

## ABALOPARATIDE (TYMLOS)

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### MEDICATION(S)

TYMLOS

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Total parathyroid hormone analog therapy has exceeded 2 years. Being used with other osteoporosis drugs.

### REQUIRED MEDICAL INFORMATION

Osteoporosis: patient has a history of a broken bone not due to trauma (non-traumatic fracture) or T-score between -1.0 and -2.5 and is at high risk for fracture or T-score lower than -2.5 AND one of the following: trial of a bisphosphonate (e.g. alendronate, ibandronate, risedronate, or zoledronic acid) or Prolia, OR side effect to bisphosphonate therapy or Prolia therapy that supports discontinuation, OR Patient is at very high risk of fracture by meeting at least one of the following: non-traumatic fracture while on bisphosphonate therapy or Prolia, patient has experienced a recent fracture (within the past 12 months) or history of multiple fractures, patient experienced a fracture while on long-term glucocorticoid therapy, or T-score less than -3.0, or patient is at high risk for falls, or 10-year hip fracture probability greater than 4.5% based on FRAX score, or 10-year major osteoporosis-related fracture probability greater than 30% based on FRAX score

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## ABEMACICLIB (VERZENIO)

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### **MEDICATION(S)**

VERZENIO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **ABIRATERONE (ZYTIGA)**

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### **MEDICATION(S)**

ABIRATERONE ACETATE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ACALABRUTINIB (CALQUENCE)

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### **MEDICATION(S)**

CALQUENCE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ACETAMINOPHEN/CAFFEINE/DIHYDROCODEINE

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### MEDICATION(S)

APAP-CAFF-DIHYDROCODEINE 325-30-16 MG TAB, DVORAH

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with other short-acting opioids.

### REQUIRED MEDICAL INFORMATION

Moderate to severe pain, initial use: trial of at least two preferred short-acting narcotic drugs, and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain. Ongoing use (more than 90 days of narcotic therapy): evaluation by pain specialist, documented patient-specific treatment plan for evaluating ongoing need for narcotic pain relief, potential misuse, and monitoring plan for adverse side effects, and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

30 days

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ACYCLOVIR CREAM (ZOVIRAX)

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### **MEDICATION(S)**

ACYCLOVIR 5 % CREAM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ACYCLOVIR OINTMENT (ZOVIRAX)

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### MEDICATION(S)

ACYCLOVIR 5 % OINTMENT

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient has tried or has a medical reason for not trying a herpes antiviral drug you take by mouth (e.g. acyclovir, valacyclovir).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## ADAGRASIB (KRAZATI)

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### MEDICATION(S)

KRAZATI

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ADALIMUMAB (HUMIRA)

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### MEDICATION(S)

HUMIRA, HUMIRA PEDIATRIC CROHNS START, HUMIRA PEN, HUMIRA PEN-CD/UC/HS STARTER, HUMIRA PEN-PEDIATRIC UC START, HUMIRA PEN-PS/UV/ADOL HS START, HUMIRA PEN-PSOR/UVEIT STARTER

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Spondyloarthritis (SpA): patient is not able to take NSAIDs due to history of GI bleed or ulcer  
OR patient has tried one RX strength NSAID in combination with a PPI and had GI side effects  
OR patient's condition did not respond to a trial of two different RX strength NSAIDs.

Crohn's Disease (CD), weekly dosing: patient has tried every other week dosing and had a flare or loss in response.

Hidradenitis suppurativa (HS): patient has Hurley stage II or III HS.

Non-infectious uveitis: patient has tried a systemic corticosteroid (e.g. prednisone, dexamethasone, hydrocortisone) or has a medical reason why corticosteroids cannot be used.

Plaque Psoriasis (PsO), initial use: patient has tried one DMARD or has a medical reason why methotrexate (MTX), cyclosporine, and acitretin cannot be used AND baseline PASI score 10 or more OR BSA 3% or more OR sensitive areas are involved OR disease affects daily living.

Ongoing use: PASI or BSA improved on Humira.

Psoriatic Arthritis (PsA): patient has tried one DMARD or has a medical reason why MTX, leflunomide, and sulfasalazine cannot be used.

Rheumatoid Arthritis (RA) and Juvenile Idiopathic Arthritis (JIA): treatment failure or side effect with methotrexate (MTX) or patient has a medical reason why MTX cannot be used.

### AGE RESTRICTION

N/A

**PRESCRIBER RESTRICTION**

RA, PsA, JIA, AS: Rheumatologist. PsO: Rheumatologist or Dermatologist. HS: Dermatologist.  
Non-infectious uveitis: Ophthalmologist.

**COVERAGE DURATION**

PsO, initial: 16 weeks - ongoing use: plan year All other indications: plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## ADAPALENE (DIFFERIN)

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### **MEDICATION(S)**

ADAPALENE 0.1 % CREAM, ADAPALENE 0.1 % GEL, ADAPALENE 0.3 % GEL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

40 years of age or older. No prior authorization required for less than 40 years old.

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## AFATINIB DIMALEATE (GILOTRIF)

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### MEDICATION(S)

GILOTRIF

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan Year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ALECTINIB (ALECENSA)

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### MEDICATION(S)

ALECENSA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan Year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ALISKIREN (TEKTURNA)

---

### MEDICATION(S)

ALISKIREN FUMARATE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Trial and failure of at least one ARB (e.g. losartan) and one of the following drug classes: ACE inhibitor (e.g. lisinopril), CCB (e.g. diltiazem, amlodipine), thiazide (e.g. hydrochlorothiazide), or beta blocker (e.g. atenolol).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ALITRETINOIN (PANRETIN)

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### **MEDICATION(S)**

PANRETIN

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **ALOSETRON (LOTRONEX)**

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### **MEDICATION(S)**

ALOSETRON HCL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Irritable bowel syndrome with diarrhea (IBS-D), initial use: patient is female, and condition did not get better with use of an antispasmodic or antidiarrheal drug. Ongoing use: IBS symptoms improved with alosetron and patient does not have constipation problems.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Initial: 2 months

Ongoing use: plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ALPELISIB (PIQRAY)

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### **MEDICATION(S)**

PIQRAY (200 MG DAILY DOSE), PIQRAY (250 MG DAILY DOSE), PIQRAY (300 MG DAILY DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ALPROSTADIL (MUSE)

---

### MEDICATION(S)

MUSE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another ED drug.

### REQUIRED MEDICAL INFORMATION

Erectile dysfunction (ED) is caused by a drug that has been reported in the medical literature to cause erectile dysfunction or erectile dysfunction is caused by an underlying medical condition.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Limited to 6 tablets per month.

### PART B PREREQUISITE

N/A

## AMBRISANTAN (LETAIRIS)

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### MEDICATION(S)

AMBRISANTAN

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Confirmation of Pulmonary Arterial Hypertension (WHO Group I) by right heart catheterization test.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## AMIFAMPRIDINE (RUZURGI)

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### MEDICATION(S)

RUZURGI

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another aminopyridine drug (e.g. dalfampridine, Firdapse). History of seizures. Allergy to aminopyridine.

### REQUIRED MEDICAL INFORMATION

Diagnosis confirmed by proximal muscle weakness and one of the following tests: positive anti-P/Q-type voltage-gated calcium channel (VGCC) antibody test OR compound muscle action potential (CMAP).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## AMIFAMPRIDINE PHOSPHATE (FIRDAPSE)

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### MEDICATION(S)

FIRDAPSE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another aminopyridine drug (e.g. dalfampridine, Ruzurgi).

History of seizures.

Allergy to aminopyridine.

### REQUIRED MEDICAL INFORMATION

Diagnosis confirmed by proximal muscle weakness and one of the following tests: positive anti-P/Q-type voltage-gated calcium channel (VGCC) antibody test OR compound muscle action potential (CMAP).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## APALUTAMIDE (ERLEADA)

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### **MEDICATION(S)**

ERLEADA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## APOMORPHINE (APOKYN)

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### MEDICATION(S)

APOMORPHINE HCL

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Loss of control of body movements due to advanced Parkinson's disease (hypomobility):  
Treatment failure to at least one antiparkinsonian drug (i.e., pramipexole, entacapone, rasagiline, etc.).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## APREMILAST (OTEZLA)

---

### MEDICATION(S)

OTEZLA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Behcet's Disease: treatment failure or side effect with one of the following: triamcinolone dental paste (Kenalog, Oralone) or colchicine.

Plaque Psoriasis (PsO), initial use: treatment failure or side effect with one of the following: a DMARD, a topical corticosteroid (i.e., betamethasone), a calcineurin inhibitor (i.e., tacrolimus ointment), or calcipotriene OR has a medical reason why methotrexate, cyclosporine, acitretin, or topical agents cannot be used.

Psoriatic arthritis (PsA): treatment failure or side effect with one DMARD drug or medical reason why methotrexate, leflunomide, and sulfasalazine cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

PsO: Dermatologist or Rheumatologist. PsA: Rheumatologist.

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## APREPITANT 40MG CAPSULE (EMEND)

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### **MEDICATION(S)**

APREPITANT 40 MG CAP

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Prevention of post-surgery nausea and vomiting (PONV): patient cannot use other antiemetics (e.g. ondansetron, promethazine, transdermal scopolamine) prior to surgery because of history of treatment failure or side effects and dose will be given within 3 hours of surgery.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

PONV: once per surgery.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **ARIPIPRAZOLE (ABILIFY ASIMTUFII)**

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### **MEDICATION(S)**

ABILIFY ASIMTUFII

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## **ARIPIPRAZOLE (ABILIFY MAINTENA ER)**

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### **MEDICATION(S)**

ABILIFY MAINTENA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## **ARIPIPRAZOLE LAUROXIL (ARISTADA AND ARISTADA INITIO)**

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### **MEDICATION(S)**

ARISTADA, ARISTADA INITIO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Schizophrenia: Psychiatrist

### **COVERAGE DURATION**

Aristada: plan year

Aristada Initio: one time to start/restart Aristada treatment

### **OTHER CRITERIA**

Aristada Initio: single use along with oral aripiprazole

### **PART B PREREQUISITE**

N/A

## ARMODAFINIL (NUVIGIL)

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### MEDICATION(S)

ARMODAFINIL

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Bipolar disorder

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Narcolepsy: patient has positive sleep study (polysomnography) for narcolepsy. Trial and failure or side effect with modafinil or there is a medical reason why modafinil cannot be used. Obstructive sleep apnea/hypopnea syndrome (OSAHS): patient has a positive sleep study for OSAHS. Trial and failure or side effect with modafinil or there is a medical reason why modafinil cannot be used. Shift work sleep disorder: patient is a night shift worker working between 11pm and 7am OR patient is an early morning shift worker with starting hours between 4am and 7am OR patient is a rotating shift worker with night shifts. Trial and failure or side effect with modafinil or there is a medical reason why modafinil cannot be used. Bipolar Disorder: being added to current treatment regimen. Trial and failure or side effect with modafinil or there is a medical reason why modafinil cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Bipolar Disorder: Psychiatrist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

**PART B PREREQUISITE**

N/A

## ASCIMINIB (SCEMBLIX)

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### MEDICATION(S)

SCEMBLIX

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## **ASENAPINE (SAPHRIS)**

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### **MEDICATION(S)**

ASENAPINE MALEATE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial and failure or side effect with one preferred atypical antipsychotic agent (e.g. aripiprazole, olanzapine) or there is a medical reason why all the preferred agents cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **ASENAPINE (SECUADO)**

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### **MEDICATION(S)**

SECUADO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial and failure or side effect with one preferred atypical antipsychotic agent (e.g. aripiprazole, olanzapine) or there is a medical reason why all the preferred agents cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ASFOTASE ALFA (STRENSIQ)

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### MEDICATION(S)

STRENSIQ

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used for odonto- or pseudo- HPP or adult-onset HPP.

### REQUIRED MEDICAL INFORMATION

Documented history of one or more signs of HPP and lab test confirms low alkaline phosphatase (ALP) activity for age and gender.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

HPP: Endocrinologist, Geneticist, or Pediatric Specialist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ATOVAQUONE (MEPRON)

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### MEDICATION(S)

ATOVAQUONE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Toxoplasmosis prevention or treatment: patient is immunocompromised or at high risk of infection. Toxoplasmosis primary prevention: patient has failed or had a side effect to tmp/smx or has a medical reason (contraindication) for not using tmp/smx. PCP prevention or treatment: patient is immunocompromised or at high risk of infection and patient has failed or had a side effect to tmp/smx or has a medical reason (contraindication) for not using tmp/smx. Babesiosis treatment: active infection confirmed by blood smear test that is positive for Babesia microti parasites, PCR blood sample positive for Babesia microti DNA, or FISH test positive for Babesia microti RNA.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

PCP: 21days, Toxo: 6wks, Babesiosis: 10 days, PCP/Toxo prevention: Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## AVAPRITINIB (AYVAKIT)

---

### **MEDICATION(S)**

AYVAKIT

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## AXITINIB (INLYTA)

---

### MEDICATION(S)

INLYTA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## AZACITIDINE (ONUREG)

---

### **MEDICATION(S)**

ONUREG

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## AZTREONAM LYSINE (CAYSTON)

---

### **MEDICATION(S)**

CAYSTON

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used for acute treatment of an infection.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **BECAPLERMIN (REGRANEX)**

---

### **MEDICATION(S)**

REGRANEX

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Treating pressure ulcers or venous stasis ulcers.

### **REQUIRED MEDICAL INFORMATION**

Diabetic ulcer has not responded to standard therapy for wound management (i.e. debridement, dressing changes, pressure relief).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

3 months

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BEDAQUILINE (SIRTURO)**

---

### **MEDICATION(S)**

SIRTURO

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

PLAN YEAR

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BELIMUMAB (BENLYSTA)**

---

### **MEDICATION(S)**

BENLYSTA 200 MG/ML SOLN A-INJ, BENLYSTA 200 MG/ML SOLN PRSYR

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Diagnosis is severe CNS lupus. Benlysta is being used with Rituxan, other biologics, or IV cyclophosphamide.

