIRA/CYLTEZO/YUFLYMA/HADLIMA, XELJANZ IR, ORENCIA, RINVOQ. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, PJIA, SJIA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.

PA Criteria	Criteria Details
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TOFACITINIB

Products Affected

- *tofacitinib*
- XELJANZ
- XELJANZ XR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), POLYARTICULAR COURSE JUVENILE IDIOPATHIC ARTHRITIS (PCJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AS: TRIAL OF OR CONTRAINDICATION TO AN NSAID. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, PSA, AS, PCJIA: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

TOLVAPTAN

Products Affected

- JYNARQUE ORAL TABLET
- *tolvaptan (polycyst kidney dis) oral tablets, sequential*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD): INITIAL: CONFIRMED POLYCYSTIC KIDNEY DISEASE VIA CT, MRI, OR ULTRASOUND.
Age Restrictions	
Prescriber Restrictions	ADPKD: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A NEPHROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	ADPKD: INITIAL: DOES NOT HAVE ESRD (I.E., RECEIVING DIALYSIS). RENEWAL: HAS NOT PROGRESSED TO ESRD/DIALYSIS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOPICAL TRETINOIN

Products Affected

- ALTRENO
- *tretinoin topical cream*

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

TORIPALIMAB-TPZI

Products Affected

- LOQTORZI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	NASOPHARYNGEAL CARCINOMA (NPC): FIRST LINE TREATMENT: 24 MOS, PREVIOUSLY TREATED: LIFETIME.
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TOVORAFENIB

Products Affected

- OJEMDA ORAL SUSPENSION FOR RECONSTITUTION
- OJEMDA ORAL TABLET

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

TRAMETINIB SOLUTION

Products Affected

- MEKINIST ORAL RECON SOLN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	UNRESECTABLE OR METASTATIC MELANOMA, MELANOMA, METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC), UNRESECTABLE OR METASTATIC SOLID TUMOR, LOW-GRADE GLIOMA (LGG): UNABLE TO SWALLOW MEKINIST TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TRAMETINIB TABLET

Products Affected

- MEKINIST ORAL TABLET 0.5 MG, 2 MG

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

TRASTUZUMAB-DKST

Products Affected

- OGIVRI

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

TRASTUZUMAB-HYALURONIDASE-OYSK

Products Affected

- HERCEPTIN HYLECTA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	ADJUVANT BREAST CANCER, METASTATIC BREAST CANCER: TRIAL OF OR CONTRAINDICATION TO ONE OF THE FOLLOWING PREFERRED AGENTS: HERZUMA, OGIVRI, ONTRUZANT, TRAZIMERA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TRAZODONE

Products Affected

- RALDESY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MAJOR DEPRESSIVE DISORDER (MDD): CONTRAINDICATION TO OR UNABLE TO SWALLOW TRAZODONE TABLETS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TREMELIMUMAB-ACTL

Products Affected

- IMJUDO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	UHCC: 30 DAYS. METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC): 5 MONTHS.
Other Criteria	UNRESECTABLE HEPATOCELLULAR CARCINOMA (UHCC): HAS NOT RECEIVED PRIOR TREATMENT WITH IMJUDO. NSCLC: HAS NOT RECEIVED A TOTAL OF 5 DOSES OF IMJUDO.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TREPROSTINIL-YUTREPIA

Products Affected

- YUTREPIA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PULMONARY ARTERIAL HYPERTENSION (PAH), PULMONARY HYPERTENSION-INTERSTITIAL LUNG DISEASE (PH-ILD): DIAGNOSIS CONFIRMED BY RIGHT HEART CATHETERIZATION WITH THE FOLLOWING PARAMETERS: 1) MEAN PULMONARY ARTERY PRESSURE (PAP) GREATER THAN 20 MMHG, 2) PULMONARY CAPILLARY WEDGE PRESSURE (PCWP) OF 15 MMHG OR LESS, AND 3) PULMONARY VASCULAR RESISTANCE (PVR) GREATER THAN 2 WOOD UNITS.
Age Restrictions	
Prescriber Restrictions	INITIAL: PAH, PH-ILD: PRESCRIBED BY OR IN CONSULTATION WITH A CARDIOLOGIST OR PULMONOLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PAH: TRIAL OF OR CONTRAINDICATION TO TWO OF THE FOLLOWING AGENTS FROM DIFFERENT DRUG CLASSES: 1) ORAL ENDOTHELIN RECEPTOR ANTAGONIST, 2) ORAL PHOSPHODIESTERASE TYPE-5 INHIBITOR FOR PAH, 3) ORAL CGMP STIMULATOR, 4) IV/SQ PROSTACYCLIN.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

TRIENTINE CAPSULE

Products Affected

- *trientine oral capsule 250 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	WILSONS DISEASE: INITIAL: LEIPZIG SCORE OF 4 OR GREATER.
Age Restrictions	
Prescriber Restrictions	WILSONS DISEASE: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A HEPATOLOGIST OR GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 12 MONTHS, RENEWAL: LIFETIME.
Other Criteria	WILSONS DISEASE: INITIAL: TRIAL OF OR CONTRAINDICATION TO FORMULARY VERSION OF PENICILLAMINE TABLET. RENEWAL: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

TRIFLURIDINE/TIPIRACIL

Products Affected

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

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

TRIPTORELIN-TRELSTAR

Products Affected

- TRELSTAR INTRAMUSCULAR
SUSPENSION FOR RECONSTITUTION

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

TUCATINIB

Products Affected

- TUKYSA ORAL TABLET 150 MG, 50 MG

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

UBROGEPANT

Products Affected

- UBRELVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	ACUTE MIGRAINE TREATMENT: INITIAL: 1) TRIAL OF OR CONTRAINDICATION TO ONE TRIPTAN (E.G., SUMATRIPTAN, RIZATRIPTAN), AND 2) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT. RENEWAL: 1) NO CONCURRENT USE WITH OTHER CGRP INHIBITORS FOR ACUTE MIGRAINE TREATMENT, AND 2) ONE OF THE FOLLOWING: (A) IMPROVEMENT FROM BASELINE IN A VALIDATED ACUTE TREATMENT PATIENT-REPORTED OUTCOME QUESTIONNAIRE, OR (B) THERAPY WORKS CONSISTENTLY IN MAJORITY OF MIGRAINE ATTACKS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

UPADACITINIB

Products Affected

- RINVOQ
- RINVOQ LQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS (NR-AXSPA): 1) C-REACTIVE PROTEIN LEVELS ABOVE THE UPPER LIMIT OF NORMAL, OR 2) SACROILIITIS ON MAGNETIC RESONANCE IMAGING (MRI).
Age Restrictions	
Prescriber Restrictions	INITIAL: RHEUMATOID ARTHRITIS (RA), ANKYLOSING SPONDYLITIS (AS), NR-AXSPA, POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST. PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A RHEUMATOLOGIST OR DERMATOLOGIST. AD: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST, ALLERGIST, OR IMMUNOLOGIST. ULCERATIVE COLITIS (UC), CROHNS DISEASE (CD): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.

PA Criteria	Criteria Details
Other Criteria	INITIAL: RA: TRIAL OF OR CONTRAINDICATION TO 3 MONTHS OF TREATMENT WITH ONE CONVENTIONAL SYNTHETIC DMARD (DISEASE-MODIFYING ANTIRHEUMATIC DRUG) - IF A PATIENT TRIED METHOTREXATE, THEN TRIAL AT A DOSE OF AT LEAST 20 MG PER WEEK OR MAXIMALLY TOLERATED DOSE IS REQUIRED. AD: 1) INTRACTABLE PRURITUS OR CRACKING/OOZING/BLEEDING OF AFFECTED SKIN, AND 2) TRIAL OF OR CONTRAINDICATION TO A TOPICAL CORTICOSTEROID, TOPICAL CALCINEURIN INHIBITOR, TOPICAL PDE4 INHIBITOR, OR TOPICAL JAK INHIBITOR. AS, NR-AXSPA: TRIAL OF OR CONTRAINDICATION TO AN NSAID (NON-STEROIDAL ANTI-INFLAMMATORY DRUG). GIANT CELL ARTERITIS (GCA): HAS COMPLETED, STARTED, OR WILL SOON START A TAPERING COURSE OF GLUCOCORTICOIDS. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: RA, PSA, AS, NR-AXSPA, PJIA: CONTINUES TO BENEFIT FROM THE MEDICATION. AD: IMPROVEMENT WHILE ON THERAPY.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

USTEKINUMAB-AAUZ SQ

Products Affected

- *ustekinumab-aauz*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

USTEKINUMAB-AEKN IV

Products Affected

- SELARSDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

USTEKINUMAB-AEKN SQ

Products Affected

- SELARSDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

USTEKINUMAB-KFCE IV

Products Affected

- YESINTEK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

USTEKINUMAB-KFCE SQ

Products Affected

- YESINTEK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	INITIAL: PLAQUE PSORIASIS (PSO): PSORIASIS COVERING 3 PERCENT OR MORE OF BODY SURFACE AREA OR PSORIATIC LESIONS AFFECTING THE HANDS, FEET, GENITAL AREA, SCALP OR FACE.
Age Restrictions	
Prescriber Restrictions	INITIAL: PSORIATIC ARTHRITIS (PSA): PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST OR RHEUMATOLOGIST. PSO: PRESCRIBED BY OR IN CONSULTATION WITH A DERMATOLOGIST. CROHNS DISEASE (CD), ULCERATIVE COLITIS (UC): PRESCRIBED BY OR IN CONSULTATION WITH A GASTROENTEROLOGIST.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: 12 MONTHS.
Other Criteria	INITIAL: PSO: 1) AT LEAST A 3 MONTH TRIAL OF ONE ORAL IMMUNOSUPPRESSANT (CYCLOSPORINE, METHOTREXATE, TACROLIMUS) OR PUVA (PHOTOTHERAPY), 2) CONTRAINDICATION OR INTOLERANCE TO BOTH IMMUNOSUPPRESSANT AND PUVA, OR 3) PATIENT IS SWITCHING FROM A DIFFERENT BIOLOGIC, PDE-4 INHIBITOR, OR JAK INHIBITOR FOR THE SAME INDICATION. INITIAL/RENEWAL: ALL INDICATIONS: NO CONCURRENT USE WITH ANOTHER SYSTEMIC BIOLOGIC OR TARGETED SMALL MOLECULES (E.G., JAK INHIBITOR, PDE-4 INHIBITOR) FOR THE SAME INDICATION. RENEWAL: PSA, PSO: CONTINUES TO BENEFIT FROM THE MEDICATION.
Indications	All FDA-approved Indications.
Off Label Uses	

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

VALBENAZINE

Products Affected

- INGREZZA
- INGREZZA INITIATION PK(TARDIV)
- INGREZZA SPRINKLE

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

VANDETANIB

Products Affected

- CAPRELSA ORAL TABLET 100 MG, 300 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	CURRENTLY STABLE ON CAPRELSA REQUIRES NO EXTRA CRITERIA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

VANZACAFTOR-TEZACAFTOR- DEUTIVACAFTOR

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	CF: INITIAL: PRESCRIBED BY OR IN CONSULTATION WITH A PULMONOLOGIST OR CYSTIC FIBROSIS EXPERT.
Coverage Duration	INITIAL: 6 MONTHS. RENEWAL: LIFETIME.
Other Criteria	CF: INITIAL: NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR. RENEWAL: 1) IMPROVEMENT IN CLINICAL STATUS, AND 2) NO CONCURRENT USE WITH ANOTHER CFTR MODULATOR.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

VEMURAFENIB

Products Affected

- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MELANOMA: ZELBORAF WILL BE USED ALONE OR IN COMBINATION WITH COTELLIC
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

VENETOCLAX

Products Affected

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

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

VERICIGUAT

Products Affected

- VERQUVO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL/RENEWAL:12 MONTHS.
Other Criteria	HEART FAILURE (HF): INITIAL: 1) TRIAL OF, CONTRAINDICATION, OR INTOLERANCE TO ONE PREFERRED SGLT-2 INHIBITOR, AND 2) TRIAL OF, CONTRAINDICATION, OR INTOLERANCE TO ONE AGENT FROM ANY OF THE FOLLOWING STANDARD OF CARE CLASSES: (A) ACE INHIBITOR, ARB, OR ARNI, (B) BETA BLOCKER (BISOPROLOL, CARVEDILOL, METOPROLOL SUCCINATE), OR (C) ALDOSTERONE ANTAGONIST (SPIRONOLACTONE, EPLERENONE). INITIAL/RENEWAL: NO CONCURRENT USE WITH RIOCIGUAT OR PDE-5 INHIBITORS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VIGABATRIN

Products Affected

- *vigabatrin*
- *vigadrone*
- *vigpoder*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	INITIAL: REFRACTORY COMPLEX PARTIAL SEIZURES (CPS), INFANTILE SPASMS: PRESCRIBED BY OR IN CONSULTATION WITH A NEUROLOGIST.
Coverage Duration	INITIAL/RENEWAL: 12 MONTHS
Other Criteria	INITIAL: CPS: TRIAL OF OR CONTRAINDICATION TO TWO ANTIEPILEPTIC AGENTS.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VIMSELTINIB

Products Affected

- ROMVIMZA

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

VISMODEGIB

Products Affected

- ERIVEDGE

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

VONOPRAZAN

Products Affected

- VOQUEZNA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	INITIAL: H PYLORI: 30 DAYS. EE: 8 WEEKS. NERD: 4 WEEKS. RENEWAL: EE: 24 WEEKS.
Other Criteria	INITIAL: EROSION ESOPHAGITIS (EE): TRIAL OF OR CONTRAINDICATION TO TWO PROTON PUMP INHIBITORS AT MAXIMUM DOSE FOR 8 WEEKS EACH. NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE (NERD): 1) NO PREVIOUS TREATMENT FAILURE WITH VOQUEZNA IN THE LAST 12 MONTHS, AND 2) TRIAL OF OR CONTRAINDICATION TO ONE PROTON PUMP INHIBITOR AT MAXIMUM DOSE FOR 8 WEEKS. RENEWAL: EE: MAINTAINED A CLINICAL RESPONSE ON VOQUEZNA.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

VORASIDENIB

Products Affected

- VORANIGO

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

VORICONAZOLE SUSPENSION

Products Affected

- *voriconazole oral suspension for reconstitution*

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

ZANIDATAMAB-HRII

Products Affected

- ZIIHERA

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

ZANUBRUTINIB

Products Affected

- BRUKINSA ORAL CAPSULE
- BRUKINSA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 MONTHS
Other Criteria	MANTLE CELL LYMPHOMA: INTOLERANCE TO CALQUENCE. CHRONIC LYMPHOCYTIC LEUKEMIA, SMALL LYMPHOCYTIC LYMPHOMA: INTOLERANCE TO CALQUENCE OR IMBRUVICA. WALDENSTROMS MACROGLOBULINEMIA: NO STEP REQUIRED.
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	Yes

ZENOCUTUZUMAB-ZBCO

Products Affected

- BIZENGRI

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

ZIFTOMENIB

Products Affected

- KOMZIFTI

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

ZOLBETUXIMAB-CLZB

Products Affected

- VYLOY

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

ZONGERTINIB

Products Affected

- HERNEXEOS

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

ZURANOLONE

Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	14 DAYS
Other Criteria	
Indications	All FDA-approved Indications.
Off Label Uses	
Part B Prerequisite	No
Prerequisite Therapy Required	No

INDEX

1ST TIER UNIFINE PENTP 5MM 31G...	179	ASSURE ID DUO PRO NDL 31G 5MM..	179
1ST TIER UNIFINE PNTIP 4MM 32G	179	ASSURE ID DUO-SHIELD 30GX3/16" ...	179
1ST TIER UNIFINE PNTIP 6MM 31G	179	ASSURE ID DUO-SHIELD 30GX5/16" ...	179
1ST TIER UNIFINE PNTIP 8MM 31G		ASSURE ID INSULIN SAFETY	
STRL,SINGLE-USE,SHRT	179	SYRINGE 1 ML 29 GAUGE X 1/2"	179
1ST TIER UNIFINE PNTIP 29GX1/2"	179	ASSURE ID PEN NEEDLE 30GX5/16" ...	179
1ST TIER UNIFINE PNTIP 31GX3/16	179	ASSURE ID PRO PEN NDL 30G 5MM...	179
1ST TIER UNIFINE PNTIP 32GX5/32	179	ASSURE ID SYR 0.5 ML 31GX15/64"	179
<i>abigale</i>	157	ASSURE ID SYR 1 ML 31GX15/64"	179
<i>abiraterone</i>	7	ATTRUBY	10
<i>abiraterone, submicronized</i>	8	AUGTYRO ORAL CAPSULE 160 MG,	
<i>abirtega</i>	7	40 MG	294
ACTIMMUNE	192	AUSTEDO ORAL TABLET 12 MG, 6	
<i>adalimumab-aaty</i>	14	MG, 9 MG	88
<i>adalimumab-aaty(cf) ai crohns</i>	14	AUSTEDO XR ORAL TABLET	
ADEMPAS	305	EXTENDED RELEASE 24 HR 12 MG, 18	
ADVOCATE INS 0.3 ML 30GX5/16"	179	MG, 24 MG, 30 MG, 36 MG, 42 MG, 48	
ADVOCATE INS 0.3 ML 31GX5/16"	179	MG, 6 MG	88
ADVOCATE INS 0.5 ML 30GX5/16"	179	AUSTEDO XR TITRATION KT(WK1-4)..	88
ADVOCATE INS 0.5 ML 31GX5/16"	179	AVMAPKI	40
ADVOCATE INS 1 ML 31GX5/16"	179	AVMAPKI-FAKZYNJA	40
ADVOCATE INS SYR 1 ML 30GX5/16..	179	AVONEX INTRAMUSCULAR PEN	
ADVOCATE PEN NEEDLE 32G 4MM...	179	INJECTOR KIT	189
ADVOCATE PEN NEEDLE 4MM 33G...	179	AVONEX INTRAMUSCULAR	
ADVOCATE PEN NEEDLES 5MM 31G.	179	SYRINGE KIT	189
ADVOCATE PEN NEEDLES 8MM 31G.	179	AYVAKIT	39
AIMOVIG AUTOINJECTOR	121	BALVERSA ORAL TABLET 3 MG, 4	
AKEEGA	251	MG, 5 MG	120
ALCOHOL PADS	179	BD AUTOSHIELD DUO NDL	
ALECENSA	21	5MMX30G	179
ALTRENO	373	BD ECLIPSE 30GX1/2" SYRINGE	179
ALUNBRIG ORAL TABLET 180 MG, 30		BD INS SYR 0.3 ML 8MMX31G(1/2)	179
MG, 90 MG	58	BD INS SYR UF 0.3 ML 12.7MMX30G..	179
ALUNBRIG ORAL TABLETS,DOSE		BD INS SYR UF 0.5 ML 12.7MMX30G	
PACK	58	NOT FOR RETAIL SALE	179
ALVAIZ	108	BD INSULIN SYR 1 ML 27GX12.7MM..	179
ALYFTREK ORAL TABLET 10-50-125		BD INSULIN SYR 1 ML 27GX5/8"	
MG, 4-20-50 MG	402	MICRO-FINE	179
<i>alyq</i>	344	BD SAFETGLD INS 0.3 ML 29G 13MM.	179
ANKTIVA	257	BD SAFETYGLD INS 0.3 ML 31G 8MM	179
AQINJECT PEN NEEDLE 31G 5MM	179	BD SAFETYGLD INS 0.5 ML 30G 8MM	179
AQINJECT PEN NEEDLE 32G 4MM	179	BD SAFETYGLD INS 1 ML 29G 13MM.	179
ARCALYST	301	BD SAFETYGLD INS 1 ML 6MMX31G	179
ARIKAYCE	23	BD SAFETYGLIDE SYRINGE 27GX5/8	179
<i>armodafinil</i>	241	BD SAFTYGLD INS 0.3 ML 6MMX31G	179

BD SAFTYGLD INS 0.5 ML 29G 13MM	179	CAMCEVI (6 MONTH)	209
BD SAFTYGLD INS 0.5 ML 6MMX31G	179	CAMZYOS	227
BD SINGLE USE SWAB	179	CAPRELSA ORAL TABLET 100 MG, 300 MG	401
BD UF MICRO PEN NEEDLE 6MMX32G	179	CAREFINE PEN NEEDLE 12.7MM 29G HRI	179
BD UF MINI PEN NEEDLE 5MMX31G	179	CAREFINE PEN NEEDLE 4MM 32G	179
BD UF NANO PEN NEEDLE 4MMX32G	179	CAREFINE PEN NEEDLE 5MM 32G	179
BD UF ORIG PEN NDL 12.7MMX29G	179	CAREFINE PEN NEEDLE 6MM 31G HRI	179
BD UF SHORT PEN NEEDLE 8MMX31G	179	CAREFINE PEN NEEDLE 8MM 30G HRI	179
BD VEO INS 0.3 ML 6MMX31G (1/2)	179	CAREFINE PEN NEEDLES 6MM 32G	179
BD VEO INS SYRING 1 ML 6MMX31G	179	CAREFINE PEN NEEDLES 8MM 31G	179
BD VEO INS SYRN 0.3 ML 6MMX31G	179	CARETOUCH ALCOHOL 70% PREP PAD	179
BD VEO INS SYRN 0.5 ML 6MMX31G	179	CARETOUCH PEN NEEDLE 29G 12MM	179
<i>bendamustine intravenous recon soln</i>	49	CARETOUCH PEN NEEDLE 31GX1/4"	179
BENDAMUSTINE INTRAVENOUS SOLUTION	49	CARETOUCH PEN NEEDLE 31GX3/16"	179
BENDEKA	49	CARETOUCH PEN NEEDLE 31GX5/16"	179
BENLYSTA SUBCUTANEOUS	46	CARETOUCH PEN NEEDLE 32GX3/16"	179
BESREMI	313	CARETOUCH PEN NEEDLE 32GX5/32"	179
<i>betaine</i>	51	CARETOUCH SYR 0.3 ML 31GX5/16"	179
BETASERON SUBCUTANEOUS KIT	190	CARETOUCH SYR 0.5 ML 30GX5/16"	179
<i>bexarotene</i>	53	CARETOUCH SYR 0.5 ML 31GX5/16"	179
BIZENGRI	414	CARETOUCH SYR 1 ML 28GX5/16"	179
BORDERED GAUZE 2"X2"	179	CARETOUCH SYR 1 ML 29GX5/16"	179
<i>bortezomib injection</i>	55	CARETOUCH SYR 1 ML 30GX5/16"	179
BORUZU	55	CARETOUCH SYR 1 ML 31GX5/16"	179
<i>bosentan oral tablet</i>	56	<i>carglumic acid</i>	65
BOSULIF ORAL CAPSULE 100 MG, 50 MG	57	CAYSTON	44
BOSULIF ORAL TABLET 100 MG, 400 MG, 500 MG	57	CIMZIA	67
<i>bosutinib oral tablet 100 mg, 400 mg, 500 mg</i>	57	CIMZIA POWDER FOR RECONST	67
BRAFTOVI	112	CIMZIA STARTER KIT	67
BRUKINSA ORAL CAPSULE	413	<i>cladribine 10 mg x 10 tab pk outer</i>	70
BRUKINSA ORAL TABLET	413	<i>cladribine(multiple sclerosis) oral tablet 10 mg (10 pack), 10 mg (4 pack), 10 mg (5 pack), 10 mg (6 pack), 10 mg (7 pack), 10 mg (8 pack), 10 mg (9 pack)</i>	70
<i>butalbital-acetaminop-caf-cod oral capsule 50-325-40-30 mg</i>	155	CLICKFINE PEN NEEDLE 32GX5/32"	179
<i>butalbital-acetaminophen-caff oral capsule</i>	155	32GX4MM, STERILE	179
<i>butalbital-acetaminophen-caff oral tablet</i>	155		
CABOMETYX ORAL TABLET 20 MG, 40 MG, 60 MG	61		
CALQUENCE (ACALABRUTINIB MAL)	9		