### **REQUIRED MEDICAL INFORMATION**

Systemic Lupus Erythematosus (SLE) initial use: patient is currently taking one or more of the following: prednisone, methylprednisolone, azathioprine, methotrexate, mycophenolate, chloroquine, hydroxychloroquine. Ongoing use requires prescriber statement to show patients clinical status has not worsened and patient has benefited from treatment as evidenced by one or more of the following: less number or severity of SLE flares, daily steroid dose has been lowered, or improvement in physician global assessment.

Lupus Nephritis: being added to standard SLE therapy (e.g. corticosteroids, immunomodulators).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

SLE: Rheumatologist

Lupus Nephritis: Rheumatologist or Nephrologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## **BELUMOSUDIL (REZUROCK)**

---

### **MEDICATION(S)**

REZUROCK

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BELZUTIFAN (WELIREG)**

---

### **MEDICATION(S)**

WELIREG

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan Year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BEXAROTENE (TARGRETIN TOPICAL GEL)**

---

### **MEDICATION(S)**

BEXAROTENE 1 % GEL

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BEXAROTENE CAPSULE (TARGRETIN)**

---

### **MEDICATION(S)**

BEXAROTENE 75 MG CAP

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **BINIMETINIB (MEKTOVI)**

---

### **MEDICATION(S)**

MEKTOVI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BOSENTAN (TRACLEER)**

---

### **MEDICATION(S)**

BOSENTAN, TRACLEER 32 MG TAB SOL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Confirmation of Pulmonary Arterial Hypertension (WHO Group I) by right heart catheterization test.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BOSUTINIB (BOSULIF)**

---

### **MEDICATION(S)**

BOSULIF

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

First line therapy for CML and ALL: medical reason why imatinib cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BREXPIRAZOLE (REXULTI)**

---

### **MEDICATION(S)**

REXULTI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Depression: being used as a single agent

### **REQUIRED MEDICAL INFORMATION**

Trial of aripiprazole or medical reason for not using aripiprazole.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## BRIGATINIB (ALUNBRIG)

---

### MEDICATION(S)

ALUNBRIG

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **BUDESONIDE (ENTOCORT EC)**

---

### **MEDICATION(S)**

BUDESONIDE 3 MG CP DR PART

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Microscopic colitis (aka lymphocytic and collagenous colitis), Autoimmune hepatitis

### **EXCLUSION CRITERIA**

Being used for severe Crohn's disease (CD). Autoimmune hepatitis: patient has liver cirrhosis.

### **REQUIRED MEDICAL INFORMATION**

CD, initial use: budesonide will be used to induce remission.

CD, ongoing use: prescriber states patient responded to initial therapy and needs to continue therapy to maintain remission and there is a medical reason why guideline supported therapies (e.g. infliximab, mesalamine, azathioprine) for maintaining CD remission cannot be used.

Autoimmune hepatitis: being used with azathioprine and has a medical reason not to use prednisone or prednisolone or had severe side effect to prednisone or prednisolone that is not also seen with budesonide and initial dose is not more than 9 mg per day.

Microscopic colitis: initial dose is not more than 9 mg per day. For ongoing use: responded to initial therapy but symptoms returned after therapy was completed and dose is not more than 6 mg per day.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Microscopic colitis: gastroenterologist or infectious disease specialist. Autoimmune hepatitis: gastroenterologist, hepatologist, or infectious disease specialist.

### **COVERAGE DURATION**

CD initial: adults 8 weeks. CD maintenance: 3 months. Autoimmune hepatitis: plan year

**OTHER CRITERIA**

Coverage duration: Microscopic colitis – initial: 8 weeks, ongoing use: plan year

**PART B PREREQUISITE**

N/A

## **BUDESONIDE ER TABLET (UCERIS)**

---

### **MEDICATION(S)**

BUDESONIDE ER

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Microscopic colitis (aka lymphocytic and collagenous colitis), Autoimmune hepatitis

### **EXCLUSION CRITERIA**

Autoimmune hepatitis: patient has liver cirrhosis.

### **REQUIRED MEDICAL INFORMATION**

Ulcerative colitis (UC): being used to start remission of active UC, and patient has tried or has a medical reason for not trying one drug from the mesalamine class (e.g. balsalazide, mesalamine), and for moderate disease, medical reason why patient cannot use a generic corticosteroid drug that is taken by mouth (e.g. prednisone, methylprednisolone, hydrocortisone, and dexamethasone).

Autoimmune hepatitis: being used with azathioprine has a medical reason not to use prednisone or prednisolone or had severe side effect to prednisone or prednisolone that is not also seen with budesonide and dose is not more than 9 mg per day.

Microscopic colitis: being used to start remission of symptoms and dose is not more than 9 mg per day. For ongoing use: responded to initial therapy but symptoms returned after therapy was completed.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

UC: Gastroenterologist. Microscopic colitis: Gastroenterologist, Infectious Disease.

Autoimmune hepatitis: Gastroenterologist, Hepatologist or Infectious Disease.

### **COVERAGE DURATION**

UC: 8 weeks. Autoimmune hepatitis: plan year.



**OTHER CRITERIA**

Coverage duration: Microscopic colitis – initial: 8 weeks, ongoing use: plan year

**PART B PREREQUISITE**

N/A

## **BUPRENORPHINE PATCH (BUTRANS) – NARCOTIC SAFETY INITIATIVE**

---

### **MEDICATION(S)**

BUPRENORPHINE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other long-acting narcotic drugs.

### **REQUIRED MEDICAL INFORMATION**

Cancer pain: dose has been consolidated to the least number of higher strength forms.

Non-cancer pain, initial: cause of pain cannot be removed or treated with other treatment options, and pain occurs daily and has lasted for at least 3 months, and pain is severe enough to need daily around-the-clock long-term narcotic use, and total daily dose across all narcotic drugs is less than 90 MME, and dose has been consolidated to the least number of higher strength forms and patient has tried at least one short-acting narcotic drug, and chart notes document pain history including baseline pain intensity score and functional interference score, a plan for monitoring side effects and misuse, and a plan to taper down narcotics.

Non-cancer pain, reauth: total daily dose across all narcotic drugs is less than 90 MME per day, and dose has been consolidated to the least number of higher strength forms, and chart notes document current pain intensity score, functional interference score, any side effects and/or misuse with current pain treatment regimen, and plan to taper narcotic use.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Cancer pain: Oncologist or Pain Specialist.

### **COVERAGE DURATION**

Cancer pain: plan year

Non-cancer pain: initial 30 days, 1st reauth 3mos, ongoing reauths plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## **BUPROPION XL (FORFIVO XL)**

---

### **MEDICATION(S)**

BUPROPION HCL ER (XL) 450 MG TAB ER 24H

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient currently on either bupropion immediate-release (Wellbutrin) or bupropion XL (generic Wellbutrin XL) and want to lower pill burden.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **BUROSUMAB-TWZA (CRYSVITA SQ)**

---

### **MEDICATION(S)**

CRYSVITA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Adult patients (18 years and older), initial use: patient has osteomalacia-related symptoms [e.g. spontaneous or unhealed bone breaks (fractures), elevated serum bone ALP] or skeletal pain that affects daily activities and not controlled with non-narcotic pain medication. Ongoing use: improvement in patients symptoms (e.g. healing of rickets, correction of leg deformities, or increase in height for children, healing of existing fractures or lower number of new fractures, less pain with daily activities, better mobility, or ALP is lower than prior lab result).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

XLH: Endocrinologist

TIO: Endocrinologist or oncologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## BUTALBITAL CONTAINING PRODUCTS

---

### **MEDICATION(S)**

ASCOMP-CODEINE, BAC, BUTALBITAL-ACETAMINOPHEN 50-300 MG CAP, BUTALBITAL-ACETAMINOPHEN 50-325 MG TAB, BUTALBITAL-APAP-CAFF-COD 50-325-40-30 MG CAP, BUTALBITAL-APAP-CAFFEINE, BUTALBITAL-ASA-CAFF-CODEINE, BUTALBITAL-ASPIRIN-CAFFEINE, ESGIC 50-325-40 MG CAP, FIORICET, ZEBUTAL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Tension Headache: trial of two prescription strength non-steroidal anti-inflammatory drugs (NSAIDs) and amount requested does not exceed the amount needed to treat the number of headache days per month and if 65 years of age and older, prescriber confirms the benefits of the drug outweigh any risks and will monitor for side effects.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

If more than 8 headache days per month: neurologist or headache or pain specialist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## C1 ESTERASE INHIBITOR (BERINERT)

---

### MEDICATION(S)

BERINERT

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **C1 ESTERASE INHIBITOR (CINRYZE)**

---

### **MEDICATION(S)**

CINRYZE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other Hereditary Angioedema (HAE) preventive therapies (e.g. danazol, Haegarda).

### **REQUIRED MEDICAL INFORMATION**

Prevention: chart documentation or labs that show C4 and C1-INH (antigenic or functional) levels confirm HAE type I or II, and prescriber states that patient has symptomatic disease.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **C1 ESTERASE INHIBITOR (HAEGARDA)**

---

### **MEDICATION(S)**

HAEGARDA

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other Hereditary Angioedema (HAE) preventive therapies (e.g. danazol, Cinryze).

### **REQUIRED MEDICAL INFORMATION**

Prevention: chart documentation or labs that show C4 and C1-INH (antigenic or functional) levels confirm HAE type I or II, and prescriber states that patient has symptomatic disease.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## C1 ESTERASE INHIBITOR (RUCONEST)

---

### **MEDICATION(S)**

RUCONEST

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CABOZANTINIB (CABOMETYX)**

---

### **MEDICATION(S)**

CABOMETYX

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CABOZANTINIB S-MALATE (COMETRIQ)**

---

### **MEDICATION(S)**

COMETRIQ (100 MG DAILY DOSE), COMETRIQ (140 MG DAILY DOSE), COMETRIQ (60 MG DAILY DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CALCIPOTRIENE/BETAMETHASONE OINTMENT (TACLONEX)**

---

### **MEDICATION(S)**

CALCIPOTRIENE-BETAMETH DIPROP 0.005-0.064 % OINTMENT

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial and failure of either calcipotriene or a topical steroid from the high or very high potency class.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CANAKINUMAB (ILARIS)

---

### **MEDICATION(S)**

ILARIS

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

SJIA and AOSD: Rheumatologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CANNABIDIOL (CBD) EXTRACT (EPIDIOLEX)**

---

### **MEDICATION(S)**

EPIDIOLEX

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Dravet syndrome: trial of one of the following anti-seizure drugs: valproic acid or divalproex, topiramate, and clobazam.

Lennox-Gastaut syndrome: trial of two of the following anti-seizure drugs: clonazepam, felbamate, lamotrigine, and topiramate.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CAPLACIZUMAB-YHDP (CABLIVI)

---

### **MEDICATION(S)**

CABLIVI

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Continuation from inpatient hospital Cablivi treatment.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

58 days for inpatient or 103 days for outpatient plasma exchange

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## CAPMATINIB (TABRECTA)

---

### MEDICATION(S)

TABRECTA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## CARGLUMIC ACID (CARBAGLU)

---

### **MEDICATION(S)**

CARGLUMIC ACID

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CARIPRAZINE HYDROCHLORIDE (VRAYLAR)

---

### MEDICATION(S)

VRAYLAR

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Major Depressive Disorder: Being used as single agent therapy.

### REQUIRED MEDICAL INFORMATION

Treatment failure or side effect to one preferred atypical antipsychotic agent (e.g. aripiprazole, clozapine, olanzapine, quetiapine, risperidone, ziprasidone).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## CASPOFUNGIN (CANCIDAS)

---

### MEDICATION(S)

CASPOFUNGIN ACETATE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Oropharyngeal or Esophageal Candidiasis: patient has tried fluconazole OR fungal culture confirms infection is resistant to azole antifungals.

Invasive Aspergillosis: patient has tried an oral or IV azole antifungal or fungal culture confirms infection is resistant to azole antifungals.

Antifungal prophylaxis in cancer patients at high risk of febrile neutropenia [e.g. due to chemotherapy regimen, AML/MDS patient, undergoing HCST]: patient has tried fluconazole, voriconazole, or posaconazole or has a medical reason (contraindications) to azole antifungals.

Pulmonary Aspergillosis: patient has tried itraconazole or voriconazole.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

2 months

### OTHER CRITERIA

Ongoing use: continued neutropenia, culture remains positive, or ongoing symptoms.

**PART B PREREQUISITE**

N/A

## **CENOBAMATE (XCOPRI)**

---

### **MEDICATION(S)**

XCOPRI, XCOPRI (250 MG DAILY DOSE), XCOPRI (350 MG DAILY DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Treatment failure or side effect with two preferred partial seizure drugs (e.g. carbamazepine, clonazepam, divalproex, felbamate, lamotrigine, levetiracetam, oxcarbazepine, phenytoin, topiramate, zonisamide) OR medical reason why the preferred partial seizure drugs cannot be used (contraindication).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CERITINIB (ZYKADIA)

---

### **MEDICATION(S)**

ZYKADIA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CHOLIC ACID (CHOLBAM)

---

### MEDICATION(S)

CHOLBAM

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## **CLOBAZAM (ONFI)**

---

### **MEDICATION(S)**

CLOBAZAM

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Dravet syndrome: trial of valproic acid or divalproex.

Lennox-Gastaut syndrome: trial of two of the following anti-seizure drugs: clonazepam, felbamate, lamotrigine, and topiramate.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CLOBAZAM ORAL FILM (SYMPAZAN)**

---

### **MEDICATION(S)**

SYMPAZAN

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Seizures due to Dravet Syndrome: Trial and failure or side effect with valproic acid or divalproex AND side effect to clobazam (Onfi) tablet that is not seen with Sympazan.  
Lennox-Gastaut Syndrome: side effect to clobazam (Onfi) tablet that is not seen with Sympazan.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CLOBETASOL FOAM (OLUX, OLUX-E)**

---

### **MEDICATION(S)**

CLOBETASOL PROPIONATE 0.05 % FOAM, CLOBETASOL PROPIONATE EMULSION, TOVET 0.05 % FOAM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried clobetasol 0.05% topical solution.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## CLOZAPINE SUSPENSION (VERSACLOZ)

---

### **MEDICATION(S)**

VERSACLOZ

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Parkinson's psychosis disorder

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has a medical reason not to use clozapine tablets.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## COBIMETINIB (COTELLIC)

---

### MEDICATION(S)

COTELLIC

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## CORTICOTROPIN (CORTROPHIN)

---

### **MEDICATION(S)**

CORTROPHIN

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Ongoing use for treating infantile spasm: medical records show continued diagnosis (e.g. EEG confirms ongoing spasm).