CLICKFINE UNIVERSAL 31G X 1/4"	
6MM, STORE BRAND.....	179
COMETRIQ ORAL CAPSULE 100	
MG/DAY(80 MG X1-20 MG X1), 140	
MG/DAY(80 MG X1-20 MG X3), 60	
MG/DAY (20 MG X 3/DAY).....	60
COMFORT EZ 0.3 ML 31G 15/64"	179
COMFORT EZ 0.5 ML 31G 15/64"	179
COMFORT EZ INS 0.3 ML 30GX1/2"	179
COMFORT EZ INS 0.3 ML 30GX5/16" ...	179
COMFORT EZ INS 1 ML 31G 15/64"	179
COMFORT EZ INS 1 ML 31GX5/16"	179
COMFORT EZ INSULIN SYR 0.3 ML	179
COMFORT EZ INSULIN SYR 0.5 ML	179
COMFORT EZ PEN NEEDLE 12MM	
29G.....	179
COMFORT EZ PEN NEEDLES 4MM	
32G SINGLE USE, MICRO.....	179
COMFORT EZ PEN NEEDLES 4MM	
33G.....	179
COMFORT EZ PEN NEEDLES 5MM	
31G MINI.....	179
COMFORT EZ PEN NEEDLES 5MM	
32G SINGLE USE,MINI,HRI.....	179
COMFORT EZ PEN NEEDLES 5MM	
33G.....	179
COMFORT EZ PEN NEEDLES 6MM	
31G.....	179
COMFORT EZ PEN NEEDLES 6MM	
32G.....	179
COMFORT EZ PEN NEEDLES 6MM	
33G.....	179
COMFORT EZ PEN NEEDLES 8MM	
31G SHORT.....	179
COMFORT EZ PEN NEEDLES 8MM	
32G.....	179
COMFORT EZ PEN NEEDLES 8MM	
33G.....	179
COMFORT EZ PRO PEN NDL 30G 8MM	
.....	179
COMFORT EZ PRO PEN NDL 31G 4MM	
.....	179
COMFORT EZ PRO PEN NDL 31G 5MM	
.....	179
COMFORT EZ SYR 0.3 ML 29GX1/2"	179
COMFORT EZ SYR 0.5 ML 28GX1/2"	179
COMFORT EZ SYR 0.5 ML 29GX1/2"	179
COMFORT EZ SYR 0.5 ML 30GX1/2"	179
COMFORT EZ SYR 1 ML 27G 12.7MM.	179
COMFORT EZ SYR 1 ML 28GX1/2"	179
COMFORT EZ SYR 1 ML 29GX1/2"	179
COMFORT EZ SYR 1 ML 30GX1/2"	179
COMFORT EZ SYR 1 ML 30GX5/16"	179
COMFORT POINT PEN NDL 31GX1/3".	179
COMFORT POINT PEN NDL 31GX1/6".	179
COMFORT TOUCH PEN NDL 31G 4MM	
.....	179
COMFORT TOUCH PEN NDL 31G 5MM	
.....	179
COMFORT TOUCH PEN NDL 31G 6MM	
.....	179
COMFORT TOUCH PEN NDL 31G 8MM	
.....	179
COMFORT TOUCH PEN NDL 32G 4MM	
.....	179
COMFORT TOUCH PEN NDL 32G 5MM	
.....	179
COMFORT TOUCH PEN NDL 32G 6MM	
.....	179
COMFORT TOUCH PEN NDL 32G 8MM	
.....	179
COMFORT TOUCH PEN NDL 33G 4MM	
.....	179
COMFORT TOUCH PEN NDL 33G 6MM	
.....	179
COMFORT TOUCH PEN NDL	
33GX5MM.....	179
COPIKTRA.....	100
CORTROPHIN GEL INJECTION.....	73
COSENTYX (2 SYRINGES).....	317
COSENTYX PEN (2 PENS).....	317
COSENTYX SUBCUTANEOUS	
SYRINGE 75 MG/0.5 ML.....	317
COSENTYX UNOREADY PEN.....	317
COTELLIC.....	72
CRESEMBA ORAL.....	194
CURITY ALCOHOL PREPS 2	
PLY,MEDIUM.....	179
CURITY GAUZE PADS.....	179
CURITY GAUZE SPONGES (12 PLY)-	
200/BAG.....	179
CYLTEZO(CF).....	16
CYLTEZO(CF) PEN.....	16
CYLTEZO(CF) PEN CROHN'S-UC-HS.....	16

CYLTEZO(CF) PEN PSORIASIS-UV	16	DROPLET INS 0.5 ML 30G 12.7MM	
<i>dalfampridine</i>	80	OUTER	179
DANYELZA	242	DROPLET INS 0.5 ML 30GX6MM(1/2) ..	179
DANZITEN	247	DROPLET INS 0.5 ML 30GX8MM(1/2) ..	179
<i>dasatinib oral tablet 100 mg, 140 mg, 20</i>		DROPLET INS 0.5 ML 31GX6MM(1/2) ..	179
<i>mg, 50 mg, 70 mg, 80 mg</i>	82	DROPLET INS 0.5 ML 31GX8MM(1/2) ..	179
DATROWAY	83	DROPLET INS SYR 0.3 ML 30GX6MM.	179
DAURISMO ORAL TABLET 100 MG, 25		DROPLET INS SYR 0.3 ML 30GX8MM.	179
MG	146	DROPLET INS SYR 0.3 ML 31GX6MM.	179
<i>deferasirox oral granules in packet</i>	85	DROPLET INS SYR 0.3 ML 31GX8MM.	179
<i>deferasirox oral tablet</i>	85	DROPLET INS SYR 0.5 ML 30G 8MM	
DERMACEA 2"X2" GAUZE 12 PLY,		OUTER	179
USP TYPE VII	179	DROPLET INS SYR 0.5 ML 31G 6MM	
DERMACEA GAUZE 2"X2" SPONGE 8		OUTER	179
PLY	179	DROPLET INS SYR 0.5 ML 31G 8MM	
DERMACEA NON-WOVEN 2"X2"		OUTER	179
SPNGE	179	DROPLET INS SYR 1 ML 29G 12.7MM	
<i>dermacinrx lidocan 5% patch outer</i>	217	OUTER	179
DIACOMIT ORAL CAPSULE 250 MG,		DROPLET INS SYR 1 ML 30G 12.5MM.	179
500 MG	342	DROPLET INS SYR 1 ML 30G 6MM	179
DIACOMIT ORAL POWDER IN		DROPLET INS SYR 1 ML 30G 8MM	
PACKET 250 MG, 500 MG	342	OUTER	179
<i>diclofenac epolamine</i>	90	DROPLET INS SYR 1 ML 31G 6MM	
<i>diclofenac sodium topical solution in</i>		OUTER	179
<i>metered-dose pump</i>	89	DROPLET INS SYR 1 ML 31G 8MM	179
<i>dimethyl fumarate oral capsule, delayed</i>		DROPLET MICRON 34G X 9/64"	179
<i>release(dr/ec) 120 mg, 120 mg (14)- 240</i>		DROPLET PEN NEEDLE 29G 10MM	179
<i>mg (46), 240 mg</i>	91	DROPLET PEN NEEDLE 29G 12MM	179
<i>diphenoxylate-atropine oral tablet</i>	165	DROPLET PEN NEEDLE 30G 8MM	179
<i>dipyridamole oral</i>	156	DROPLET PEN NEEDLE 31G 5MM	179
<i>dronabinol oral capsule</i>	95	DROPLET PEN NEEDLE 31G 6MM	179
DROPLET 0.3 ML 29G 12.7MM(1/2)		DROPLET PEN NEEDLE 31G 8MM	179
OUTER	179	DROPLET PEN NEEDLE 32G 4MM	179
DROPLET 0.3 ML 30G 12.7MM(1/2)		DROPLET PEN NEEDLE 32G 5MM	179
OUTER	179	DROPLET PEN NEEDLE 32G 6MM	179
DROPLET 0.5 ML 29GX12.5MM(1/2)	179	DROPLET PEN NEEDLE 32G 8MM	179
DROPLET 0.5 ML 30GX12.5MM(1/2)	179	DROPSAFE ALCOHOL 70% PREP	
DROPLET INS 0.3 ML 29GX12.5MM	179	PADS	179
DROPLET INS 0.3 ML 30G 8MM(1/2)		DROPSAFE INS SYR 0.3 ML 31G 6MM	179
OUTER	179	DROPSAFE INS SYR 0.3 ML 31G 8MM	179
DROPLET INS 0.3 ML 30GX12.5MM	179	DROPSAFE INS SYR 0.5 ML 31G 6MM	179
DROPLET INS 0.3 ML 31G 6MM(1/2)		DROPSAFE INS SYR 0.5 ML 31G 8MM	179
OUTER	179	DROPSAFE INSUL SYR 1 ML 31G	
DROPLET INS 0.3 ML 31G 8MM(1/2)		6MM	179
OUTER	179	DROPSAFE INSUL SYR 1 ML 31G	
DROPLET INS 0.5 ML 29G 12.7MM		8MM	179
OUTER	179		

DROPSAFE INSULN 1 ML 29G 12.5MM	EASY TOUCH 1 ML SYR 29GX1/2"	179
.....	EASY TOUCH 1 ML SYR 30GX1/2"	179
DROPSAFE PEN NEEDLE 31G 4MM.....	EASY TOUCH ALCOHOL 70% PADS	
DROPSAFE PEN NEEDLE 31G 5MM.....	GAMMA-STERILIZED.....	179
DROPSAFE PEN NEEDLE 31G 8MM.....	EASY TOUCH AUTO 0.5 ML 30G 6MM	179
DROPSAFE PEN NEEDLE 31GX1/4"	EASY TOUCH AUTO 0.5 ML 30G 8MM	179
<i>droxidopa</i>	EASY TOUCH AUTORET 1 ML 30G	
DUPIXENT PEN	6MM.....	179
DUPIXENT SYRINGE.....	EASY TOUCH AUTORET 1 ML 30G	
EASY CMFT SFTY PEN NDL 31G 5MM	8MM.....	179
EASY CMFT SFTY PEN NDL 31G 6MM	EASY TOUCH FLIPLOK 1 ML 27GX0.5	179
EASY CMFT SFTY PEN NDL 32G 4MM	EASY TOUCH INS 0.3 ML 30G 8MM.....	179
EASY COMFORT 0.3 ML 31G 1/2"	EASY TOUCH INS 0.3 ML 31G 8MM.....	179
EASY COMFORT 0.3 ML 31G 5/16"	EASY TOUCH INS 0.5 ML 30G 8MM.....	179
EASY COMFORT 0.5 ML 30GX1/2"	EASY TOUCH INS 0.5 ML 31G 6MM.....	179
EASY COMFORT 0.5 ML 31GX5/16"	EASY TOUCH INS 0.5 ML 31G 8MM.....	179
EASY COMFORT 0.5 ML 32GX5/16"	EASY TOUCH INS 1 ML 27G 1/2"	179
EASY COMFORT 0.5 ML SYRINGE.....	EASY TOUCH INS 1 ML 28G 12.7MM...	179
EASY COMFORT 1 ML 31GX5/16"	EASY TOUCH INS SYR 1 ML 30G 8MM	
EASY COMFORT 1 ML 32GX5/16"		179
EASY COMFORT ALCOHOL 70% PAD	EASY TOUCH INS SYR 1 ML 31G 6MM	
EASY COMFORT INSULIN 1 ML SYR..		179
EASY COMFORT PEN NDL 29G 4MM..	EASY TOUCH INS SYR 1 ML 31G 8MM	
EASY COMFORT PEN NDL 29G 5MM..		179
EASY COMFORT PEN NDL 31GX1/4" ..	EASY TOUCH INSULIN 1 ML 29GX1/2	179
EASY COMFORT PEN NDL 31GX3/16" ..	EASY TOUCH INSULIN 1 ML 30GX1/2	179
EASY COMFORT PEN NDL 31GX5/16" ..	EASY TOUCH INSULIN SYR 1 ML.....	179
EASY COMFORT PEN NDL 32GX5/32" ..	EASY TOUCH INSULIN SYR 1 ML	
EASY COMFORT PEN NDL 33G 4MM..	RETRACTABLE.....	179
EASY COMFORT PEN NDL 33G 5MM..	EASY TOUCH INSULN 1 ML 29GX1/2" ..	179
EASY COMFORT PEN NDL 33G 6MM..	EASY TOUCH INSULN 1 ML 30GX1/2" ..	179
EASY COMFORT SYR 0.5 ML 29G	EASY TOUCH INSULN 1 ML 30GX5/16	179
8MM.....	EASY TOUCH INSULN 1 ML 31GX5/16	179
EASY COMFORT SYR 1 ML 29G 8MM..	EASY TOUCH LUER LOK INSUL 1 ML	179
EASY COMFORT SYR 1 ML 30GX1/2" ..	EASY TOUCH PEN NEEDLE 29GX1/2" ..	179
EASY GLIDE INS 0.3 ML 31GX6MM....	EASY TOUCH PEN NEEDLE 30GX5/16	179
EASY GLIDE INS 0.5 ML 31GX6MM....	EASY TOUCH PEN NEEDLE 31GX1/4" ..	179
EASY GLIDE INS 1 ML 31GX6MM.....	EASY TOUCH PEN NEEDLE 31GX3/16	179
EASY GLIDE PEN NEEDLE 4MM 33G..	EASY TOUCH PEN NEEDLE 31GX5/16	179
EASY TOUCH 0.3 ML 30G 8MM(1/2)....	EASY TOUCH PEN NEEDLE 32GX1/4" ..	179
EASY TOUCH 0.3 ML 31G 6MM(1/2)....	EASY TOUCH PEN NEEDLE 32GX3/16	179
EASY TOUCH 0.3 ML 31G 8MM(1/2)....	EASY TOUCH PEN NEEDLE 32GX5/32	179
EASY TOUCH 0.3 ML SYR 30GX1/2"	EASY TOUCH SAF PEN NDL 29G 5MM	
EASY TOUCH 0.5 ML SYR 27GX1/2"		179
EASY TOUCH 0.5 ML SYR 29GX1/2"	EASY TOUCH SAF PEN NDL 29G 8MM	
EASY TOUCH 0.5 ML SYR 30GX1/2"		179
EASY TOUCH 0.5 ML SYR 30GX5/16... ..		

EASY TOUCH SAF PEN NDL 30G 5MM.....	179	EMBRACE PEN NEEDLE 31G 5MM.....	179
EASY TOUCH SAF PEN NDL 30G 8MM.....	179	EMBRACE PEN NEEDLE 31G 6MM.....	179
EASY TOUCH SYR 0.5 ML 28G 12.7MM.....	179	EMBRACE PEN NEEDLE 31G 8MM.....	179
EASY TOUCH SYR 0.5 ML 29G 12.7MM.....	179	EMBRACE PEN NEEDLE 32G 4MM.....	179
EASY TOUCH SYR 1 ML 27G 16MM.....	179	EMGALITY PEN.....	141
EASY TOUCH UNI-SLIP SYR 1 ML.....	179	EMGALITY SYRINGE	
EASYLIFE ALCOHOL 70% PADS.....	179	SUBCUTANEOUS SYRINGE 120	
EASYLIFE INS PEN NDL 29G 12MM.....	179	MG/ML, 300 MG/3 ML (100 MG/ML X 3).....	141
EASYLIFE INS PEN NDL 31G 4MM.....	179	EMRELIS.....	354
EASYLIFE INS PEN NDL 31G 5MM.....	179	ENBREL MINI.....	124
EASYLIFE INS PEN NDL 31G 6MM.....	179	ENBREL SUBCUTANEOUS SOLUTION.....	124
EASYLIFE INS PEN NDL 31G 8MM.....	179	ENBREL SUBCUTANEOUS SYRINGE.....	124
EASYLIFE INS PEN NDL 32G 4MM.....	179	ENBREL SURECLICK.....	124
EASYLIFE INS PEN NDL 32G 5MM.....	179	ENSACOVE ORAL CAPSULE 100 MG, 25 MG.....	113
EASYLIFE INS PEN NDL 32G 6MM.....	179	EPCLUSA ORAL PELLETS IN PACKET 150-37.5 MG, 200-50 MG.....	330
EASYLIFE INS PEN NDL 32G 8MM.....	179	EPCLUSA ORAL TABLET.....	330
EASYLIFE INS PEN NDL 33G 4MM.....	179	EPIDIOLEX.....	62
EASYLIFE INS PEN NDL 33G 5MM.....	179	EPKINLY.....	117
EASYLIFE INS PEN NDL 33G 6MM.....	179	ERBITUX.....	69
EASYLIFE INS PEN NDL 33G 8MM.....	179	ERIVEDGE.....	408
EASYLIFE INS SYR 0.5 ML 30G 8MM.....	179	ERLEADA ORAL TABLET 240 MG, 60 MG.....	28
EASYLIFE INS SYR 1 ML 30G 8MM.....	179	<i>erlotinib oral tablet 100 mg, 150 mg, 25 mg.....</i>	122
EASYLIFE INS SYR 1 ML 31G 8MM.....	179	<i>estradiol-norethindrone acet.....</i>	157
EASYLIFE SAFTY PEN NDL 31G 4MM.....	179	<i>everolimus (antineoplastic) oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg.....</i>	126
EASYLIFE SAFTY PEN NDL 31G 5MM.....	179	<i>everolimus (antineoplastic) oral tablet for suspension.....</i>	127
EASYLIFE SYR 1 ML 30G 12.7MM.....	179	EXEL U100 0.3 ML 29GX1/2".....	179
EASYTOUCH SAF PEN NDL 30G 6MM.....	179	EXEL U100 0.3 ML 30GX5/16".....	179
ELAHERE.....	238	EXXUA ORAL TABLET EXTENDED RELEASE 24 HR.....	144
ELIGARD.....	210	EXXUA ORAL TABLET, EXT REL 24HR DOSE PACK.....	144
ELIGARD (3 MONTH).....	210	FAKZYNJA.....	40
ELIGARD (4 MONTH).....	210	FASENRA.....	50
ELIGARD (6 MONTH).....	210	FASENRA PEN.....	50
ELREXFIO 44 MG/1.1 ML VIAL INNER, SUV, P/F.....	107	<i>fentanyl citrate buccal lozenge on a handle.....</i>	131
ELREXFIO SUBCUTANEOUS SOLUTION 40 MG/ML.....	107	<i> fingolimod.....</i>	137
<i>eltrombopag olamine oral powder in packet 12.5 mg, 25 mg.....</i>	109	FINTEPLA.....	130
<i>eltrombopag olamine oral tablet 12.5 mg, 25 mg, 50 mg, 75 mg.....</i>	109	FOTIVDA.....	365
EMBRACE PEN NEEDLE 29G 12MM.....	179		
EMBRACE PEN NEEDLE 30G 5MM.....	179		
EMBRACE PEN NEEDLE 30G 8MM.....	179		

FRUZAQLA ORAL CAPSULE 1 MG, 5 MG.....	139	HEALTHWISE INS 1 ML 30GX5/16".....	179
FT STERILE PADS 2" X 2"	179	HEALTHWISE INS 1 ML 31GX5/16".....	179
FYARRO	327	HEALTHWISE PEN NEEDLE 31G 5MM	179
GAUZE PAD TOPICAL BANDAGE 2 X 2 ".....	179	HEALTHWISE PEN NEEDLE 31G 8MM	179
GAVRETO.....	288	HEALTHWISE PEN NEEDLE 32G 4MM	179
<i>gefitinib</i>	143	HEALTHY ACCENTS PENTIP 5MM 31G.....	179
GILOTRIF	20	HEALTHY ACCENTS PENTIP 6MM 31G.....	179
<i>glatiramer subcutaneous syringe 20 mg/ml, 40 mg/ml</i>	147	HEALTHY ACCENTS PENTIP 8MM 31G.....	179
<i>glatopa subcutaneous syringe 20 mg/ml, 40 mg/ml</i>	147	HEALTHY ACCENTS PENTIP 12MM 29G.....	179
<i>glutamine (sickle cell)</i>	215	HEB INCONTROL ALCOHOL 70% PADS.....	179
<i>glyburide</i>	159	HERCEPTIN HYLECTA.....	379
<i>glyburide micronized</i>	159	HERNEXEOS.....	417
<i>glyburide-metformin</i>	159	HUMIRA PEN.....	12
GNP ALCOHOL SWAB STERILE, TWO PLY.....	179	HUMIRA PEN CROHNS-UC-HS START.	12
GNP PEN NEEDLE 31G 5MM.....	179	HUMIRA SUBCUTANEOUS SYRINGE KIT 40 MG/0.8 ML.....	12
GNP PEN NEEDLE 32G 6MM.....	179	HUMIRA(CF).....	12
GNP SIMPLI PEN NEEDLE 32G 4MM...	179	HUMIRA(CF) PEDI CROHNS STARTER	12
GNP ULT C 0.3 ML 29GX1/2" (1/2) 1/2 UNIT.....	179	HUMIRA(CF) PEN.....	12
GNP ULT CMFRT 0.5 ML 29GX1/2"	179	HUMIRA(CF) PEN CROHNS-UC-HS.....	12
GNP ULTR CMFRT 0.5 ML 30GX5/16...	179	HUMIRA(CF) PEN PEDIATRIC UC.....	12
GNP ULTRA COMFORT 1 ML SYRINGE.....	179	HUMIRA(CF) PEN PSOR-UV-ADOL HS.	12
GNP ULTRA COMFORT 3/10 ML SYR..	179	HYRNUO.....	323
GOMEKLI ORAL CAPSULE 1 MG, 2 MG.....	237	IBRANCE.....	268
GOMEKLI ORAL TABLET FOR SUSPENSION.....	237	IBTROZI.....	349
GS PEN NEEDLE 31G X 8MM.....	179	<i>icatibant</i>	170
HADLIMA.....	18	ICLUSIG.....	286
HADLIMA PUSHTOUCH.....	18	IDHIFA.....	111
HADLIMA(CF).....	18	<i>imatinib oral tablet 100 mg, 400 mg</i>	172
HADLIMA(CF) PUSHTOUCH.....	18	IMBRUVICA ORAL CAPSULE 140 MG, 70 MG.....	169
HAEGARDA SUBCUTANEOUS RECON SOLN 2,000 UNIT, 3,000 UNIT	59	IMBRUVICA ORAL SUSPENSION.....	169
HARVONI ORAL PELLETS IN PACKET 33.75-150 MG, 45-200 MG.....	203	IMBRUVICA ORAL TABLET.....	169
HARVONI ORAL TABLET.....	203	IMDELLTRA.....	351
HEALTHWISE INS 0.3 ML 30GX5/16"..	179	IMJUDO.....	381
HEALTHWISE INS 0.3 ML 31GX5/16"..	179	IMKELDI.....	173
HEALTHWISE INS 0.5 ML 30GX5/16"..	179	IMPAVIDO.....	236
HEALTHWISE INS 0.5 ML 31GX5/16"..	179	INCONTROL PEN NEEDLE 12MM 29G	179
		INCONTROL PEN NEEDLE 4MM 32G..	179
		INCONTROL PEN NEEDLE 5MM 31G..	179
		INCONTROL PEN NEEDLE 6MM 31G..	179
		INCONTROL PEN NEEDLE 8MM 31G..	179