Multiple Sclerosis (MS): patient is on a maintenance drug for MS (e.g. Tecfidera, Betaseron, glatiramer, Gilenya, Aubagio) but has an acute flare up and has had a side effect or contraindication to corticosteroids that is not seen with the use of Cortrophin.

Idiopathic or lupus erythematosus associated nephrotic syndrome, first use: patient's condition has not gotten better while using at least one immunosuppressive drug (cyclophosphamide, cyclosporine, and mycophenolate), and patient's condition responded to corticosteroid therapy but has had a side effect with the therapy that would not be seen with the use of Cortrophin. Ongoing use requires prescriber statement that patient's condition has gotten better while using Cortrophin.

All other FDA approved indications, first use: patient has not seen improvement of symptoms despite trying at least one different FDA approved drug for the condition other than corticosteroids, and has had a side effect to corticosteroids that is not seen with the use of Cortrophin. Ongoing use requires prescriber statement that patient's condition has gotten better while using Cortrophin.

### **AGE RESTRICTION**

N/A

**PRESCRIBER RESTRICTION**

Infantile spasms: Neurologist or Neonatologist MS: Neurologist or MS specialist Idiopathic or lupus erythematosus associated nephrotic syndrome: Nephrologist Rheumatic disorders, Collagen diseases: Rheumatologist Skin Diseases: Dermatologist Eye diseases: Ophthalmologist Symptomatic sarcoidosis: Pulmonologist

**COVERAGE DURATION**

MS flare: 3 weeks, Other FDA approved uses: 1 month

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## CRIZOTINIB (XALKORI)

---

### **MEDICATION(S)**

XALKORI 200 MG CAP, XALKORI 250 MG CAP

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **CYSTEAMINE (CYSTAGON)**

---

### **MEDICATION(S)**

CYSTAGON

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CYSTEAMINE (CYSTARAN)**

---

### **MEDICATION(S)**

CYSTARAN

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **CYSTEAMINE DELAYED RELEASE (PROCYSBI)**

---

### **MEDICATION(S)**

PROCYSBI

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DABRAFENIB (TAFINLAR)

---

### MEDICATION(S)

TAFINLAR

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **DACOMITINIB (VIZIMPRO)**

---

### **MEDICATION(S)**

VIZIMPRO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **DALFAMPRIDINE EXTENDED-RELEASE TABLET (AMPYRA)**

---

### **MEDICATION(S)**

DALFAMPRIDINE ER

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Multiple sclerosis, initial use: 25-foot walking test score. Ongoing use: updated timed 25-foot walking test shows improvement from prior or baseline test.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Neurologist or Multiple Sclerosis specialist

### **COVERAGE DURATION**

Initial use: 3 months. Ongoing use: plan year.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **DAPSONE GEL (ACZONE)**

---

### **MEDICATION(S)**

DAPSONE 5 % GEL, DAPSONE 7.5 % GEL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried a topical retinoin agent (i.e., adapalene, tretinoin) AND either a topical benzoyl peroxide containing agent or a topical anti-infective (e.g., clindamycin).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **DARBEPOETIN ALFA (ARANESP)**

---

### **MEDICATION(S)**

ARANESP (ALBUMIN FREE)

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

low red blood cells (anemia) due to Myelodysplastic Syndrome (MDS), anemia in patients with cancer who are undergoing palliative treatment.

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Chronic kidney disease (CKD), initial: anemia confirmed by a hemoglobin (Hgb) is 10g/dL. Ongoing use: Hgb level of less than or equal to 11g/dl. Myelosuppressive chemo related anemia: Hgb is less than 10g/dl AND one of the following: patient is on chemo or completed last dose within last 8 wks or patient has multiple myeloma (MM) on Revlimid tx. MDS: Hgb is less than or equal to 10g/dL (symptomatic anemia), and EPO level is less than or equal to 500U/ml. Anemia in cancer patients undergoing palliative treatment: Hgb is less than or equal to 10g/dL. For all indications: target Hgb level has not been met or maintained with at least 8 weeks of max dose Retacrit OR patient has a medical reason (contraindication) not to use Retacrit OR had a side effect with Retacrit that is not seen with Aranesp OR patient has a religious belief that does not allow treatment with drugs that contain human albumin.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

MDS, CKD: 6 months. Anemia due to chemo: 12 weeks no more than 8 weeks after last dose of chemo



**OTHER CRITERIA**

Excluded under Part D if covered by Part B. Duration for anemia in cancer patients receiving palliative treatment: plan year.

**PART B PREREQUISITE**

N/A

## DAROLUTAMIDE (NUBEQA)

---

### **MEDICATION(S)**

NUBEQA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **DASATINIB (SPRYCEL)**

---

### **MEDICATION(S)**

SPRYCEL

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DECITABINE/CEDAZURIDINE (INQOVI)

---

### MEDICATION(S)

INQOVI

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DEFERIPRONE (FERRIPROX)

---

### MEDICATION(S)

DEFERIPRONE, FERRIPROX 100 MG/ML SOLUTION

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Treatment failure or side effect with deferasirox.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DENOSUMAB (PROLIA)

---

### MEDICATION(S)

PROLIA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with other osteoporosis drugs.

### REQUIRED MEDICAL INFORMATION

Treatment or prevention of postmenopausal osteoporosis in women OR to increase bone mass in men: one of the following: trial of a bisphosphonate (e.g. alendronate, ibandronate, risedronate, or zoledronic acid), OR side effect to bisphosphonate therapy that supports discontinuation, OR Patient is at very high risk of fracture by meeting at least one of the following: non-traumatic fracture while on bisphosphonate therapy, patient has experienced a recent fracture (within the past 12 months) or history of multiple fractures, patient experienced a fracture while on long-term glucocorticoid therapy, or T-score less than -3.0, or patient is at high risk for falls, or 10-year hip fracture probability of greater than 4.5% based on FRAX score, or 10-year major osteoporosis-related fracture probability greater than 30% based on FRAX score.

Glucocorticoid-induced osteoporosis: initiating or continuing long-term glucocorticoid treatment (e.g. prednisone, dexamethasone) and either has history of a non-traumatic fracture or is at high risk for fracture.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## DENOSUMAB (XGEVA)

---

### MEDICATION(S)

XGEVA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Bone metastases from solid tumors or multiple myeloma: documentation of metastatic bone disease by scan or x-ray. Treatment of high calcium due to cancer: patient tried intravenous bisphosphonate therapy (e.g. zoledronic acid, pamidronate) within the last 30 days but did not respond well enough or had a side effect.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## **DESONIDE 0.05% GEL (DESONATE)**

---

### **MEDICATION(S)**

DESONIDE 0.05 % GEL, DESRX

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial of desonide 0.05% cream, lotion, or ointment

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DEUTETRABENAZINE (AUSTEDO)

---

### MEDICATION(S)

AUSTEDO, AUSTEDO XR, AUSTEDO XR PATIENT TITRATION

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Tardive Dyskinesia: Neurologist or Psychiatrist

Huntington's Chorea: Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DEXAMETHASONE TABLET (HEMADY)

---

### MEDICATION(S)

HEMADY

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Medical reason why patient cannot use preferred dexamethasone tablet.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DEXTROMETHORPHAN HBR- BUPROPION HCL ER (AUVELITY)

---

### MEDICATION(S)

AUVELITY

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Treatment failure or side effect with at least two generic antidepressants (e.g., fluoxetine, bupropion, duloxetine, sertraline, venlafaxine, etc.).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DEXTROMETHORPHAN HYDROBROMIDE/QUINIDINE SULFATE (NUEDEXTA)

---

### MEDICATION(S)

NUEDEXTA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **DICLOFENAC TOPICAL GEL (SOLARAZE)**

---

### **MEDICATION(S)**

DICLOFENAC SODIUM 3 % GEL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Treatment failure or side effect with one preferred topical Actinic Keratosis agent (e.g. fluorouracil cream or solution, imiquimod cream) OR has a medical reason for not using all preferred topical Actinic Keratosis agents.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

90 days

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DIHYDROERGOTAMINE INJECTION (D.H.E. 45)

---

### **MEDICATION(S)**

DIHYDROERGOTAMINE MESYLATE 1 MG/ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another triptan or ergot-type drug.

### **REQUIRED MEDICAL INFORMATION**

Migraine Headache: total number of doses matches the amount needed to treat the number of headache days per month, and trial of at least two preferred triptans or has a medical reason (contraindication) for not using triptans, for more than 8 headache days per month: prescribed by a Neurologist or headache specialist and currently taking a migraine prevention drug OR has a contraindication to all of the following migraine prevention drugs: divalproex, valproate, topiramate, amitriptyline, venlafaxine, atenolol, and nadolol. Cluster Headache: total number of doses matches the amount needed to treat the number of headache days per month, and trial of sumatriptan and zolmitriptan, and currently on prophylactic drugs supported for preventing cluster headaches including prednisone, dexamethasone, verapamil, lithium, or topiramate, OR has a contraindication to the supported prophylactic drugs.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Cluster Headache: Neurologist or headache specialist.

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A



## DIHYDROERGOTAMINE MESYLATE (MIGRANAL NASAL SPRAY)

---

### MEDICATION(S)

DIHYDROERGOTAMINE MESYLATE 4 MG/ML SOLUTION

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another triptan or ergot-type drug.

### REQUIRED MEDICAL INFORMATION

Migraine Headache: total number of doses matches the amount needed to treat the number of headache days per month, and trial of at least two preferred triptans or has a medical reason (contraindication) for not using triptans.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **DIMETHYL FUMARATE (TECFIDERA)**

---

### **MEDICATION(S)**

DIMETHYL FUMARATE, DIMETHYL FUMARATE STARTER PACK

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other disease-modifying therapies for relapsing Multiple Sclerosis.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DOXYCYCLINE MONOHYDRATE (ORACEA)

---

### MEDICATION(S)

DOXYCYCLINE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Acne Rosacea: patient has tried and failed or had a side effect to any two of the following: a topical sulfacetamide sodium product, a topical metronidazole product, topical azelaic acid, and topical ivermectin OR there is a medical reason why patient cannot use all of these drugs.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## DRONABINOL

---

### **MEDICATION(S)**

DRONABINOL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## **DROXIDOPA (NORTHERA)**

---

### **MEDICATION(S)**

DROXIDOPA

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Orthostatic hypotension is caused by primary anatomic failure such as Parkinson's disease, multiple system neuropathy or pure autonomic failure, dopamine beta-hydroxylase deficiency, or non-diabetic autonomic neuropathy. For ongoing use: patient has had clinical improvement in symptoms (i.e. dizziness, lightheadedness, vision, weakness, fatigue, concentration, head/neck discomfort) or daily living activities.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Cardiologist or Neurologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DULAGLUTIDE (TRULICITY)

---

### MEDICATION(S)

TRULICITY

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another GLP-1 agent

### REQUIRED MEDICAL INFORMATION

Type 2 diabetes: Confirmed by one of the following tests: Hemoglobin A1C of 6.5% or higher, or fasting blood sugar (glucose) of at least 126 mg/dl , or blood glucose of greater than or equal to 200 mg/dl either by Oral Glucose Tolerance Test or Plasma (blood) test AND one of the following: trial and failure or side effect with metformin or medical reason why metformin cannot be used, or A1C of 7.5% or greater and being used with another diabetes drug, or patient has heart (cardiovascular) disease. or more than one risk factor for heart (cardiovascular) disease (i.e., high blood pressure, high cholesterol, chronic kidney disease, etc.).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Dose and duration is not more than the FDA labeled maximum.

### PART B PREREQUISITE

N/A

## DULOXETINE (DRIZALMA SPRINKLE)

---

### **MEDICATION(S)**

DRIZALMA SPRINKLE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical reason why patient is not able to use duloxetine delayed-release capsule.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## DUVELISIB (COPIKTRA)

---

### **MEDICATION(S)**

COPIKTRA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## ELACESTRANT (ORSERDU)

---

### **MEDICATION(S)**

ORSERDU

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **ELEXACAFTOR/TEZACAFTOR/IVACAFTOR (TRIKAFTA)**

---

### **MEDICATION(S)**

TRIKAFTA 100-50-75 & 150 MG TAB THPK, TRIKAFTA 50-25-37.5 & 75 MG TAB THPK

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another CFTR modulator agent (e.g. Kalydeco, Symdeko, Orkambi)

### **REQUIRED MEDICAL INFORMATION**

Documentation that confirms there is at least one CFTR gene mutation sensitive to Trikafta.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ELIGLUSTAT (CERDELGA)

---

### MEDICATION(S)

CERDELGA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another therapy that treats Type-1 Gauchers disease. Patient is an ultra-rapid CYP2D6 metabolizer.