INCRELEX.....	228	INSUPEN PEN NEEDLE 32G 6MM (RX)	179
<i>indomethacin oral capsule</i>	166		179
<i>infliximab</i>	177	ITOVEBI ORAL TABLET 3 MG, 9 MG..	176
INGREZZA.....	400	IV ANTISEPTIC WIPES.....	179
INGREZZA INITIATION PK(TARDIV)..	400	IWILFIN.....	101
INGREZZA SPRINKLE.....	400	JAKAFI.....	315
INLURIYO.....	175	JAKAFI XR.....	315
INLYTA ORAL TABLET 1 MG, 5 MG.....	42	JASCAYD.....	243
INQOVI.....	84	<i>javygtor oral tablet,soluble</i>	316
INREBIC.....	129	JAYPIRCA ORAL TABLET 100 MG, 50	
INSULIN 1 ML SYRINGE.....	179	MG.....	284
INSULIN 1/2 ML SYRINGE.....	179	JEMPERLI.....	94
INSULIN 3/10 ML SYRINGE.....	179	JYNARQUE ORAL TABLET.....	372
INSULIN SYR 0.3 ML 31GX1/4(1/2).....	179	KALYDECO.....	195
INSULIN SYRIN 0.5 ML 28GX1/2"		KENDALL ALCOHOL 70% PREP PAD.	179
(OTC).....	179	KERENDIA.....	135
INSULIN SYRIN 0.5 ML 29GX1/2"		KESIMPTA PEN.....	258
(OTC).....	179	<i>ketorolac oral</i>	160
INSULIN SYRIN 0.5 ML 30GX1/2" (RX)	179	KEYTRUDA.....	275
INSULIN SYRIN 0.5 ML 30GX5/16"		KEYTRUDA QLEX.....	276
SHORT NEEDLE (OTC).....	179	KIMMTRAK.....	352
INSULIN SYRING 0.5 ML 27G 1/2"		KINERET.....	26
INNER.....	179	KISQALI 200 MG DAILY DOSE.....	298
INSULIN SYRINGE 0.3 ML.....	179	KISQALI FEMARA CO-PACK ORAL	
INSULIN SYRINGE 0.3 ML 31GX1/4.....	179	TABLET 200 MG/DAY(200 MG X 1)-2.5	
INSULIN SYRINGE 0.5 ML.....	179	MG, 400 MG/DAY(200 MG X 2)-2.5 MG,	
INSULIN SYRINGE 0.5 ML 31GX1/4.....	179	600 MG/DAY(200 MG X 3)-2.5 MG.....	299
INSULIN SYRINGE 1 ML.....	179	KISQALI ORAL TABLET 200 MG/DAY	
INSULIN SYRINGE 1 ML 27G 1/2"		(200 MG X 1), 400 MG/DAY (200 MG X	
INNER.....	179	2), 600 MG/DAY (200 MG X 3).....	298
INSULIN SYRINGE 1 ML 28GX1/2"		KOMZIFTI.....	415
(OTC).....	179	KOSELUGO ORAL CAPSULE 10 MG,	
INSULIN SYRINGE 1 ML 30GX1/2"		25 MG.....	322
SHORT NEEDLE (OTC).....	179	KOSELUGO ORAL CAPSULE,	
INSULIN SYRINGE 1 ML 30GX5/16"		SPRINKLE 5 MG, 7.5 MG.....	322
SHORT NEEDLE (OTC).....	179	KRAZATI.....	11
INSULIN SYRINGE 1 ML 31GX1/4".....	179	<i>lanreotide subcutaneous syringe 120</i>	
INSULIN SYRINGE 1 ML 31GX5/16"		<i>mg/0.5 ml</i>	199
SHORT NEEDLE,THIN II (OTC).....	179	<i>lapatinib</i>	200
INSULIN SYRINGE-NEEDLE U-100		LAZCLUZE ORAL TABLET 240 MG, 80	
SYRINGE 0.3 ML 29 GAUGE, 1 ML 29		MG.....	202
GAUGE X 1/2", 1/2 ML 28 GAUGE.....	179	<i>lenalidomide</i>	204
INSULIN U-500 SYRINGE-NEEDLE.....	179	LENVIMA.....	205
INSUPEN PEN NEEDLE 29GX1/2".....	179	<i>leuprolide acetate (3 month)</i>	208
INSUPEN PEN NEEDLE 31G 8MM.....	179	<i>leuprolide subcutaneous kit</i>	207
INSUPEN PEN NEEDLE 31GX3/16".....	179	<i>lidocaine topical adhesive patch,medicated</i>	
INSUPEN PEN NEEDLE 32G 4MM.....	179	5 %.....	217

<i>lidocaine topical ointment</i>	216	MAGELLAN INSULIN SYRINGE 1 ML	179
<i>lidocaine-prilocaine topical cream</i>	218	MARGENZA.....	225
<i>lidocan iii</i>	217	MAXICOMFORT II PEN NDL	
LISCO SPONGES 100/BAG.....	179	31GX6MM.....	179
LITE TOUCH 31GX1/4" PEN NEEDLE..	179	MAXICOMFORT INS 0.5 ML 27GX1/2"	179
LITE TOUCH INSULIN 0.5 ML SYR.....	179	MAXI-COMFORT INS 0.5 ML 28G.....	179
LITE TOUCH INSULIN 1 ML SYR.....	179	MAXICOMFORT INS 1 ML 27GX1/2" ...	179
LITE TOUCH INSULIN SYR 1 ML.....	179	MAXI-COMFORT INS 1 ML 28GX1/2"..	179
LITE TOUCH PEN NEEDLE 29G.....	179	MAXICOMFORT PEN NDL 29G X 5MM	
LITE TOUCH PEN NEEDLE 31G.....	179		179
LITETOUCH INS 0.3 ML 29GX1/2".....	179	MAXICOMFORT PEN NDL 29G X 8MM	
LITETOUCH INS 0.3 ML 30GX5/16".....	179		179
LITETOUCH INS 0.3 ML 31GX5/16".....	179	MAYZENT ORAL TABLET 0.25 MG, 1	
LITETOUCH INS 0.5 ML 31GX5/16".....	179	MG, 2 MG.....	326
LITETOUCH SYR 0.5 ML 28GX1/2".....	179	MAYZENT STARTER(FOR 1MG	
LITETOUCH SYR 0.5 ML 29GX1/2".....	179	MAINT).....	326
LITETOUCH SYR 0.5 ML 30GX5/16" ...	179	MAYZENT STARTER(FOR 2MG	
LITETOUCH SYRIN 1 ML 28GX1/2".....	179	MAINT).....	326
LITETOUCH SYRIN 1 ML 29GX1/2".....	179	<i>megestrol oral suspension 400 mg/10 ml</i>	
LITETOUCH SYRIN 1 ML 30GX5/16" ...	179	<i>(40 mg/ml), 625 mg/5 ml (125 mg/ml)</i>	167
LIVTENCITY.....	226	<i>megestrol oral tablet</i>	167
LONSURF ORAL TABLET 15-6.14 MG,		MEKINIST ORAL RECON SOLN.....	376
20-8.19 MG.....	384	MEKINIST ORAL TABLET 0.5 MG, 2	
LOQTORZI.....	374	MG.....	377
LORBRENA ORAL TABLET 100 MG,		MEKTOVI.....	54
25 MG.....	221	<i>metyrosine</i>	233
LUMAKRAS ORAL TABLET 120 MG,		MICRODOT PEN NEEDLE 31GX6MM..	179
240 MG, 320 MG.....	341	MICRODOT PEN NEEDLE 32GX4MM..	179
LUNSUMIO.....	240	MICRODOT PEN NEEDLE 33GX4MM..	179
LUNSUMIO VELO.....	240	MICRODOT READYGARD NDL 31G	
LUPRON DEPOT.....	211	5MM OUTER.....	179
LUPRON DEPOT (3 MONTH).....	211	<i>mifepristone oral tablet 300 mg</i>	235
LUPRON DEPOT (4 MONTH).....	211	<i>mimvey</i>	157
LUPRON DEPOT (6 MONTH).....	211	MINI PEN NEEDLE 32G 5MM.....	179
LUPRON DEPOT-PED (3 MONTH).....	213	MINI PEN NEEDLE 32G 8MM.....	179
LUPRON DEPOT-PED		MINI PEN NEEDLE 33G 5MM.....	179
INTRAMUSCULAR SYRINGE KIT.....	213	MINI PEN NEEDLE 33G 6MM.....	179
LUTRATE DEPOT (3 MONTH).....	208	MINI ULTRA-THIN II PEN NDL 31G	
LYNOZYFIC INTRAVENOUS		STERILE.....	179
SOLUTION 2 MG/ML, 20 MG/ML.....	219	MIPLYFFA.....	32
LYNPARZA.....	259	<i>modafinil oral tablet 100 mg, 200 mg</i>	241
LYTGOBI.....	140	MODEYSO.....	93
<i>macitentan</i>	224	MONOJECT 0.5 ML SYRN 28GX1/2".....	179
MAGELLAN INSUL SYRINGE 0.3 ML..	179	MONOJECT 1 ML SYRN 27X1/2".....	179
MAGELLAN INSUL SYRINGE 0.5 ML..	179	MONOJECT 1 ML SYRN 28GX1/2"	
MAGELLAN INSULIN SYR 0.3 ML.....	179	(OTC).....	179
MAGELLAN INSULIN SYR 0.5 ML.....	179	MONOJECT INSUL SYR U100 (OTC)....	179

MONOJECT INSUL SYR U100		NYVEPRIA.....	271
.5ML,29GX1/2" (OTC).....	179	ODOMZO.....	337
MONOJECT INSUL SYR U100 0.5 ML		OGIVRI.....	378
CONVERTS TO 29G (OTC).....	179	OGSIVEO ORAL TABLET 100 MG, 150	
MONOJECT INSUL SYR U100 1 ML.....	179	MG, 50 MG.....	252
MONOJECT INSUL SYR U100 1 ML 3'S,		OJEMDA ORAL SUSPENSION FOR	
29GX1/2" (OTC).....	179	RECONSTITUTION.....	375
MONOJECT INSUL SYR U100 1 ML		OJEMDA ORAL TABLET.....	375
W/O NEEDLE (OTC).....	179	OJJAARA.....	239
MONOJECT INSULIN SYR 0.3 ML.....	179	ONAPGO.....	29
MONOJECT INSULIN SYR 0.3 ML		ONUREG.....	43
(OTC).....	179	OPDIVO.....	254
MONOJECT INSULIN SYR 0.5 ML.....	179	OPDIVO QVANTIG.....	255
MONOJECT INSULIN SYR 0.5 ML		OPDUALAG.....	256
(OTC).....	179	OPSUMIT.....	224
MONOJECT INSULIN SYR 1 ML 3'S		ORENCIA.....	4
(OTC).....	179	ORENCIA (WITH MALTOSE).....	2
MONOJECT INSULIN SYR U-100.....	179	ORENCIA CLICKJECT.....	4
MONOJECT SYRINGE 0.3 ML.....	179	ORFADIN ORAL SUSPENSION.....	253
MONOJECT SYRINGE 0.5 ML.....	179	ORGOVYX.....	293
MONOJECT SYRINGE 1 ML.....	179	ORILISSA ORAL TABLET 150 MG, 200	
<i>morphine concentrate oral solution</i>	154	MG.....	103
MOUNJARO.....	150	ORKAMBI ORAL TABLET.....	223
NANO 2 GEN PEN NEEDLE 32G 4MM.....	179	ORSERDU ORAL TABLET 345 MG, 86	
NANO PEN NEEDLE 32G 4MM.....	179	MG.....	102
NERLYNX.....	245	OSENVELT.....	87
NIKTIMVO.....	41	OTEZLA.....	30
<i>nilotinib hcl oral capsule 150 mg, 200 mg,</i>		OTEZLA STARTER.....	30
<i>50 mg</i>	246	OTEZLA XR.....	30
NINLARO.....	197	OTEZLA XR INITIATION.....	30
<i>nintedanib</i>	248	<i>oxandrolone</i>	266
<i>nitisinone</i>	253	OZEMPIC ORAL.....	149
NIVESTYM.....	134	OZEMPIC SUBCUTANEOUS.....	149
NORDITROPIN FLEXPPO.....	334	<i>paroxetine hcl</i>	168
NOVOFINE 32G NEEDLES.....	179	<i>pazopanib oral tablet 200 mg, 400 mg</i>	270
NOVOFINE AUTOCOVER 30G		PEGASYS.....	273
NEEDLE.....	179	PEMAZYRE.....	277
NOVOFINE PLUS PEN NDL 32GX1/6".....	179	PEN NEEDLE 30G 5MM OUTER.....	179
NUBEQA.....	81	PEN NEEDLE 30G 8MM INNER.....	179
NUCALA SUBCUTANEOUS AUTO-		PEN NEEDLE 30G X 5/16".....	179
INJECTOR.....	230	PEN NEEDLE 31G X 1/4" HRI.....	179
NUCALA SUBCUTANEOUS RECON		PEN NEEDLE 33G 4MM.....	179
SOLN.....	230	PEN NEEDLE, DIABETIC NEEDLE 29	
NUCALA SUBCUTANEOUS SYRINGE		GAUGE X 1/2".....	179
100 MG/ML, 40 MG/0.4 ML.....	230	PEN NEEDLES 12MM 29G	
NUPLAZID.....	281	29GX12MM,STRL.....	179
NURTEC ODT.....	303	PEN NEEDLES 4MM 32G.....	179

PEN NEEDLES 5MM 31G		PRO COMFORT PEN NDL 4MM 32G....	179
31GX5MM,STRL,MINI (OTC).....	179	PRO COMFORT PEN NDL 5MM 32G....	179
PEN NEEDLES 8MM 31G		PRO-COMFORT ALCOHOL 70% PADS	179
31GX8MM,STRL,SHORT (OTC).....	179	PRODIGY INS SYR 1 ML 28GX1/2".....	179
<i>penicillamine oral tablet</i>	278	PRODIGY SYRNG 0.5 ML 31GX5/16"...	179
PENTIPS PEN NEEDLE 29G 1/2".....	179	PRODIGY SYRNGE 0.3 ML 31GX5/16".....	179
PENTIPS PEN NEEDLE 31G 1/4".....	179	<i>promethazine injection solution 25 mg/ml</i>	162
PENTIPS PEN NEEDLE 31GX3/16"		<i>promethazine oral tablet</i>	162
MINI, 5MM.....	179	<i>promethazine rectal suppository 25 mg</i>	162
PENTIPS PEN NEEDLE 31GX5/16"		<i>promethazine rectal suppository 12.5 mg,</i>	
SHORT, 8MM.....	179	<i>25 mg</i>	162
PENTIPS PEN NEEDLE 32G 1/4".....	179	PURE CMFT SFTY PEN NDL 31G 5MM	179
PENTIPS PEN NEEDLE 32GX5/32".....	179	PURE CMFT SFTY PEN NDL 31G 6MM	179
<i>phenobarbital</i>	161	PURE CMFT SFTY PEN NDL 32G 4MM	179
PIP PEN NEEDLE 31G X 5MM.....	179	PURE COMFORT ALCOHOL 70%	
PIP PEN NEEDLE 32G X 4MM.....	179	PADS.....	179
PIQRAY ORAL TABLET 200 MG/DAY		PURE COMFORT PEN NDL 32G 4MM..	179
(200 MG X 1), 250 MG/DAY (200 MG		PURE COMFORT PEN NDL 32G 5MM..	179
X1-50 MG X1), 300 MG/DAY (150 MG X		PURE COMFORT PEN NDL 32G 6MM..	179
2).....	22	PURE COMFORT PEN NDL 32G 8MM..	179
<i>pirfenidone oral capsule</i>	282	<i>pyrimethamine</i>	289
<i>pirfenidone oral tablet 267 mg, 534 mg,</i>		QINLOCK.....	307
<i>801 mg</i>	282	<i>quinine sulfate</i>	290
PLEGRIDY SUBCUTANEOUS PEN		QULIPTA.....	37
INJECTOR 125 MCG/0.5 ML, 63		RA ISOPROPYL ALCOHOL 70% WIPES	
MCG/0.5 ML- 94 MCG/0.5 ML.....	191		179
PLEGRIDY SUBCUTANEOUS		RA STERILE PADS 2"X 2".....	179
SYRINGE 125 MCG/0.5 ML, 63 MCG/0.5		RALDESY.....	380
ML- 94 MCG/0.5 ML.....	191	RAYA SURE PEN NEEDLE 29G 12MM..	179
<i>pomalidomide</i>	285	RAYA SURE PEN NEEDLE 31G 4MM..	179
POMALYST.....	285	RAYA SURE PEN NEEDLE 31G 5MM..	179
<i>posaconazole oral tablet,delayed release</i>		RAYA SURE PEN NEEDLE 31G 6MM..	179
<i>(dr/ec)</i>	287	RELION INS SYR 0.3 ML 31GX6MM....	179
PREMPHASE.....	158	RELION INS SYR 0.5 ML 31GX6MM....	179
PREMPRO.....	158	RELION INS SYR 1 ML 31GX15/64".....	179
PREVENT PEN NEEDLE 31GX1/4".....	179	RELION PEN NEEDLE 32GX5/32".....	179
PREVENT PEN NEEDLE 31GX5/16".....	179	RELION SYRING 0.3 ML 31GX5/16"	
PREVYMIS ORAL TABLET.....	206	INNER.....	179
PRO COMFORT 0.5 ML 30GX1/2".....	179	RELION SYRING 0.5 ML 31GX5/16"	
PRO COMFORT 0.5 ML 30GX5/16".....	179	INNER (OTC).....	179
PRO COMFORT 0.5 ML 31GX5/16".....	179	RETACRIT INJECTION SOLUTION	
PRO COMFORT 1 ML 30GX1/2".....	179	10,000 UNIT/ML, 2,000 UNIT/ML,	
PRO COMFORT 1 ML 30GX5/16".....	179	20,000 UNIT/2 ML, 20,000 UNIT/ML,	
PRO COMFORT 1 ML 31GX5/16".....	179	3,000 UNIT/ML, 4,000 UNIT/ML, 40,000	
PRO COMFORT ALCOHOL 70% PADS	179	UNIT/ML.....	118
PRO COMFORT PEN NDL 32G 8MM....	179	RETEVMO ORAL CAPSULE 40 MG, 80	
PRO COMFORT PEN NDL 32G X 1/4"....	179	MG.....	321

RETEVMO ORAL TABLET 120 MG, 160 MG, 40 MG, 80 MG.....	321	SELARSDI.....	392, 394
REVCovi.....	105	SEROSTIM SUBCUTANEOUS RECON SOLN 4 MG, 5 MG, 6 MG.....	336
REVUFORJ ORAL TABLET 110 MG, 160 MG, 25 MG.....	297	SIGNIFOR.....	269
REZDIFFRA.....	295	<i>sildenafil (pulm.hypertension) oral tablet..</i>	324
REZLIDHIA.....	260	SIRTURO.....	45
REZUROCK.....	47	SKY SAFETY PEN NEEDLE 30G 5MM..	179
RINVOQ.....	388	SKY SAFETY PEN NEEDLE 30G 8MM..	179
RINVOQ LQ.....	388	SKYRIZI INTRAVENOUS.....	308
RITUXAN HYCELA.....	310	SKYRIZI SUBCUTANEOUS PEN INJECTOR.....	308
ROMVIMZA.....	407	SKYRIZI SUBCUTANEOUS SYRINGE 150 MG/ML.....	308
ROZLYTREK ORAL CAPSULE 100 MG, 200 MG.....	114	SKYRIZI SUBCUTANEOUS WEARABLE INJECTOR.....	308
ROZLYTREK ORAL PELLETS IN PACKET.....	115	<i>sodium oxybate.....</i>	328
RUBRACA.....	314	SOMATULINE DEPOT SUBCUTANEOUS SYRINGE 60 MG/0.2 ML, 90 MG/0.3 ML.....	199
RYBELSUS.....	149	SOMAVERT.....	274
RYBREVANT.....	25	<i>sorafenib.....</i>	338
RYBREVANT FASPRO.....	24	SPRAVATO.....	123
RYDAPT.....	234	STIVARGA.....	292
RYTELO.....	174	STRENSIQ.....	34
SAFESNAP INS SYR UNITS-100 0.3 ML 30GX5/16",10X10.....	179	SUBVENITE ORAL SUSPENSION.....	198
SAFESNAP INS SYR UNITS-100 0.5 ML 29GX1/2",10X10.....	179	<i>sunitinib malate.....</i>	343
SAFESNAP INS SYR UNITS-100 0.5 ML 30GX5/16",10X10.....	179	SURE CMFT SFTY PEN NDL 31G 6MM..	179
SAFESNAP INS SYR UNITS-100 1 ML 28GX1/2",10X10.....	179	SURE CMFT SFTY PEN NDL 32G 4MM..	179
SAFESNAP INS SYR UNITS-100 1 ML 29GX1/2",10X10.....	179	SURE COMFORT 0.5 ML SYRINGE.....	179
SAFETY PEN NEEDLE 31G 4MM.....	179	SURE COMFORT 1 ML SYRINGE.....	179
SAFETY PEN NEEDLE 5MM X 31G.....	179	SURE COMFORT 3/10 ML SYRINGE....	179
SAFETY SYRINGE 0.5 ML 30G 1/2".....	179	SURE COMFORT 3/10 ML SYRINGE INSULIN SYRINGE.....	179
<i>sapropterin oral tablet,soluble.....</i>	316	SURE COMFORT 30G PEN NEEDLE.....	179
SCEMBLIX ORAL TABLET 100 MG, 20 MG, 40 MG.....	33	SURE COMFORT ALCOHOL PREP PADS.....	179
<i>scopolamine base.....</i>	164	SURE COMFORT INS 0.3 ML 31GX1/4..	179
SECURESAFE PEN NDL 30GX5/16" OUTER.....	179	SURE COMFORT INS 0.5 ML 31GX1/4..	179
SECURESAFE PEN NEEDLE 31G 5MM OUTER.....	179	SURE COMFORT INS 1 ML 31GX1/4" ...	179
SECURESAFE SYR 0.5 ML 29G 1/2" OUTER.....	179	SURE COMFORT PEN NDL 29GX1/2" 12.7MM.....	179
SECURESAFE SYRNG 1 ML 29G 1/2" OUTER.....	179	SURE COMFORT PEN NDL 31G 5MM..	179
		SURE COMFORT PEN NDL 31G 8MM..	179
		SURE COMFORT PEN NDL 32G 4MM..	179
		SURE COMFORT PEN NDL 32G 6MM..	179
		SURE-FINE PEN NEEDLES 12.7MM.....	179
		SURE-FINE PEN NEEDLES 5MM.....	179