### REQUIRED MEDICAL INFORMATION

Patient has at least one of the following: low red blood cell count (anemia) with a low hemoglobin for age and sex, low platelet count (thrombocytopenia) with a platelet count under 100,000 cells/mcl or bleeding episodes documented as being due to thrombocytopenia, evidence of bone disease, enlarged liver (hepatomegaly), enlarged spleen (splenomegaly), or clinical symptoms of abdominal pain, fatigue, impaired physical movements, malnutrition (cachexia), or bone pain.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ELTROMBOPAG OLAMINE (PROMACTA)

---

### **MEDICATION(S)**

PROMACTA

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Myelodysplastic syndrome (MDS)-related thrombocytopenia

### **EXCLUSION CRITERIA**

Chronic immune thrombocytopenia (ITP): being used with another thrombopoietin receptor agonist (TPO-RA). MDS: being used in high-risk MDS

### **REQUIRED MEDICAL INFORMATION**

Chronic Hepatitis C: on interferon-based therapy and platelet count is less than or equal to 75,000/mcl prior to therapy or falls to less than or equal to 50,000/mcl during therapy. Chronic ITP, initial: platelet count is less than 30,000/mcl, and patient had a side effect or did not respond well enough to one of the following treatments: corticosteroids, IVIG, anti-D, and splenectomy OR has a medical reason not to use (contraindication) corticosteroids, IVIG, and anti-D. Aplastic anemia: prior therapy did not work well enough and platelet count is less than 50,000 cells/mcl or being used with cyclosporine and antithymocyte globulin (ATG) therapy for initial treatment. Thrombocytopenia due to MDS: treatment failure or side effect to at least one supported first line therapy for low risk MDS (e.g. decitabine, cyclosporine, ATG, lenalidomide). Ongoing use: platelet count has improved since starting Promacta but is not more than 400,000 and for MDS only disease has not progressed to acute leukemia.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

ITP, initial: 3 months all other conditions: 6 months Ongoing use: 6 months

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## ENASIDENIB MESYLATE (IDHIFA)

---

### MEDICATION(S)

IDHIFA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ENCORAFENIB (BRAFTOVI)

---

### MEDICATION(S)

BRAFTOVI

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ENTRECTINIB (ROZLYTREK)

---

### **MEDICATION(S)**

ROZLYTREK 100 MG CAP, ROZLYTREK 200 MG CAP

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## ENZALUTAMIDE (XTANDI)

---

### **MEDICATION(S)**

XTANDI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## EPOETIN ALFA-EPBX (RETACRIT)

---

### MEDICATION(S)

RETACRIT

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

low red blood cells (anemia) due to Myelodysplastic Syndrome (MDS), anemia due to rheumatoid arthritis (RA), anemia due to ribavirin therapy, anemia in patients with cancer who are undergoing palliative treatment.

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Chronic kidney disease (CKD), initial use: anemia confirmed by a hemoglobin level of 10g/dL. Ongoing use: Hgb level of less than or equal to 11g/dl. Anemia due to cancer drug therapy (myelosuppressive chemotherapy) patient is on chemo or completed last dose within last 8 wks for patient has multiple myeloma (MM) on Revlimid tx, Hgb is less than 10g/dl. MDS: Hgb is less than or equal to 10g/dL (symptomatic anemia), and EPO level is less than or equal to 500U/ml. HIV: currently on zidovudine and Hgb is less than or equal to 10g/dl. Anemia prior to a planned surgery: Hgb is less than 13g/dl and patient is likely to have significant blood loss and need of blood transfusions during surgery. Anemia due to RA, initial: current Hgb less than or equal to 10 g/dL, and anemia is not acute or caused by correctable etiology (e.g. occult blood loss due to gastritis). Ongoing use: current Hgb is 12 g/dL or less. Hepatitis C on ribavirin therapy: at least a 3 g/dL drop in Hgb within one month on ribavirin, or Hgb is 12 g/dL or less or Hct is 36% or lower. Anemia in cancer patients undergoing palliative treatment: Hgb is less than or equal to 10g/dL.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

**COVERAGE DURATION**

MDS/CKD: 6mos Anemia d/t zidovudine & chemo: 12 wks Surgery: 1mo All other: refer to other criteria

**OTHER CRITERIA**

Coverage duration for anemia due to chemo: 12 weeks and no more than 8 weeks after last dose of chemo. Coverage duration for planned surgery: 1 month, RA: 6months, anemia due to ribavirin: duration of ribavirin therapy, anemia in cancer patients receiving palliative treatment: plan year. Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## ERDAFITINIB (BALVERSA)

---

### **MEDICATION(S)**

BALVERSA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ERENUMAB-AOOE (AIMOVIG)

---

### MEDICATION(S)

AIMOVIG

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Migraine headache prevention: documentation of 4 or more headache days per month, treatment failure or side effects with at least 2 preventive therapies from the following drug classes: beta blockers, antidepressants, anticonvulsants or there is a medical reason why the patient cannot use the AAN level A or B guideline endorsed preventive drugs.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ERLOTINIB (TARCEVA)

---

### **MEDICATION(S)**

ERLOTINIB HCL

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ESTROGEN PRODUCTS (HIGH RISK MEDICATION)

---

### MEDICATION(S)

DUAVEE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

To help with symptoms of dryness, discomfort, pain in the vaginal area due to menopause: patient has tried the safer drugs estradiol vaginal cream and estradiol vaginal ring (Estring), and prescriber confirms the benefits of the drug outweigh any risks and will monitor for side effects. Other FDA indications: prescriber confirms the benefits of the drug outweigh any risks and will monitor for side effects.

### AGE RESTRICTION

65 years and older. No prior authorization required for less than 65 years old.

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan Year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ETANERCEPT (ENBREL – KIT, SYRINGE, SURECLICK)

---

### MEDICATION(S)

ENBREL, ENBREL SURECLICK

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

hidradenitis suppurativa and graft vs host disease (GVHD)

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Spondyloarthritis (SpA): patient is not able to use NSAIDs due to history of GI bleed or ulcer OR patient has tried one RX strength NSAID in combination with a PPI and had GI side effects OR patient's condition did not respond to a trial of two different RX strength NSAIDs.

Plaque Psoriasis (PsO), initial use: patient has tried one DMARD or has a medical reason why MTX, cyclosporine, and acitretin cannot be used AND baseline PASI score 10 or more OR BSA 3% or more OR sensitive areas are involved OR disease affects daily living. PsO, ongoing use: PASI or BSA improved on Enbrel. Psoriatic Arthritis (PsA): patient has tried one DMARD or has a medical reason why MTX, leflunomide, and sulfasalazine cannot be used.

Rheumatoid Arthritis (RA) and Polyarticular Juvenile Idiopathic Arthritis (pJIA): Treatment failure or side effect with methotrexate or patient has a medical reason why MTX cannot be used.

Hidradenitis suppurativa (HS): Hurley stage II or III HS and failed or had a side effect with Humira failed or has a medical reason not to use Humira. For ongoing use: clinical response seen with use of Enbrel.

GVHD: treatment failure or side effect to injectable or oral corticosteroids (e.g. prednisone, methylprednisolone).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

RA, pJIA, PsA and AS: Rheumatologist. PsO: Dermatologist or Rheumatologist. HS: Dermatologist.



**COVERAGE DURATION**

PsO: initial 12 weeks, ongoing plan year,

HS: 6 months,

Other indications: plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## EVEROLIMUS (AFINITOR DISPERZ)

---

### MEDICATION(S)

EVEROLIMUS 2 MG TAB SOL, EVEROLIMUS 3 MG TAB SOL, EVEROLIMUS 5 MG TAB SOL

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## EVEROLIMUS (AFINITOR)

---

### **MEDICATION(S)**

EVEROLIMUS 10 MG TAB, EVEROLIMUS 2.5 MG TAB, EVEROLIMUS 5 MG TAB, EVEROLIMUS 7.5 MG TAB

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## EVEROLIMUS (ZORTRESS)

---

### **MEDICATION(S)**

EVEROLIMUS 0.25 MG TAB, EVEROLIMUS 0.5 MG TAB, EVEROLIMUS 0.75 MG TAB, EVEROLIMUS 1 MG TAB

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial of or medical reason for not using mycophenolate and tacrolimus.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## **EVOLOCUMAB (REPATHA)**

---

### **MEDICATION(S)**

REPATHA, REPATHA PUSHTRONEX SYSTEM, REPATHA SURECLICK

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Primary Hyperlipidemia [including Heterozygous Familial Hypercholesterolemia (HeFH) or reduction of death due to Cardiovascular Disease (CVD)]: current LDL cholesterol (LDL-C) is at or above 70mg/dl (or at or above 55mg/dl if prescriber states extreme risk for heart disease) on lipid lowering therapy (such as statins and/or ezetimibe), and being used with a high-intensity statin like atorvastatin 40-80mg or rosuvastatin 20-40mg unless patient cannot use statins due to a medical reason (contraindication) or is intolerant to statins as defined by statin related rhabdomyolysis or has had skeletal-related muscle symptoms with the use of two different statins.

Homozygous Familial Hypercholesterolemia (HoFH): a positive genetic test for LDL-R genetic mutations OR clinical evidence that confirms HoFH, current lipid-lowering regimen has not worked well enough and being used with other lipid lowering therapies (e.g. statins, ezetimibe, LDL apheresis).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

HoFH: Cardiologist or Endocrinologist

### **COVERAGE DURATION**

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## EXENATIDE (BYETTA)

---

### MEDICATION(S)

BYETTA 10 MCG PEN, BYETTA 5 MCG PEN

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another GLP-1 agent

### REQUIRED MEDICAL INFORMATION

Type 2 diabetes: Confirmed by one of the following tests: Hemoglobin A1C of 6.5% or higher, or fasting blood sugar (glucose) of at least 126 mg/dl , or blood glucose of greater than or equal to 200 mg/dl either by Oral Glucose Tolerance Test or Plasma (blood) test AND one of the following: trial and failure or side effect with metformin or medical reason why metformin cannot be used, or A1C of 7.5% or greater and being used with another diabetes drug.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Dose and duration is not more than the FDA labeled maximum.

### PART B PREREQUISITE

N/A

## FEDRATINIB (INREBIC)

---

### MEDICATION(S)

INREBIC

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Not being used with another agent that treats myelofibrosis.

### REQUIRED MEDICAL INFORMATION

Myelofibrosis: platelet count of at least 50,000 cells/mcl, trial and failure of Jakafi or has a medical reason for not using Jakafi (contraindication).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## FENFLURAMINE (FINTEPLA)

---

### MEDICATION(S)

FINTEPLA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Dravet Syndrome: Trial of one of the following anti-seizure drugs: valproic acid or divalproex, topiramate, and clobazam. Lennox-Gastaut syndrome: trial of two of the following anti-seizure drugs: clonazepam, felbamate, lamotrigine, and topiramate

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## FERRIC CITRATE (AURYXIA)

---

### MEDICATION(S)

AURYXIA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient has high blood phosphate levels and is on dialysis due to CKD AND Trial and failure or side effect to both calcium acetate (PhosLo) and sevelamer carbonate (Renvela), or medical reason why calcium acetate and sevelamer carbonate cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **FIDAXOMICIN (DIFICID)**

---

### **MEDICATION(S)**

DIFICID

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Clostridium difficile: evidence of current infection AND trial of oral vancomycin or has a medical reason why oral vancomycin cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

10 days

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## FILGRASTIM-SNDZ (ZARXIO)

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### MEDICATION(S)

ZARXIO

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Cyclic neutropenia, agranulocytosis, Febrile neutropenia, Drug-Induced neutropenia, Myelodysplastic Syndrome (MDS), AIDS - neutropenia

### EXCLUSION CRITERIA

Being used to treat low white bloods called neutrophils (neutropenia) due to autoimmune disorders, burn victims, or chronic infections.

### REQUIRED MEDICAL INFORMATION

Agranulocytosis, congenital neutropenia, cyclic neutropenia, or idiopathic neutropenia: neutropenia is recurring or does not go away and there is a history of recurring infections (e.g. multiple episodes of infections requiring antibiotics) or at least one hospitalization for an infection within the past year.

Febrile neutropenia, neutropenia due to HIV/AIDs, or neutropenia caused by drugs other than cancer drugs: no use of pegfilgrastim within the past 14 days and absolute neutrophil count (ANC) is less than 800/mm<sup>3</sup> or ANC is less than 1000/mm<sup>3</sup> with neutropenia expected to last more than 5 days.

Neutropenia due to cancer drug therapy: not being used with pegfilgrastim.

Neutropenia due to radiation therapy: not being used with pegfilgrastim.

Acute myeloid leukemia (AML): being used to prevent or reduce neutropenia due to use of cancer drug therapy.

MDS: Hgb less than 10 gm/dL AND EPO level less than or equal to 500 mU/mL AND One of the following: ANC is less than 800/mm<sup>3</sup> or ANC is less than 1000/mm<sup>3</sup> with neutropenia expected to last more than 5 days or being used with epoetin (e.g. Retacrit) to improve symptoms of low red blood cells (anemia).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

**COVERAGE DURATION**

Febrile neutropenia, peripheral blood cell collection: 2 mos. HIV: plan year. (see other criteria)

**OTHER CRITERIA**

Coverage duration:

Congenital, cyclic, idiopathic neutropenia and agranulocytosis: plan year Neutropenia due to cancer drug therapy and AML: duration of cancer drug therapy

Neutropenia due to radiation: duration of radiation therapy

MDS: 3 months

Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## **FINERENONE (KERENDIA)**

---

### **MEDICATION(S)**

KERENDIA

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Treatment failure or side effect to Farxiga or there is a medical reason why Farxiga cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## FINGOLIMOD HCL (GILENYA)

---

### **MEDICATION(S)**

FINGOLIMOD HCL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other disease-modifying therapies for relapsing Multiple Sclerosis.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan Year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **FLURANDRENOLIDE CREAM, LOTION, OINTMENT (CORDRAN, NOLIX)**

---

### **MEDICATION(S)**

FLURANDRENOLIDE 0.05 % LOTION, FLURANDRENOLIDE 0.05 % OINTMENT, NOLIX 0.05 % LOTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried two generically available topical corticosteroids in the same or higher potency class.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## FOSTAMATINIB (TAVALISSE)

---

### MEDICATION(S)

TAVALISSE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Chronic immune thrombocytopenia (ITP), initial use: platelet count is less than 30,000 cells/mcl, and patient has tried one of the following treatments: corticosteroids (e.g. prednisone), IVIG, anti-D, and splenectomy or has a medical reason not to use (contraindication) corticosteroids, IVIG, and anti-D. Ongoing use: patient's platelet count has increased from baseline.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **FUTIBATINIB (LYTGOBI)**

---

### **MEDICATION(S)**

LYTGOBI (12 MG DAILY DOSE), LYTGOBI (16 MG DAILY DOSE), LYTGOBI (20 MG DAILY DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## GANAXOLONE SUSPENSION (ZTALMY)

---

### MEDICATION(S)

ZTALMY

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

A genetic test confirms CDKL5 (cyclin-dependent kinase-like 5) deficiency disorder

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## GEFITINIB (IRESSA)

---

### MEDICATION(S)

GEFITINIB

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **GILTERITINIB FUMARATE (XOSPATA)**

---

### **MEDICATION(S)**

XOSPATA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## GLASDEGIB MALEATE (DAURISMO)

---

### MEDICATION(S)

DAURISMO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## GLATIRAMER (COPAXONE)

---

### **MEDICATION(S)**

COPAXONE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other disease-modifying therapies for relapsing Multiple Sclerosis.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## GLECAPREVIR/PIBRENTASVIR (MAVYRET)

---

### **MEDICATION(S)**

MAVYRET

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current AASLD/IDSA guidelines.