SURE-FINE PEN NEEDLES 8MM.....	179	<i>testosterone cypionate</i>	359
SURE-JECT INSU SYR U100 0.3 ML.....	179	<i>testosterone enanthate</i>	360
SURE-JECT INSU SYR U100 0.5 ML.....	179	<i>testosterone transdermal gel in metered-</i>	
SURE-JECT INSU SYR U100 1 ML.....	179	<i>dose pump 12.5 mg/ 1.25 gram (1 %),</i>	
SURE-JECT INSUL SYR U100 1 ML.....	179	<i>20.25 mg/1.25 gram (1.62 %).....</i>	358
SURE-JECT INSULIN SYRINGE 1 ML..	179	<i>testosterone transdermal gel in packet 1 %</i>	
SURE-PREP ALCOHOL PREP PADS.....	179	<i>(25 mg/2.5gram), 1 % (50 mg/5 gram).....</i>	358
SYMPAZAN.....	71	<i>tetrabenazine</i>	361
SYNRIBO.....	261	TEVIMBRA.....	363
TABRECTA.....	64	THALOMID ORAL CAPSULE 100 MG,	
<i>tadalafil oral tablet 2.5 mg, 5 mg.....</i>	345	150 MG, 200 MG, 50 MG.....	362
TAFINLAR ORAL CAPSULE.....	77	THINPRO INS SYRIN U100-0.3 ML.....	179
TAFINLAR ORAL TABLET FOR		THINPRO INS SYRIN U100-0.5 ML.....	179
SUSPENSION.....	78	THINPRO INS SYRIN U100-1 ML.....	179
TAGRISSO.....	265	TIBSOVO.....	196
TALVEY.....	350	TIVDAK.....	364
TALZENNA.....	347	<i>tofacitinib</i>	370
TAVNEOS.....	38	<i>tolvaptan (polycys kidney dis) oral tablets,</i>	
TECHLITE 0.3 ML 29GX12MM (1/2).....	179	<i>sequential</i>	372
TECHLITE 0.3 ML 30GX8MM (1/2).....	179	<i>torpenz oral tablet 10 mg, 2.5 mg, 5 mg,</i>	
TECHLITE 0.3 ML 31GX6MM (1/2).....	179	<i>7.5 mg.....</i>	126
TECHLITE 0.3 ML 31GX8MM (1/2).....	179	TRELSTAR INTRAMUSCULAR	
TECHLITE 0.5 ML 30GX12MM (1/2).....	179	SUSPENSION FOR RECONSTITUTION	385
TECHLITE 0.5 ML 30GX8MM (1/2).....	179	TREMFYA INTRAVENOUS.....	152
TECHLITE 0.5 ML 31GX6MM (1/2).....	179	TREMFYA ONE-PRESS.....	152
TECHLITE 0.5 ML 31GX8MM (1/2).....	179	TREMFYA PEN INDUCTION PK(2PEN)	
TECHLITE INS SYR 1 ML 29GX12MM.	179		152
TECHLITE INS SYR 1 ML 30GX12MM.	179	TREMFYA PEN SUBCUTANEOUS PEN	
TECHLITE INS SYR 1 ML 31GX6MM...	179	INJECTOR 200 MG/2 ML.....	152
TECHLITE INS SYR 1 ML 31GX8MM...	179	TREMFYA SUBCUTANEOUS	
TECHLITE PEN NEEDLE 29GX1/2".....	179	SYRINGE.....	152
TECHLITE PEN NEEDLE 29GX3/8".....	179	<i>tretinoin topical cream</i>	373
TECHLITE PEN NEEDLE 31GX1/4".....	179	<i>tridacaine iii</i>	217
TECHLITE PEN NEEDLE 31GX3/16"....	179	<i>trientine oral capsule 250 mg</i>	383
TECHLITE PEN NEEDLE 31GX5/16"....	179	TRIKAFTA ORAL GRANULES IN	
TECHLITE PEN NEEDLE 32GX1/4".....	179	PACKET, SEQUENTIAL.....	106
TECHLITE PEN NEEDLE 32GX5/16"....	179	TRIKAFTA ORAL TABLETS,	
TECHLITE PEN NEEDLE 32GX5/32"....	179	SEQUENTIAL.....	106
TECHLITE PLUS PEN NDL 32G 4MM...	179	TRUE CMFRT PRO 0.5 ML 30G 5/16"....	179
TECVAYLI.....	353	TRUE CMFRT PRO 0.5 ML 31G 5/16"....	179
TEPMETKO.....	356	TRUE CMFRT PRO 0.5 ML 32G 5/16"....	179
<i>teriparatide subcutaneous pen injector 20</i>		TRUE CMFT SFTY PEN NDL 31G 5MM	179
<i>mcg/dose (560mcg/2.24ml).....</i>	357	TRUE CMFT SFTY PEN NDL 31G 6MM	179
TERUMO INS SYRINGE U100-1 ML.....	179	TRUE CMFT SFTY PEN NDL 32G 4MM	179
TERUMO INS SYRINGE U100-1/2 ML..	179	TRUE COMFORT 0.5 ML 30G 1/2".....	179
TERUMO INS SYRINGE U100-1/3 ML..	179	TRUE COMFORT 0.5 ML 30G 5/16".....	179
TERUMO INS SYRNG U100-1/2 ML.....	179	TRUE COMFORT 0.5 ML 31G 5/16".....	179

TRUE COMFORT 0.5 ML 31GX5/16".....	179	TRUEPLUS SYR 1 ML 31GX5/16".....	179
TRUE COMFORT 1 ML 31GX5/16".....	179	TRULICITY.....	148
TRUE COMFORT ALCOHOL 70% PADS.....	179	TRUQAP.....	63
TRUE COMFORT PEN NDL 31G 8MM..	179	TRUXIMA.....	311
TRUE COMFORT PEN NDL 31GX5MM	179	TUKYSA ORAL TABLET 150 MG, 50 MG.....	386
TRUE COMFORT PEN NDL 31GX6MM	179	TURALIO.....	280
TRUE COMFORT PEN NDL 32G 5MM..	179	TYENNE.....	366, 368
TRUE COMFORT PEN NDL 32G 6MM..	179	TYENNE AUTOINJECTOR.....	368
TRUE COMFORT PEN NDL 32GX4MM	179	TYMLOS.....	1
TRUE COMFORT PEN NDL 33G 4MM..	179	UBRELVY.....	387
TRUE COMFORT PEN NDL 33G 5MM..	179	UDENYCA ONBODY.....	272
TRUE COMFORT PEN NDL 33G 6MM..	179	ULT CFT 0.3 ML 31GX5/16" (1/2) 1/2 UNIT.....	179
TRUE COMFORT PRO 1 ML 30G 1/2"...	179	ULTICAR INS 0.3 ML 31GX1/4(1/2).....	179
TRUE COMFORT PRO 1 ML 30G 5/16".	179	ULTICARE INS 0.3 ML 31GX1/4".....	179
TRUE COMFORT PRO 1 ML 31G 5/16".	179	ULTICARE INS 0.5 ML 31GX1/4".....	179
TRUE COMFORT PRO 1 ML 32G 5/16".	179	ULTICARE INS 1 ML 31GX1/4".....	179
TRUE COMFORT PRO ALCOHOL PADS.....	179	ULTICARE INS SYR 1 ML 30GX1/2".....	179
TRUE COMFORT SFTY 1 ML 30G 1/2".	179	ULTICARE PEN NEEDLE 31GX3/16".....	179
TRUE COMFRT PRO 0.5 ML 30G 1/2"...	179	ULTICARE PEN NEEDLE 6MM 31G.....	179
TRUE COMFRT SFTY 1 ML 30G 5/16"..	179	ULTICARE PEN NEEDLE 8MM 31G.....	179
TRUE COMFRT SFTY 1 ML 31G 5/16"..	179	ULTICARE PEN NEEDLES 12MM 29G.	179
TRUE COMFRT SFTY 1 ML 32G 5/16"..	179	ULTICARE PEN NEEDLES 4MM 32G MICRO, 32GX4MM.....	179
TRUE-CMFRT PRO PEN NDL 31G 5MM	179	ULTICARE PEN NEEDLES 6MM 32G...	179
TRUE-CMFRT PRO PEN NDL 31G 6MM	179	ULTICARE SAFE PEN NDL 30G 8MM..	179
TRUE-CMFRT PRO PEN NDL 31G 8MM	179	ULTICARE SAFE PEN NDL 5MM 30G..	179
TRUE-CMFRT PRO PEN NDL 32G 4MM	179	ULTICARE SYR 0.3 ML 30GX1/2".....	179
TRUEPLUS PEN NEEDLE 29GX1/2".....	179	ULTICARE SYR 0.3 ML 31GX5/16" SHORT NDL.....	179
TRUEPLUS PEN NEEDLE 31G X 1/4"...	179	ULTICARE SYR 0.5 ML 30GX1/2".....	179
TRUEPLUS PEN NEEDLE 31GX3/16"...	179	ULTICARE SYR 0.5 ML 31GX5/16" SHORT NDL.....	179
TRUEPLUS PEN NEEDLE 31GX5/16"...	179	ULTICARE SYR 1 ML 31GX5/16".....	179
TRUEPLUS PEN NEEDLE 32GX5/32"...	179	ULTIGUARD SAFE 1 ML 30G 12.7MM.	179
TRUEPLUS SYR 0.3 ML 29GX1/2".....	179	ULTIGUARD SAFE0.3 ML 30G 12.7MM	179
TRUEPLUS SYR 0.3 ML 30GX5/16".....	179	ULTIGUARD SAFE0.5 ML 30G 12.7MM	179
TRUEPLUS SYR 0.3 ML 31GX5/16".....	179	ULTIGUARD SAFEPACK 1 ML 31G 8MM.....	179
TRUEPLUS SYR 0.5 ML 28GX1/2".....	179	ULTIGUARD SAFEPACK 29G 12.7MM	179
TRUEPLUS SYR 0.5 ML 29GX1/2".....	179	ULTIGUARD SAFEPACK 31G 5MM.....	179
TRUEPLUS SYR 0.5 ML 30GX5/16".....	179	ULTIGUARD SAFEPACK 31G 6MM.....	179
TRUEPLUS SYR 0.5 ML 31GX5/16".....	179	ULTIGUARD SAFEPACK 31G 8MM.....	179
TRUEPLUS SYR 1 ML 28GX1/2".....	179	ULTIGUARD SAFEPACK 32G 4MM.....	179
TRUEPLUS SYR 1 ML 29GX1/2".....	179		
TRUEPLUS SYR 1 ML 30GX5/16".....	179		