### **REQUIRED MEDICAL INFORMATION**

Required medical information will be aligned with current AASLD/IDSA guidelines.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Hepatologist, Gastroenterologist, or Infectious Disease.

### **COVERAGE DURATION**

Length of therapy will be based on current AASLD/IDSA guidelines and FDA labeling.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## GLYCEROL PHENYLBUTYRATE (RAVICTI)

---

### **MEDICATION(S)**

RAVICTI

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## GLYCOPYRROLATE SOLUTION (CUVPOSA)

---

### **MEDICATION(S)**

GLYCOPYRROLATE 1 MG/5ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## HIGH RISK MEDICATION

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### MEDICATION(S)

AMITRIPTYLINE HCL, CARBINOXAMINE MALEATE 4 MG TAB, CARBINOXAMINE MALEATE 4 MG/5ML SOLUTION, CARISOPRODOL, CARISOPRODOL-ASPIRIN, CARISOPRODOL-ASPIRIN-CODEINE, CHLORDIAZEPOXIDE HCL, CHLORDIAZEPOXIDE-AMITRIPTYLINE, CLEMASTINE FUMARATE 2.68 MG TAB, CLOMIPRAMINE HCL, CYCLOBENZAPRINE HCL, CYPROHEPTADINE HCL, DICYCLOMINE HCL 10 MG CAP, DICYCLOMINE HCL 10 MG/5ML SOLUTION, DICYCLOMINE HCL 20 MG TAB, DOXEPIN HCL 10 MG CAP, DOXEPIN HCL 10 MG/ML CONC, DOXEPIN HCL 100 MG CAP, DOXEPIN HCL 150 MG CAP, DOXEPIN HCL 25 MG CAP, DOXEPIN HCL 50 MG CAP, DOXEPIN HCL 75 MG CAP, FEXMID, HYDROXYZINE HCL 10 MG TAB, HYDROXYZINE HCL 10 MG/5ML SYRUP, HYDROXYZINE HCL 25 MG TAB, HYDROXYZINE HCL 50 MG TAB, HYDROXYZINE PAMOATE, IMIPRAMINE HCL, KETOROLAC TROMETHAMINE 10 MG TAB, METAXALL, METAXALONE, METHOCARBAMOL 500 MG TAB, METHOCARBAMOL 750 MG TAB, ORPHENADRINE CITRATE ER, PERPHENAZINE-AMITRIPTYLINE, PHENADOZ, PHENOBARBITAL, PROMETHAZINE HCL, PROMETHAZINE VC, PROMETHAZINE VC/CODEINE, PROMETHAZINE-CODEINE, PROMETHAZINE-DM, PROMETHAZINE-PHENYLEPH-CODEINE, PROMETHAZINE-PHENYLEPHRINE, PROMETHEGAN, PROPANTHELINE BROMIDE, SCOPOLAMINE, THIORIDAZINE HCL, TRIMIPRAMINE MALEATE, VANADOM

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Prescriber confirms the benefits of the drug outweigh any risks and will monitor for side effects.

### AGE RESTRICTION

65 years and older. No prior authorization required for less than 65 years old.

### PRESCRIBER RESTRICTION

N/A

**COVERAGE DURATION**

Plan Year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## **IBRUTINIB (IMBRUVICA)**

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### **MEDICATION(S)**

IMBRUVICA 140 MG CAP, IMBRUVICA 280 MG TAB, IMBRUVICA 420 MG TAB, IMBRUVICA 560 MG TAB, IMBRUVICA 70 MG CAP, IMBRUVICA 70 MG/ML SUSPENSION

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ICATIBANT (FIRAZYR)

---

### **MEDICATION(S)**

ICATIBANT ACETATE, SAJAZIR

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **IDELALISIB (ZYDELIG)**

---

### **MEDICATION(S)**

ZYDELIG

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ILOPERIDONE (FANAPT)

---

### **MEDICATION(S)**

FANAPT, FANAPT TITRATION PACK

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried one preferred atypical antipsychotic drug (e.g. aripiprazole, clozapine, olanzapine, quetiapine, risperidone, ziprasidone) or has a medical reason not to use preferred atypical antipsychotic drugs.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## IMATINIB MESYLATE (GLEEVEC)

---

### **MEDICATION(S)**

IMATINIB MESYLATE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Hypereosinophilic syndrome (HES): Allergist, Immunologist, or Hematologist.

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## IMIGLUCERASE (CEREZYME)

---

### MEDICATION(S)

CEREZYME

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another therapy that treats Type-1 Gaucher's disease.

### REQUIRED MEDICAL INFORMATION

Disease confirmed by either glucocerebrosidase enzyme activity in the white blood cells or skin fibroblasts less or equal to 30% of normal activity or genetic analysis identifying two copies of a mutant glucocerebrosidase encoding allele, AND patient has at least one of the following: low red blood cell count (anemia) with a low hemoglobin for age and sex, low platelet count (thrombocytopenia) with a platelet count under 100,000 cells/mcl or bleeding episodes documented as being due to thrombocytopenia, evidence of bone disease, enlarged liver (hepatomegaly), enlarged spleen (splenomegaly), or clinical symptoms of abdominal pain, fatigue, impaired physical movements, malnutrition (cachexia), or bone pain.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Excluded under Part D if covered by Part B.

### PART B PREREQUISITE

N/A

## **IMMUNE GLOBULIN, GAMM(IGG)/GLYCINE/GLUCOSE/IGA (GAMMAGARD)**

---

### **MEDICATION(S)**

GAMMAGARD, GAMMAGARD S/D LESS IGA

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Autoimmune mucocutaneous blistering disease (AMBD), Guillian-Barre syndrome, Bone marrow transplant, Autoimmune Hemolytic anemia, Multiple myeloma, Polymyositis and dermatomyositis, Solid organ transplants, Bone marrow transplants, Hemopoietic stem cell transplant, Small lymphocytic leukemia

### **EXCLUSION CRITERIA**

AMBD: being used with another immunomodulator

### **REQUIRED MEDICAL INFORMATION**

Primary Immunodeficiency Disorder (PIDD), SQ and IV administration: current IgG is less than 200mg/dL or ALL of the following: history of recurrent bacterial infections, and failure to respond to antigenic challenge test with diphtheria and tetanus toxoids or pneumococcal polysaccharide vaccine, and history of IgG less than 500mg/dL documented on two occasions or diagnosed by an allergist or immunologist. Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Multifocal Acquired Demyelinating Polyneuropathy, or pure sensory Chronic Inflammatory Demyelinating Polyneuropathy (CIDP): IV administration, diagnosis confirmed by electrodiagnostic criteria and two of the following criteria: motor or sensory dysfunction in more than one limb lasting at least 2 months, no reflexes (areflexia), nerve biopsy shows evidence of demyelination and remyelination, or CSF cell count is less than 10cells/mm<sup>3</sup> (if HIV positive then CSF count less than 50cells/mm<sup>3</sup>). Primary immune thrombocytopenia (ITP): IV administration, platelet count is less than 30,000cells/mm<sup>3</sup>. For ongoing use: continued thrombocytopenia with prior response to IVIG or is scheduled for surgery or invasive procedure. Myasthenia Gravis (MG): IV administration, treatment failure, side effect, or medical reason for not using one of the following: a corticosteroid, mycophenolate, azathioprine, cyclosporine, or cyclophosphamide. Multifocal Motor Neuropathy (MMN): IV administration and condition confirmed with nerve conduction studies (electrodiagnostic testing).

### **AGE RESTRICTION**

N/A

**PRESCRIBER RESTRICTION**

CIDP, MMN, MG: Neurologist

**COVERAGE DURATION**

MG: 3 months ITP: 6 months GBS: 5 days all other conditions: plan year

**OTHER CRITERIA**

AMBD (pemphigus, epidermolysis bullosa acquisita): IV administration, condition is confirmed by testing the sore or blister (lesional tissue biopsy or serology) and did not respond to trial of an immunosuppressant drug (e.g. azathioprine, cyclophosphamide) and an oral or IV corticosteroid (e.g. prednisone) or has a medical reason not to use these types of drugs.

Autoimmune hemolytic anemia, Polymyositis, or Dermatomyositis: IV administration, trial and failure of high dose corticosteroids. Bone marrow transplant or HSCT: IV administration, being used to prevent bacterial infections and one of the following: within 100 days post-transplant, immunoglobulin G (IgG) level is less than 400 mg/dl, IgG is below normal and chronic graft vs host disease (GVHD) on steroids or GVHD with lung infection, or has cytomegalovirus (CMV).

Chronic lymphocytic leukemia/small lymphocytic leukemia: history of hypogammaglobulinemia (IgG below 500 mg/dl) or recurrent bacterial infections. Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## IMMUNE GLOBULIN, GAMMA (IGG)/PROLINE/IGA (HIZENTRA)

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### MEDICATION(S)

HIZENTRA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Multifocal Acquired Demyelinating Polyneuropathy, or pure sensory Chronic Inflammatory Demyelinating Polyneuropathy (CIDP): diagnosis confirmed by electrodiagnostic criteria (nerve conduction studies), and patient has been started on IVIG and is switching to Hizentra for ongoing therapy.

Primary Immunodeficiency Disorder (PID): current IgG is less than 200mg/dL or ALL of the following: history of recurrent bacterial infections, and failure to respond to antigenic challenge test with diphtheria and tetanus toxoids or pneumococcal polysaccharide vaccine, and history of IgG less than 500mg/dL documented on two occasions or diagnosed by an allergist or immunologist.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

CIDP, Multifocal acquired Demyelinating Polyneuropathy, or pure sensory CIDP: Neurologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## IMMUNE GLOBULIN, GAMMA(IGG)/GLYCINE/IGA (GAMMAKED)

---

### MEDICATION(S)

GAMMAKED

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Autoimmune mucocutaneous blistering disease (AMBD), Guillian-Barre syndrome, Bone marrow transplant, Autoimmune Hemolytic anemia, Kawasaki disease, Multiple myeloma, Polymyositis and dermatomyositis, Solid organ transplants, Bone marrow transplants, Chronic lymphocytic leukemia, Small lymphocytic leukemia, Hemopoietic stem cell transplant

### EXCLUSION CRITERIA

AMBD: being used with another immunomodulator

### REQUIRED MEDICAL INFORMATION

Primary Immunodeficiency Disorder (PIDD), SQ and IV administration: current IgG is less than 200mg/dL or ALL of the following: history of recurrent bacterial infections, and failure to respond to antigenic challenge test with diphtheria and tetanus toxoids or pneumococcal polysaccharide vaccine, and history of IgG less than 500mg/dL documented on two occasions or diagnosed by an allergist or immunologist. Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Multifocal Acquired Demyelinating Polyneuropathy, or pure sensory Chronic Inflammatory Demyelinating Polyneuropathy (CIDP): IV administration, diagnosis confirmed by electrodiagnostic criteria and two of the following criteria: motor or sensory dysfunction in more than one limb lasting at least 2 months, no reflexes (areflexia), nerve biopsy shows evidence of demyelination and remyelination, or CSF cell count is less than 10cells/mm<sup>3</sup> (if HIV positive then CSF count less than 50cells/mm<sup>3</sup>). Primary immune thrombocytopenia (ITP): IV administration, platelet count is less than 30,000cells/mm<sup>3</sup>. For ongoing use: continued thrombocytopenia with prior response to IVIG or is scheduled for surgery or invasive procedure. Myasthenia Gravis (MG): IV administration, treatment failure, side effect, or medical reason for not using one of the following: a corticosteroid, mycophenolate, azathioprine, cyclosporine, or cyclophosphamide. Multifocal Motor Neuropathy (MMN): IV administration and condition confirmed with nerve conduction studies (electrodiagnostic testing).

### AGE RESTRICTION

N/A

**PRESCRIBER RESTRICTION**

CIDP, MMN, MG: Neurologist

**COVERAGE DURATION**

MG: 3 months, ITP: 6 months, GBS: 5 days, all other conditions: plan year

**OTHER CRITERIA**

AMBD (pemphigus, epidermolysis bullosa acquisita): IV administration, condition is confirmed by testing the sore or blister (lesional tissue biopsy or serology) and did not respond to trial of an immunosuppressant drug (e.g. azathioprine, cyclophosphamide) and an oral or IV corticosteroid (e.g. prednisone) or has a medical reason not to use these types of drugs.

Autoimmune hemolytic anemia, Polymyositis, or Dermatomyositis: IV administration, trial and failure of high dose corticosteroids. Bone marrow transplant or HSCT: IV administration, being used to prevent bacterial infections and one of the following: within 100 days post-transplant, immunoglobulin G (IgG) level is less than 400 mg/dl, IgG is below normal and chronic graft vs host disease (GVHD) on steroids or GVHD with lung infection, or has cytomegalovirus (CMV).

Chronic lymphocytic leukemia/small lymphocytic leukemia: history of hypogammaglobulinemia (IgG below 500 mg/dl) or recurrent bacterial infections. Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A



## IMMUNE GLOBULIN, GAMMA(IGG)/GLYCINE/IGA (GAMUNEX-C)

---

### MEDICATION(S)

GAMUNEX-C

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Autoimmune mucocutaneous blistering disease (AMBD), Guillian-Barre syndrome, Bone marrow transplant, Autoimmune Hemolytic anemia, Kawasaki disease, Multiple myeloma, Polymyositis and dermatomyositis, Solid organ transplants, Bone marrow transplants, Chronic lymphocytic leukemia, Small lymphocytic leukemia, Hemopoietic stem cell transplant

### EXCLUSION CRITERIA

AMBD: being used with another immunomodulator

### REQUIRED MEDICAL INFORMATION

Primary Immunodeficiency Disorder (PIDD), SQ and IV administration: current IgG is less than 200mg/dL or ALL of the following: history of recurrent bacterial infections, and failure to respond to antigenic challenge test with diphtheria and tetanus toxoids or pneumococcal polysaccharide vaccine, and history of IgG less than 500mg/dL documented on two occasions or diagnosed by an allergist or immunologist.

Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Multifocal Acquired Demyelinating Polyneuropathy, or pure sensory Chronic Inflammatory Demyelinating Polyneuropathy (CIDP): IV administration, diagnosis confirmed by electrodiagnostic criteria and two of the following criteria: motor or sensory dysfunction in more than one limb lasting at least 2 months, no reflexes (areflexia), nerve biopsy shows evidence of demyelination and remyelination, or CSF cell count is less than 10cells/mm<sup>3</sup> (if HIV positive then CSF count less than 50cells/mm<sup>3</sup>).

Primary immune thrombocytopenia (ITP): IV administration, platelet count is less than 30,000cells/mm<sup>3</sup>. For ongoing use: continued thrombocytopenia with prior response to IVIG or is scheduled for surgery or invasive procedure.

Myasthenia Gravis (MG): IV administration, treatment failure, side effect, or medical reason for not using one of the following: a corticosteroid, mycophenolate, azathioprine, cyclosporine, or cyclophosphamide.

Multifocal Motor Neuropathy (MMN): IV administration and condition confirmed with nerve conduction studies (electrodiagnostic testing).

**AGE RESTRICTION**

N/A

**PRESCRIBER RESTRICTION**

CIDP, MMN, MG: Neurologist

**COVERAGE DURATION**

MG: 3 months ITP: 6 months GBS: 5 days all other conditions: plan year

**OTHER CRITERIA**

AMBD (pemphigus, epidermolysis bullosa acquisita): IV administration, condition is confirmed by testing the sore or blister (lesional tissue biopsy or serology) and did not respond to trial of an immunosuppressant drug (e.g. azathioprine, cyclophosphamide) and an oral or IV corticosteroid (e.g. prednisone) or has a medical reason not to use these types of drugs.

Autoimmune hemolytic anemia, Polymyositis, or Dermatomyositis: IV administration, trial and failure of high dose corticosteroids.

Bone marrow transplant or HSCT: IV administration, being used to prevent bacterial infections and one of the following: within 100 days post-transplant, immunoglobulin G (IgG) level is less than 400 mg/dl, IgG is below normal and chronic graft vs host disease (GVHD) on steroids or GVHD with lung infection, or has cytomegalovirus (CMV).

Chronic lymphocytic leukemia/small lymphocytic leukemia: history of hypogammaglobulinemia (IgG below 500 mg/dl) or recurrent bacterial infections.

Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## INFIGRANTINIB (TRUSELTIQ)

---

### **MEDICATION(S)**

TRUSELTIQ (100MG DAILY DOSE), TRUSELTIQ (125MG DAILY DOSE), TRUSELTIQ (50MG DAILY DOSE), TRUSELTIQ (75MG DAILY DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## INTERFERON BETA-1B (BETASERON)

---

### **MEDICATION(S)**

BETASERON

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other disease-modifying therapies for relapsing Multiple Sclerosis.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## INTERFERON GAMMA-1B (ACTIMMUNE)

---

### MEDICATION(S)

ACTIMMUNE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## INTRAVENOUS IMMUNE GLOBULIN (IVIG)

---

### **MEDICATION(S)**

BIVIGAM, CARIMUNE NF, FLEBOGAMMA DIF, GAMMAPLEX, PRIVIGEN

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Autoimmune mucocutaneous blistering disease (AMBD), Guillian-Barre syndrome, Bone marrow transplant, Autoimmune Hemolytic anemia, Multiple myeloma, Polymyositis and dermatomyositis, Solid organ transplants, Bone marrow transplants, Hemopoietic stem cell transplant, Small lymphocytic leukemia

### **EXCLUSION CRITERIA**

AMBD: being used with another immunomodulator

### **REQUIRED MEDICAL INFORMATION**

Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Multifocal Acquired Demyelinating Polyneuropathy, or pure sensory Chronic Inflammatory Demyelinating Polyneuropathy (CIDP): diagnosis confirmed by electrodiagnostic criteria and two of the following criteria: motor or sensory dysfunction in more than one limb lasting at least 2 months, no reflexes (areflexia), nerve biopsy shows evidence of demyelination and remyelination, or CSF cell count is less than 10cells/mm<sup>3</sup> (if HIV positive then CSF count less than 50cells/mm<sup>3</sup>).

Primary Immune Thrombocytopenia (ITP): platelet count is less than 30,000cells/mm<sup>3</sup>. For ongoing use: continued thrombocytopenia with prior response to IVIG or is scheduled for surgery or invasive procedure.

Myasthenia Gravis (MG): treatment failure, side effect, or medical reason for not using one of the following: a corticosteroid, mycophenolate, azathioprine, cyclosporine, or cyclophosphamide.

Primary Immunodeficiency Disorder (PIDD): current IgG is less than 200mg/dL or ALL of the following: history of recurrent bacterial infections, and failure to respond to antigenic challenge test with diphtheria and tetanus toxoids or pneumococcal polysaccharide vaccine, and history of IgG less than 500mg/dL documented on two occasions or diagnosed by an allergist or immunologist.

Multifocal Motor Neuropathy (MMN): condition confirmed by nerve conduction studies (electrodiagnostic testing)

**AGE RESTRICTION**

N/A

**PRESCRIBER RESTRICTION**

CIDP, MMN, MG: Neurologist

**COVERAGE DURATION**

MG: 3 months ITP: 6 months GBS: 5 days all other conditions: plan year

**OTHER CRITERIA**

AMBD (pemphigus, epidermolysis bullosa acquisita): condition is confirmed by testing the sore or blister (lesional tissue biopsy or serology) and did not respond to trial of an immunosuppressant drug (e.g. azathioprine, cyclophosphamide) and an oral or IV corticosteroid (e.g. prednisone) or has a medical reason not to use these types of drugs.

Autoimmune hemolytic anemia, Polymyositis, or Dermatomyositis: trial and failure of high dose corticosteroids.

Bone marrow transplant or HSCT: being used to prevent bacterial infections and one of the following: within 100 days post-transplant, immunoglobulin G (IgG) level is less than 400 mg/dl, IgG is below normal and chronic graft vs host disease (GVHD) on steroids or GVHD with lung infection, or has cytomegalovirus (CMV).

Chronic lymphocytic leukemia/small lymphocytic leukemia: history of hypogammaglobulinemia (IgG below 500 mg/dl) or recurrent bacterial infections.

Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## ISAVUCONAZONIUM (CRESEMBA)

---

### MEDICATION(S)

CRESEMBA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Invasive aspergillosis: blood or tissue culture positive for Aspergillus, and patient has a medical reason for not using voriconazole.

Invasive mucormycosis: culture is positive for mucormycosis pathogens (e.g. Rhizopus, Rhizomucor, Lichtheimia, Mucormycetes) or being prescribed by infectious disease specialist.

Esophageal candidiasis: patient has HIV infection and patient has a medical reason for not using oral fluconazole.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

3 months

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## ITRACONAZOLE (SPORANOX) B

---

### MEDICATION(S)

ITRACONAZOLE

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Systemic infections due to sporotrichosis (cutaneous, lymphonodular, osteoarticular, pulmonary, disseminated, or meningeal), coccidiomycosis, cryptococcosis, tinea corporis, cruris, pedis, manuum, capitis, versicolor, and unguium (onychomycosis), allergic bronchopulmonary aspergillosis (ABPA). Prophylaxis (primary or secondary) or maintenance treatment of talaromycosis (*Talaromyces marneffe*). treatment of pulmonary aspergillosis, chronic (cavitary or necrotizing). Prophylaxis of aspergillosis and histoplasmosis.

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Blastomycosis, Histoplasmosis, Sporotrichosis, Cryptococcosis, or Aspergillosis infection: culture confirms infection.

Tinea Capitis: patient has tried or has a medical reason for not using oral terbinafine.

Tinea Corpus, Curis, Pedis or Manuum: patient has tried or has a medical reason for not using topical antifungal or oral terbinafine.

Tinea Versicolor: patient has tried or has a medical reason for not using topical ketoconazole or oral fluconazole.

Onychomycosis: patient has tried or has a medical reason for not using oral terbinafine.

Coccidioidomycosis culture confirms infection, and patient has tried or has a medical reason for not using fluconazole.

Aspergillosis or Histoplasmosis prevention: patient is immunosuppressed/compromised.

Prophylaxis (primary or secondary) or maintenance treatment of talaromycosis (*Talaromyces marneffe*): Patient with HIV infection.

For oral solution and treatment other than oropharyngeal or esophageal candidiasis: patient has difficulty swallowing a capsule or tablet.

### AGE RESTRICTION

N/A

**PRESCRIBER RESTRICTION**

N/A

**COVERAGE DURATION**

T. Vesicolor: 1wk T. Capitis: 8wks Onyc: 3mo Other Tinea: 1mo ABPA: 4mo All other dx: plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## IVABRADINE (CORLANOR)

---

### MEDICATION(S)

CORLANOR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Left heart ventricular ejection fraction (LVEF) less than or equal to 35%, patient is in sinus rhythm with resting heart rate of at least 70 beats per minute, and patient is on the highest tolerated dose of guideline supported therapies including a renin-angiotensin inhibitor drug (e.g. ACE-Inhibitor, ARB agent, Entresto) and beta-blocker drug (e.g. bisoprolol, carvedilol, metoprolol succinate) unless there is a medical reason for not using (contraindication) the supported therapies. Pediatric patients: CHF is due to dilated cardiomyopathy.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## IVACFTOR (KALYDECO)

---

### **MEDICATION(S)**

KALYDECO

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## IVERMECTIN CREAM (SOOLANTRA)

---

### **MEDICATION(S)**

IVERMECTIN 1 % CREAM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial and failure or side effect to two of the following: topical metronidazole, oral tetracycline, oral azithromycin, and oral isotretinoin OR medical reason why all the first line acne rosacea drugs cannot be used

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## IVOSIDENIB (TIBSOVO)

---

### **MEDICATION(S)**

TIBSOVO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## IXAZOMIB CITRATE (NINLARO)

---

### **MEDICATION(S)**

NINLARO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **IXEKIZUMAB (TALTZ)**

---

### **MEDICATION(S)**

TALTZ

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another targeted immunotherapy drug.

### **REQUIRED MEDICAL INFORMATION**

Spondyloarthritis (SpA): patient is not able to take NSAIDs due to history of GI bleed or ulcer  
OR patient has tried one RX strength NSAID in combination with a PPI and had GI side effects  
OR patient's condition did not respond to a trial of two different RX strength NSAIDs.

Plaque Psoriasis (PsO), initial use: patient has tried one DMARD or has a medical reason why methotrexate (MTX), cyclosporine, and acitretin cannot be used AND baseline PASI score 10 or more OR BSA 3% or more OR sensitive areas are involved OR disease affects daily living. PSO, ongoing use: PASI or BSA improved on Taltz.

Psoriatic Arthritis (PsA): patient has tried one DMARD or has a medical reason why MTX, leflunomide, and sulfasalazine cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

PsA: Rheumatologist. PsO: Rheumatologist or Dermatologist.

### **COVERAGE DURATION**

PsO initial: 12 weeks, PsO ongoing: plan year, all other indications: plan year

### **OTHER CRITERIA**

N/A



**PART B PREREQUISITE**

N/A

## LANREOTIDE ACETATE (SOMATULINE DEPOT)

---

### **MEDICATION(S)**

LANREOTIDE ACETATE, SOMATULINE DEPOT 60 MG/0.2ML SOLUTION, SOMATULINE DEPOT 90 MG/0.3ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Acromegaly: Endocrinologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## LAPATINIB DITOSYLATE (TYKERB)

---

### **MEDICATION(S)**

LAPATINIB DITOSYLATE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LAROTRECTINIB SULFATE (VITRAKVI)

---

### **MEDICATION(S)**

VITRAKVI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LEDIPASVIR/SOFOSBUVIR (HARVONI)

---

### **MEDICATION(S)**

HARVONI, LEDIPASVIR-SOFOSBUVIR

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current AASLD/IDSA guidelines.

### **REQUIRED MEDICAL INFORMATION**

Required medical information will be aligned with current AASLD/IDSA guidelines.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Hepatologist, Gastroenterologist, or Infectious Disease.

### **COVERAGE DURATION**

Length of therapy will be based on current AASLD/IDSA guidelines and FDA labeling.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LENALIDOMIDE (REVLIMID)

---

### **MEDICATION(S)**

LENALIDOMIDE, REVLIMID

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

MDS: transfusion dependent or hemoglobin less than 10 g/dL confirming anemia associated disease.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LENVATINIB (LENVIMA)

---

### MEDICATION(S)

LENVIMA (10 MG DAILY DOSE), LENVIMA (12 MG DAILY DOSE), LENVIMA (14 MG DAILY DOSE), LENVIMA (18 MG DAILY DOSE), LENVIMA (20 MG DAILY DOSE), LENVIMA (24 MG DAILY DOSE), LENVIMA (4 MG DAILY DOSE), LENVIMA (8 MG DAILY DOSE)

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## LEVALBUTEROL SOLUTION (XOPENEX)

---

### MEDICATION(S)

LEVALBUTEROL HCL

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient has had a side effect with albuterol nebulized solution (not MDI or oral syrup) that is not seen with the use of levalbuterol.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Excluded under Part D if covered by Part B.

### PART B PREREQUISITE

N/A



## LEVETIRACETAM (SPRITAM)

---

### **MEDICATION(S)**

SPRITAM

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical reason why patient is not able to use generic levetiracetam oral solution and tablet.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LEVOMILNACIPRAN HCL (FETZIMA)

---

### **MEDICATION(S)**

FETZIMA, FETZIMA TITRATION

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Treatment failure or side effect with at least two formulary drugs that treat major depressive disorder (MDD).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LIDOCAINE PATCH (LIDODERM)

---

### **MEDICATION(S)**

LIDOCAINE 5 % PATCH

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LINEZOLID ORAL (ZYVOX)

---

### MEDICATION(S)

LINEZOLID 100 MG/5ML RECON SUSP, LINEZOLID 600 MG TAB

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

VRE, MRSA, or VISA skin or soft tissue infection confirmed by culture and sensitivity (C&S): treatment failure or side effect with one oral drug noted on the C&S to work on the bacteria causing the infection or recommended by an Infectious Disease (ID) specialist. MSSA skin or soft tissue infection: recommended by an ID specialist and treatment failure or side effect with two preferred oral drugs noted on the C&S to work on the bacteria causing the infection or medical reason why the preferred drugs cannot be used. Empiric therapy for suspected MRSA infection: prescribed or recommended by an ID specialist OR trial of one oral antibiotic supported for MRSA including clindamycin, doxycycline, or minocycline, and double strength trimethoprim/sulfamethoxazole, OR medical reason why all oral antibiotics supported for MRSA empiric therapy cannot be used. Infection of the bone or joint OR infective endocarditis: culture and sensitivity report confirm VRE, MRSA, or VISA/VRSA and prescribed or recommended by ID specialist. Multidrug-resistant tuberculosis infection (MDR-TB): being used with pretomanid and bedaquiline.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

VRE 28 days. Osteo 42 days. Endocarditis 56 days. MDR-TB 26 wks. Empiric tx/pneumonia/SSTI 14days.

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## **LIRAGLUTIDE (VICTOZA)**

---

### **MEDICATION(S)**

VICTOZA

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another GLP-1 agent

### **REQUIRED MEDICAL INFORMATION**

Type 2 diabetes: Confirmed by one of the following tests: Hemoglobin A1C of 6.5% or higher, or fasting blood sugar (glucose) of at least 126 mg/dl , or blood glucose of greater than or equal to 200 mg/dl either by Oral Glucose Tolerance Test or Plasma (blood) test AND one of the following: trial and failure or side effect with metformin or medical reason why metformin cannot be used, or A1C of 7.5% or greater and being used with another diabetes drug, or patient has heart (cardiovascular) disease.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Dose and duration is not more than the FDA labeled maximum.

### **PART B PREREQUISITE**

N/A

## LOMITAPIDE MESYLATE (JUXTAPID)

---

### MEDICATION(S)

JUXTAPID

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Homozygous Familial Hypercholesterolemia (HoFH): diagnosis confirmed by a genetic test confirming LDL-R genetic mutations or clinical evidence supporting HoFH, AND being used with a standard lipid lowering regimen (i.e. a maximally tolerated statin, a non-statin lipid lowering drug for patient with documented statin intolerance) AND patient has tried and failed or has a medical reason for not using Repatha.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Cardiologist or Endocrinologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## LONG-ACTING NARCOTIC DRUGS (NARCOTIC SAFETY INITIATIVE)

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### MEDICATION(S)

FENTANYL 100 MCG/HR PATCH 72HR, FENTANYL 12 MCG/HR PATCH 72HR, FENTANYL 25 MCG/HR PATCH 72HR, FENTANYL 50 MCG/HR PATCH 72HR, FENTANYL 75 MCG/HR PATCH 72HR, HYDROMORPHONE HCL ER, METHADONE HCL 10 MG TAB, METHADONE HCL 10 MG/5ML SOLUTION, METHADONE HCL 10 MG/ML CONC, METHADONE HCL 10 MG/ML SOLUTION, METHADONE HCL 5 MG TAB, METHADONE HCL 5 MG/5ML SOLUTION, METHADONE HCL INTENSOL, OXYCODONE HCL ER, OXYMORPHONE HCL ER

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with other long-acting narcotic drugs.

### REQUIRED MEDICAL INFORMATION

Cancer pain: dose has been consolidated to the least number of higher strength forms. Non-cancer pain, initial: cause of pain cannot be removed or treated with other treatment options, and pain occurs daily and has lasted for at least 3 months, and pain is severe enough to need daily around-the-clock long-term narcotic use, and total daily dose across all narcotic drugs is less than 90 MME, and dose has been consolidated to the least number of higher strength forms and trial of at least one short-acting and morphine sulfate ER tablet (MS Contin), and chart notes document pain history including baseline pain intensity score and functional interference score, a plan for monitoring side effects and misuse, and a plan to taper down narcotics. Non-cancer pain, reauth: total daily dose across all narcotic drugs is less than 90 MME per day, and dose has been consolidated to the least number of higher strength forms, and chart notes document current pain intensity score, functional interference score, any side effects and/or misuse with current pain treatment regimen, and plan to taper narcotic use.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Cancer pain: Oncologist or Pain Specialist.



**COVERAGE DURATION**

Cancer pain: plan year

Non-cancer pain: initial 30 days, 1st reauth 3mos, ongoing reauths plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## LORLATINIB (LORBRENA)

---

### **MEDICATION(S)**

LORBRENA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## LUMATEPERONE (CAPLYTA)

---

### MEDICATION(S)

CAPLYTA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Schizophrenia: Trial and failure or side effect with one of the following generic atypical antipsychotics: aripiprazole, clozapine, olanzapine, quetiapine, risperidone, ziprasidone OR there is a medical reason for not using the preferred atypical antipsychotics (contraindication). Bipolar Disorder: Trial and failure or side effect with one of the following generic atypical antipsychotics: aripiprazole, olanzapine, quetiapine, risperidone, ziprasidone OR there is a medical reason for not using the generic atypical antipsychotics (contraindication).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## MACITENTAN (OPSUMIT)

---

### MEDICATION(S)

OPSUMIT

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Confirmation of Pulmonary Arterial Hypertension (WHO Group I) by right heart catheterization test.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## MECASERMIN (INCRELEX)

---

### MEDICATION(S)

INCRELEX

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Severe primary IGF-1 deficiency: being used with growth hormone therapy.

### REQUIRED MEDICAL INFORMATION

Initial use: height is at or more than 3.0 standard deviations below standard range for sex and age, and basal IGF-1 is at or more than 3.0 standard deviations below standard range for sex and age, and evidence of delayed bone age, and for severe IGF-1 deficiency growth hormone level is normal or higher for sex and age. Ongoing use: response to therapy and evidence of delayed bone age.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Endocrinologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## MECHLORETHAMINE (VALCHLOR)

---

### MEDICATION(S)

VALCHLOR

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## MEGESTROL ACETATE (MEGACE)

---

### MEDICATION(S)

MEGESTROL ACETATE 20 MG TAB, MEGESTROL ACETATE 40 MG TAB, MEGESTROL ACETATE 40 MG/ML SUSPENSION, MEGESTROL ACETATE 400 MG/10ML SUSPENSION, MEGESTROL ACETATE 800 MG/20ML SUSPENSION

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## MEGESTROL ACETATE ES (MEGACE ES)

---

### MEDICATION(S)

MEGESTROL ACETATE 625 MG/5ML SUSPENSION

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient has tried megestrol acetate 200mg/5ml oral suspension.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## MEPOLIZUMAB (NUCALA)

---

### MEDICATION(S)

NUCALA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Eosinophilic asthma: being used with other targeted therapies for asthma treatment (e.g. Xolair, Cinqair, Dupixent, Fasenra).

### REQUIRED MEDICAL INFORMATION

Eosinophilic asthma, initial use: eosinophil blood count is 150 cells/microliter or more within last six weeks documented treatment failure with recent use of high-dose inhaled corticosteroid along with long-acting beta agonist or leukotriene receptor antagonists (i.e., montelukast), and patient has had at least one of the following within the past year: one or more acute asthma-related ER visit(s), one or more acute inpatient visits where asthma was the diagnosis, or two or more acute asthma exacerbations that require oral corticosteroids. Ongoing use: asthma symptoms improved and/or controlled while on Nucala. Eosinophilic granulomatosis with polyangiitis (EGPA): patients condition did not improve or has relapsed despite treatment with an oral corticosteroid and/or immunosuppressive therapy (e.g. azathioprine, methotrexate, mycophenolic acid). Hypereosinophilic syndrome (HES): patient is negative for FIP1-like 1-platelet derived growth factor receptor (FIP1L1-PDGFR) mutation and patient had an inadequate response to oral corticosteroids or hydroxyurea. Chronic rhinosinusitis with nasal polyps (CRSwNP): Treatment failure with an intranasal corticosteroid or medical rationale why intranasal corticosteroids cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Eosinophilic asthma: Immunologist, Pulmonologist, or Allergist. EGPA: Immunologist, Rheumatologist. HES: Immunologist, Allergist, Hematologist. CRSwNP: allergist, immunologist, or otolaryngologist.

**COVERAGE DURATION**

Eosinophilic asthma initial use: 6 mo, ongoing use: plan year. All other Dx: plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## MERCAPTOPURINE MONOHYDRATE (PURIXAN)

---

### **MEDICATION(S)**

PURIXAN

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical reason why patient cannot use mercaptopurine tablet.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## METHOTREXATE ORAL SOLUTION (XATMEP)

---

### **MEDICATION(S)**

XATMEP

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical reason why patient cannot take tablet form of methotrexate.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## METHYLNALTREXONE (RELISTOR SQ)

---

### **MEDICATION(S)**

RELISTOR 12 MG/0.6ML SOLUTION, RELISTOR 8 MG/0.4ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Constipation due to ongoing use of opioids for non-cancer pain: patient has tried lubiprostone (Amitiza) or Movantik.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **METHYLTESTOSTERONE (ANDROID, TESTRED)**

---

### **MEDICATION(S)**

METHYLTESTOSTERONE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **METOCLOPRAMIDE ODT (METOZOLV ODT)**

---

### **MEDICATION(S)**

METOCLOPRAMIDE HCL 10 MG TAB DISP, METOCLOPRAMIDE HCL 5 MG TAB DISP

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical condition that prevents swallowing a tablet.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## METRELEPTIN (MYALEPT)

---

### MEDICATION(S)

MYALEPT

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used for HIV-related Lipodystrophy, general obesity not associated with congenital leptin deficiency.

### REQUIRED MEDICAL INFORMATION

Generalized Lipodystrophy: Patient has one of the following: diabetes mellitus (HbA1c greater than 7% or FPG values 126 mg/dL or greater), or hypertriglyceridemia (TG 250mg/dL or greater), or history of pancreatitis associated with hypertriglyceridemia, or increased fasting insulin (greater than 25mIU/L)

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## MIDOSTAURIN (RYDAPT)

---

### **MEDICATION(S)**

RYDAPT

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **MIFEPRISTONE (KORLYM)**

---

### **MEDICATION(S)**

KORLYM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **MIGLUSTAT (ZAVESCA)**

---

### **MEDICATION(S)**

MIGLUSTAT

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used in with another therapy for Gauchers disease type-1.

### **REQUIRED MEDICAL INFORMATION**

Patient has at least one of the following: low red blood cell count (anemia) with a low hemoglobin for age and sex, low platelet count (thrombocytopenia) with a platelet count under 100,000 cells/mcl or bleeding episodes documented as being due to thrombocytopenia, evidence of bone disease, enlarged liver (hepatomegaly), enlarged spleen (splenomegaly), or clinical symptoms of abdominal pain, fatigue, impaired physical movements, malnutrition (cachexia), or bone pain.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **MOBOCERTINIB (EXKIVITY)**

---

### **MEDICATION(S)**

EXKIVITY

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## MODAFINIL (PROVIGIL)

---

### MEDICATION(S)

MODAFINIL

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Narcolepsy: sleep study (polysomnography) confirms narcolepsy.

Obstructive Sleep Apnea/Hypopnea Syndrome (OSAHS): sleep study (polysomnography) confirms OSAHS.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **MOMELOTINIB (OJJAARA)**

---

### **MEDICATION(S)**

OJJAARA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **NALOXONE AUTO-INJECTOR (EVZIO)**

---

### **MEDICATION(S)**

NALOXONE HCL 2 MG/0.4ML SOLN A-INJ

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Side effect or contraindication to naloxone prefilled syringe and nasal spray that is not also seen with naloxone auto-injector.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **NATALIZUMAB (TYSABRI)**

---

### **MEDICATION(S)**

TYSABRI

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

RRMS: Tysabri is being used in combination with another disease-modifying therapy for MS.  
Crohn's Disease: Tysabri is being used with immunosuppressants, other targeted immunotherapies, or anakinra.

### **REQUIRED MEDICAL INFORMATION**

Relapsing forms of Multiple Sclerosis: patient has tried at least one of the following therapies: Betaseron, glatiramer 20mg, Copaxone, Aubagio, Gilenya, Tecfidera.  
Crohn's Disease, initial use: patient tried and failed a TNF-blocker (e.g. Humira) or has a medical reason why TNF-blockers (e.g. Humira) cannot be used. Ongoing use: patient's condition has improved while on Tysabri.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

MS: plan year

Crohn's Disease: initial use: 3 months, ongoing use: plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A



## NERATINIB (NERLYNX)

---

### MEDICATION(S)

NERLYNX

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## NILOTINIB (TASIGNA)

---

### MEDICATION(S)

TASIGNA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

First line therapy for CML and ALL: medical reason why imatinib cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **NINTEDANIB ESYLATE (OFEV)**

---

### **MEDICATION(S)**

OFEV

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Idiopathic Pulmonary Fibrosis (IPF): being used with another IPF drug

### **REQUIRED MEDICAL INFORMATION**

IPF, initial use: patient has the following pulmonary function tests: forced vital capacity (FVC) equal or over 50%, and diffusing capacity of carbon monoxide (DLCO) equal or over 30%.

Ongoing use: patient has not received a lung transplant, sslLD (Systemic sclerosis interstitial lung disease); trial of at least one guideline supported therapy (i.e. cyclophosphamide, mycophenolate, azathioprine, or systemic glucocorticoids ) or medical reason why they cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

IPF: Pulmonologist. ILD: Rheumatologist or Pulmonologist.

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **NIRAPARIB TOSYLATE (ZEJULA)**

---

### **MEDICATION(S)**

ZEJULA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## NITAZOXANIDE (ALINIA)

---

### MEDICATION(S)

ALINIA 100 MG/5ML RECON SUSP, NITAZOXANIDE

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Cryptosporidiosis in HIV+ patients, Clostridium difficile colitis, viral gastroenteritis, amebiasis (Entamoeba histolytica), liver fluke infection (Fasciola hepatica), Cestode (tapeworm)

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

All medically accepted conditions: stool culture results confirm diagnosis.