ULTIGUARD SAFEPAK 32G 6MM.....	179	ULTRACARE PEN NEEDLE 32GX1/4" ..	179
ULTIGUARD SAFEPK 0.3 ML 31G		ULTRACARE PEN NEEDLE 32GX3/16"	179
8MM.....	179	ULTRACARE PEN NEEDLE 32GX5/32"	179
ULTIGUARD SAFEPK 0.5 ML 31G		ULTRACARE PEN NEEDLE 33GX5/32"	179
8MM.....	179	ULTRA-FINE 0.3 ML 30G 12.7MM.....	179
ULTILET ALCOHOL STERL SWAB.....	179	ULTRA-FINE 0.3 ML 31G 6MM (1/2).....	179
ULTILET INSULIN SYRINGE 0.3 ML...	179	ULTRA-FINE 0.3 ML 31G 8MM (1/2).....	179
ULTILET INSULIN SYRINGE 0.5 ML...	179	ULTRA-FINE 0.5 ML 30G 12.7MM.....	179
ULTILET INSULIN SYRINGE 1 ML.....	179	ULTRA-FINE INS SYR 1 ML 31G 6MM	179
ULTILET PEN NEEDLE	179	ULTRA-FINE INS SYR 1 ML 31G 8MM	179
ULTILET PEN NEEDLE 4MM 32G.....	179	ULTRA-FINE PEN NDL 29G 12.7MM....	179
ULTRA COMFORT 0.3 ML 29GX1/2"	179	ULTRA-FINE PEN NEEDLE 31G 5MM..	179
ULTRA COMFORT 0.3 ML SYRINGE...	179	ULTRA-FINE PEN NEEDLE 31G 8MM..	179
ULTRA COMFORT 0.5 ML 28GX1/2"		ULTRA-FINE PEN NEEDLE 32G 6MM..	179
CONVERTS TO 29G.....	179	ULTRA-FINE SYR 0.3 ML 31G 6MM.....	179
ULTRA COMFORT 0.5 ML 29GX1/2"	179	ULTRA-FINE SYR 0.3 ML 31G 8MM.....	179
ULTRA COMFORT 0.5 ML 31GX5/16" ..	179	ULTRA-FINE SYR 0.5 ML 31G 6MM.....	179
ULTRA COMFORT 0.5 ML SYRINGE...	179	ULTRA-FINE SYR 0.5 ML 31G 8MM.....	179
ULTRA COMFORT 1 ML 29GX1/2"	179	ULTRA-FINE SYR 1 ML 30G 12.7MM...	179
ULTRA COMFORT 1 ML 30GX5/16"	179	ULTRA-THIN II 1 ML 31GX5/16"	179
ULTRA COMFORT 1 ML 31GX5/16"	179	ULTRA-THIN II INS 0.3 ML 30G	179
ULTRA COMFORT 1 ML SYRINGE.....	179	ULTRA-THIN II INS 0.3 ML 31G	179
ULTRA FLO 0.3 ML 30G 1/2" (1/2).....	179	ULTRA-THIN II INS 0.5 ML 29G	179
ULTRA FLO 0.3 ML 30G 5/16"(1/2).....	179	ULTRA-THIN II INS 0.5 ML 30G	179
ULTRA FLO 0.3 ML 31G 5/16"(1/2).....	179	ULTRA-THIN II INS 0.5 ML 31G	179
ULTRA FLO PEN NEEDLE 31G 5MM...	179	ULTRA-THIN II INS SYR 1 ML 29G	179
ULTRA FLO PEN NEEDLE 31G 8MM...	179	ULTRA-THIN II INS SYR 1 ML 30G	179
ULTRA FLO PEN NEEDLE 32G 4MM...	179	ULTRA-THIN II PEN NDL 29GX1/2"	179
ULTRA FLO PEN NEEDLE 33G 4MM...	179	ULTRA-THIN II PEN NDL 31GX5/16.....	179
ULTRA FLO PEN NEEDLES 12MM 29G		UNIFINE OTC PEN NEEDLE 31G 5MM	179
.....	179	UNIFINE OTC PEN NEEDLE 32G 4MM	179
ULTRA FLO SYR 0.3 ML 29GX1/2"	179	UNIFINE PENTIPS 12MM 29G	
ULTRA FLO SYR 0.3 ML 30G 5/16"	179	29GX12MM, STRL.....	179
ULTRA FLO SYR 0.3 ML 31G 5/16"	179	UNIFINE PENTIPS 31GX3/16"	
ULTRA FLO SYR 0.5 ML 29G 1/2"	179	31GX5MM,STRL,MINI.....	179
ULTRA THIN PEN NDL 32G X 4MM.....	179	UNIFINE PENTIPS 32G 4MM.....	179
ULTRACARE INS 0.3 ML 30GX5/16"	179	UNIFINE PENTIPS 32GX1/4"	179
ULTRACARE INS 0.3 ML 31GX5/16"	179	UNIFINE PENTIPS 33GX5/32"	179
ULTRACARE INS 0.5 ML 30GX1/2"	179	UNIFINE PENTIPS 6MM 31G.....	179
ULTRACARE INS 0.5 ML 30GX5/16"	179	UNIFINE PENTIPS 8MM 31G	
ULTRACARE INS 0.5 ML 31GX5/16"	179	31GX8MM, SHORT, STRL.....	179
ULTRACARE INS 1 ML 30G X 5/16"	179	UNIFINE PENTIPS MAX 30GX3/16"	179
ULTRACARE INS 1 ML 30GX1/2"	179	UNIFINE PENTIPS PLUS 29GX1/2"	
ULTRACARE INS 1 ML 31G X 5/16"	179	12MM.....	179
ULTRACARE PEN NEEDLE 31GX1/4" ..	179	UNIFINE PENTIPS PLUS 30GX3/16"	179
ULTRACARE PEN NEEDLE 31GX3/16"	179	UNIFINE PENTIPS PLUS 31G 5MM.....	179
ULTRACARE PEN NEEDLE 31GX5/16"	179		

UNIFINE PENTIPS PLUS 31GX1/4"		VERIFINE PLUS PEN NDL 32G 4MM-	
ULTRA SHORT, 6MM.....	179	SHARPS CONTAINER.....	179
UNIFINE PENTIPS PLUS 32GX5/32"	179	VERIFINE SYRING 0.5 ML 29G 1/2"	179
UNIFINE PENTIPS PLUS 33GX5/32"	179	VERIFINE SYRING 1 ML 31G 5/16"	179
UNIFINE PROTECT 30G 5MM.....	179	VERIFINE SYRNG 0.3 ML 31G 5/16"	179
UNIFINE PROTECT 30G 8MM.....	179	VERIFINE SYRNG 0.5 ML 31G 5/16"	179
UNIFINE PROTECT 32G 4MM.....	179	VERQUVO.....	405
UNIFINE SAFECONTROL 30G 5MM.....	179	VERSALON ALL PURPOSE SPONGE	
UNIFINE SAFECONTROL 30G 8MM.....	179	25'S,N-STERILE,3PLY	179
UNIFINE SAFECONTROL 31G 5MM.....	179	VERZENIO	6
UNIFINE SAFECONTROL 31G 6MM.....	179	<i>vigabatrin</i>	406
UNIFINE SAFECONTROL 31G 8MM.....	179	<i>vigadrone</i>	406
UNIFINE SAFECONTROL 32G 4MM.....	179	<i>vigpoder</i>	406
UNIFINE ULTRA PEN NDL 31G 5MM..	179	VITRAKVI ORAL CAPSULE 100 MG,	
UNIFINE ULTRA PEN NDL 31G 6MM..	179	25 MG	201
UNIFINE ULTRA PEN NDL 31G 8MM..	179	VITRAKVI ORAL SOLUTION	201
UNIFINE ULTRA PEN NDL 32G 4MM..	179	VIVIMUSTA	49
UPTRAVI ORAL TABLET 1,000 MCG,		VIZIMPRO	79
1,200 MCG, 1,400 MCG, 1,600 MCG, 200		VONJO	267
MCG, 400 MCG, 600 MCG, 800 MCG	319	VOQUEZNA	409
UPTRAVI ORAL TABLETS,DOSE		VORANIGO	410
PACK.....	319	<i>voriconazole oral suspension for</i>	
<i>ustekinumab-aauz</i>	390	<i>reconstitution</i>	411
VALCHLOR	229	VOSEVI	332
VANFLYTA	291	VOWST	128
VANISHPOINT 0.5 ML 30GX1/2" SY		VUMERITY	92
OUTER.....	179	VYALEV	138
VANISHPOINT INS 0.5 ML 30G 8MM		VYLOY	416
OUTER.....	179	VYNDAMAX.....	346
VANISHPOINT INS 1 ML 30GX3/16"	179	WEBCOL ALCOHOL PREPS	
VANISHPOINT U-100 29X1/2 SYR.....	179	20'S,LARGE	179
VENCLEXTA ORAL TABLET 10 MG,		WELIREG	48
100 MG, 50 MG.....	404	WINREVAIR.....	339
VENCLEXTA STARTING PACK.....	404	WM UNIFINE PENTIP PLUS 8MM 31G	
VEOZAH	132	W/ SAFETY CLICK.....	179
VERIFINE INS SYR 1 ML 29G 1/2"	179	XALKORI ORAL CAPSULE.....	75
VERIFINE PEN NEEDLE 29G 12MM.....	179	XALKORI ORAL PELLET 150 MG, 20	
VERIFINE PEN NEEDLE 31G 5MM.....	179	MG, 50 MG	76
VERIFINE PEN NEEDLE 31G X 6MM... 179		XDEMVY	222
VERIFINE PEN NEEDLE 31G X 8MM... 179		XELJANZ	370
VERIFINE PEN NEEDLE 32G 6MM.....	179	XELJANZ XR.....	370
VERIFINE PEN NEEDLE 32G X 4MM... 179		XERMELO	355
VERIFINE PEN NEEDLE 32G X 5MM... 179		XIFAXAN ORAL TABLET 200 MG, 550	
VERIFINE PLUS PEN NDL 31G 5MM... 179		MG	300
VERIFINE PLUS PEN NDL 31G 8MM... 179		XOLAIR.....	262
VERIFINE PLUS PEN NDL 32G 4MM... 179		XOSPATA	145

XPOVIO ORAL TABLET 100 MG/WEEK (50 MG X 2), 40 MG/WEEK (10 MG X 4), 40 MG/WEEK (40 MG X 1), 40MG TWICE WEEK (40 MG X 2), 60 MG/WEEK (60 MG X 1), 60MG TWICE WEEK (120 MG/WEEK), 80 MG/WEEK (40 MG X 2), 80 MG/WEEK (80 MG X 1), 80MG TWICE WEEK (160 MG/WEEK).....	320
XTANDI ORAL CAPSULE.....	116
XTANDI ORAL TABLET 40 MG, 80 MG	116
YERVOY	193
YESINTEK.....	396, 398
YONSA.....	8
YUFLYMA(CF).....	14
YUFLYMA(CF) AI CROHN'S-UC-HS	14
YUFLYMA(CF) AUTOINJECTOR.....	14
<i>yulithira oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg</i>	126
YUTREPIA.....	382
ZEJULA ORAL CAPSULE.....	250
ZEJULA ORAL TABLET	250
ZELBORAF	403
ZIIHERA.....	412
ZIRABEV	52
ZOLADEX.....	151
ZTALMY	142
ZTLIDO	217
ZURZUVAE ORAL CAPSULE 20 MG, 25 MG, 30 MG.....	418
ZYDELIG	171
ZYKADIA.....	66
ZYNLONTA.....	220
ZYNYZ.....	296