Giardiasis: treatment failure or side effect with metronidazole OR medical reason for not using metronidazole (contraindication).

Clostridium difficile colitis: treatment failure or side effect with vancomycin OR medical reason for not using vancomycin (contraindication).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

One course (see other criteria)

### OTHER CRITERIA

Giardiasis: 3 days

Cryptosporidiosis: 3 days unless HIV+ then 14 days

Clostridium difficile colitis: 10 days

viral gastroenteritis: 3 days amebiasis (Entamoeba histolytica): 3 days

liver fluke infection (Fasciola hepatica): 7 days

Cestode (tapeworm): 3 days

**PART B PREREQUISITE**

N/A

## **NITISINONE (ORFADIN)**

---

### **MEDICATION(S)**

NITISINONE 10 MG CAP, NITISINONE 2 MG CAP, NITISINONE 5 MG CAP

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan Year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## NITISINONE (NITYR)

---

### MEDICATION(S)

NITYR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## NON-FORMULARY HIGH RISK MEDICATION

---

**MEDICATION(S)**

ESGIC 50-325-40 MG CAP

**PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

**OFF LABEL USES**

N/A

**EXCLUSION CRITERIA**

N/A

**REQUIRED MEDICAL INFORMATION**

Prescriber's supporting statement states the drug is medically necessary because all covered drugs on the formulary would not be as effective or would cause side effects.

**AGE RESTRICTION**

N/A

**PRESCRIBER RESTRICTION**

N/A

**COVERAGE DURATION**

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## NON-PREFERRED DOXYCYCLINE PRODUCTS

---

### MEDICATION(S)

DOXYCYCLINE HYCLATE 100 MG TAB DR, DOXYCYCLINE HYCLATE 150 MG TAB,  
DOXYCYCLINE HYCLATE 150 MG TAB DR, DOXYCYCLINE HYCLATE 200 MG TAB DR,  
DOXYCYCLINE HYCLATE 50 MG TAB DR, DOXYCYCLINE HYCLATE 75 MG TAB,  
DOXYCYCLINE HYCLATE 75 MG TAB DR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Side effect with preferred doxycycline that does not occur with the use of the non-preferred doxycycline product.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## NON-PREFERRED HYDROCODONE-ACETAMINOPHEN PRODUCTS

---

### MEDICATION(S)

HYDROCODONE-ACETAMINOPHEN 10-300 MG TAB, HYDROCODONE-ACETAMINOPHEN 5-300 MG TAB, HYDROCODONE-ACETAMINOPHEN 7.5-300 MG TAB

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with other short-acting opioid narcotics.

### REQUIRED MEDICAL INFORMATION

Moderate to severe pain, initial use: side effect to the preferred hydrocodone-acetaminophen product that would not occur with the non-preferred hydrocodone-acetaminophen product and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain. Ongoing use (more than 90 days of narcotic therapy): evaluation by pain specialist, documented patient-specific treatment plan for evaluating ongoing need for narcotic pain relief, potential misuse, and monitoring plan for adverse side effects, and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

30 days

### OTHER CRITERIA

N/A

**PART B PREREQUISITE**

N/A

## OCTREOTIDE ACETATE (SANDOSTATIN LAR DEPOT)

---

### MEDICATION(S)

SANDOSTATIN LAR DEPOT

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

AIDS-associated diarrhea, Bleeding esophageal varices, Chemotherapy-induced diarrhea, Intestinal obstruction, Neuroendocrine Tumor of the lung, Pituitary adenoma, Prevention of postoperative complications of pancreatic surgery, Pancreatic tumors (gastrinoma, glucagonoma, insulinoma), Radiation-induced diarrhea, Recurrent Meningioma, Thymoma, Zollinger-Ellison syndrome

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Diarrhea due to HIV: patient has been on anti-retroviral therapy (ART) for at least one month, and other causes (i.e. infection, underlying GI disease, malabsorption) have been ruled out, and treatment failure or side effect with diphenoxylate/atropine or loperamide. Intestinal obstruction: intestinal obstruction is due to cancer.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Acromegaly: Endocrinologist

### COVERAGE DURATION

Acromegaly: plan year

Other conditions: 6 months

### OTHER CRITERIA

Excluded under Part D if covered by Part B.

**PART B PREREQUISITE**

N/A

## OCTREOTIDE ACETATE (SANDOSTATIN)

---

### MEDICATION(S)

OCTREOTIDE ACETATE

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

AIDS-associated diarrhea, Bleeding esophageal varices, Chemotherapy-induced diarrhea, Cryptosporidiosis, Dumping syndrome, Intestinal obstruction, Neuroendocrine Tumor of the GI tract, lung, or thymus, Lymphorrhagia, Pancreatitis, necrotizing Pituitary adenoma, Prevention of postoperative complications of pancreatic surgery, Pancreatic tumors (gastrinoma, glucagonoma, insulinoma), paraganglioma, pheochromocytoma, Polycystic Ovary Syndrome (PCOS), Radiation-induced diarrhea, Recurrent Meningioma, Thymoma, Zollinger-Ellison syndrome.

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Diarrhea due to HIV: patient has been on anti-retroviral therapy (ART) for at least one month, and prescriber states other causes (i.e. infection, underlying GI disease, malabsorption) have been ruled out, and patient has tried diphenoxylate/atropine or loperamide. Intestinal obstruction: intestinal obstruction is due to cancer.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Acromegaly: Endocrinologist

### COVERAGE DURATION

Acromegaly: plan year, Other conditions: 6 months

### OTHER CRITERIA

N/A

**PART B PREREQUISITE**

N/A



## ODEVIXIBAT (BYLVAY)

---

### **MEDICATION(S)**

BYLVAY, BYLVAY (PELLETS)

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Gastroenterologist or Hepatologist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## OLANZAPINE PAMOATE (ZYPREXA RELPREVV)

---

### **MEDICATION(S)**

ZYPREXA RELPREVV

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (e.g. risperidone, ziprasidone) and medical reason why injectable risperidone (Risperdal Consta) cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## OLANZAPINE-SAMIDORPHAN (LYBALVI)

---

### **MEDICATION(S)**

LYBALVI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried at least one oral atypical antipsychotic (i.e. risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole) or there is a medical reason why all oral atypical antipsychotics cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## OLAPARIB (LYNPARZA)

---

### **MEDICATION(S)**

LYNPARZA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## OLSALAZINE SODIUM (DIPENTUM)

---

### **MEDICATION(S)**

DIPENTUM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Trial of one of the following: mesalamine 0.375g long-acting capsule, mesalamine 1.2g long-acting tablet, or balsalazide 750 mg capsule OR medical reason why these drugs cannot be used (contraindication).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## OLUTASIDENIB (REZLIDHIA)

---

### **MEDICATION(S)**

REZLIDHIA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## OMALIZUMAB (XOLAIR)

---

### MEDICATION(S)

XOLAIR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Allergic asthma: being used with other targeted therapies for asthma treatment (e.g. Nucala, Cinqair, Dupixent, Fasenra).

### REQUIRED MEDICAL INFORMATION

Allergic Asthma, initial use: recent total serum IgE level is more than 30IU/ml, documented treatment failure with recent use of high-dose inhaled corticosteroid along with long-acting beta agonist or leukotriene receptor antagonists (i.e., montelukast), and patient has had at least one of the following within the past year: one or more acute asthma-related ER visit(s), one or more acute inpatient visits where asthma was the diagnosis, or two or more acute asthma exacerbations that require oral corticosteroids. Ongoing use: asthma symptoms improved and/or controlled while on Xolair.

Chronic Idiopathic Urticaria (CIU): failure to respond to hydroxyzine, doxepin, or high dose second-generation antihistamines OR has a medical reason not to use (contraindication) or had a side effect to hydroxyzine, doxepin, and second-generation antihistamines.

Nasal polyps: treatment failure or side effect with a nasal corticosteroid (e.g. fluticasone)

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

CIU: Allergist or Immunologist Allergic asthma: Pulmonologist or Immunologist

Nasal polyps: allergist, immunologist, or otolaryngologist.

### COVERAGE DURATION

Allergic asthma, initial 6 months, ongoing use: plan year

All other Dx: plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A



## ORITAVANCIN DIPHOSPHATE (ORBACTIV)

---

### MEDICATION(S)

ORBACTIV

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Medical reason why oral antibiotics cannot be used AND culture and sensitivity report confirm vancomycin-resistant staphylococcus aureus (VRSA), vancomycin-insensitive staphylococcus aureus (VISA), or methicillin-resistant staphylococcus aureus (MRSA) and patient has an allergy or contraindication to vancomycin.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Infectious Disease Specialist

### COVERAGE DURATION

One treatment course.

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## OSIMERTINIB (TAGRISSO)

---

### MEDICATION(S)

TAGRISSO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## OSPEMIFENE (OSPHENA)

---

### MEDICATION(S)

OSPHENA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Painful sex (dyspareunia) due to menopause: patient has tried Premarin Vaginal cream.  
Vaginal dryness due to menopause: patient has tried at least two of the following: Premarin vaginal cream, estradiol vaginal cream, estradiol vaginal tablet, Yuvaferm, or Estring.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **OXYMORPHONE IMMEDIATE-RELEASE (OPANA)**

---

### **MEDICATION(S)**

OXYMORPHONE HCL

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other short-acting opioid narcotics.

### **REQUIRED MEDICAL INFORMATION**

Moderate to severe pain, initial use: trial of at least two preferred short-acting narcotic drugs, and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain. Ongoing use (more than 90 days of narcotic therapy): evaluation by pain specialist, documented patient-specific treatment plan for evaluating ongoing need for narcotic pain relief, potential misuse, and monitoring plan for adverse side effects, and total dose across all narcotics is not more than 90 MME/day or if more than 90 MME/day the prescriber states the total dose is medically necessary to treat the pain.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

30 days

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **PACRITINIB (VONJO)**

---

### **MEDICATION(S)**

VONJO

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used along with another agent for myelofibrosis

### **REQUIRED MEDICAL INFORMATION**

Platelet count is less than 50,000 cells/mcl.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **PALBOCICLIB (IBRANCE)**

---

### **MEDICATION(S)**

IBRANCE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **PALIPERIDONE ER (INVEGA)**

---

### **MEDICATION(S)**

PALIPERIDONE ER

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Trial of risperidone and one other preferred atypical antipsychotic such as aripiprazole, ziprasidone, quetiapine, or olanzapine.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **PALIPERIDONE PALMITATE (INVEGA HAFYERA)**

---

### **MEDICATION(S)**

INVEGA HAFYERA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Use of a once-a-month paliperidone palmitate ER injection (e.g., INVEGA SUSTENNA) for at least 4 months OR an every-three-month paliperidone palmitate extended release injectable suspension (e.g., INVEGA TRINZA) for at least one three-month cycle.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A



## **PALIPERIDONE PALMITATE (INVEGA SUSTENNA)**

---

### **MEDICATION(S)**

INVEGA SUSTENNA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## **PALIPERIDONE PALMITATE (INVEGA TRINZA)**

---

### **MEDICATION(S)**

INVEGA TRINZA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Older adults (65 years and older) with dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Treatment failure with at least one oral atypical antipsychotic (risperidone, ziprasidone, quetiapine, olanzapine, aripiprazole) and use of Invega Sustenna for at least 4 months.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## PARATHYROID HORMONE (NATPARA)

---

### MEDICATION(S)

NATPARA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Hypoparathyroidism: Lab tests confirm low blood calcium (hypocalcemia).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## PART D VS PART B

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### MEDICATION(S)

ABELCET, ACETYLCYSTEINE 10 % SOLUTION, ACETYLCYSTEINE 20 % SOLUTION, ACYCLOVIR SODIUM, ALBUTEROL SULFATE (2.5 MG/3ML) 0.083% NEBU SOLN, ALBUTEROL SULFATE (5 MG/ML) 0.5% NEBU SOLN, ALBUTEROL SULFATE 0.63 MG/3ML NEBU SOLN, ALBUTEROL SULFATE 1.25 MG/3ML NEBU SOLN, ALBUTEROL SULFATE 2.5 MG/0.5ML NEBU SOLN, ALDURAZYME, ALKERAN 2 MG TAB, AMINOSYN II 10 % SOLUTION, AMINOSYN-PF 10 % SOLUTION, AMPHOTERICIN B, AMPHOTERICIN B LIPOSOME, APREPITANT 125 MG CAP, APREPITANT 80 & 125 MG CAP, APREPITANT 80 & 125 MG MISC, APREPITANT 80 MG CAP, APRETUDE, ARALAST NP, ARFORMOTEROL TARTRATE, AZASAN, AZATHIOPRINE, AZATHIOPRINE SODIUM, BUDESONIDE 0.25 MG/2ML SUSPENSION, BUDESONIDE 0.5 MG/2ML SUSPENSION, BUDESONIDE 1 MG/2ML SUSPENSION, CABENUVA, CALCITRIOL 0.25 MCG CAP, CALCITRIOL 0.5 MCG CAP, CALCITRIOL 1 MCG/ML SOLUTION, CINACALCET HCL, CROMOLYN SODIUM 20 MG/2ML NEBU SOLN, CYCLOPHOSPHAMIDE 25 MG CAP, CYCLOPHOSPHAMIDE 25 MG TAB, CYCLOPHOSPHAMIDE 50 MG CAP, CYCLOPHOSPHAMIDE 50 MG TAB, CYCLOSPORINE 100 MG CAP, CYCLOSPORINE 25 MG CAP, CYCLOSPORINE 50 MG/ML SOLUTION, CYCLOSPORINE MODIFIED, DEXAMETHASONE SOD PHOSPHATE PF 10 MG/ML SOLUTION, DEXAMETHASONE SODIUM PHOSPHATE 10 MG/ML SOLUTION, DEXAMETHASONE SODIUM PHOSPHATE 100 MG/10ML SOLUTION, DOXERCALCIFEROL, ELAPRASE, ENGERIX-B, FABRAZYME, FORMOTEROL FUMARATE, GENGRAF, GLASSIA, GRANISETRON HCL, HEPARIN SODIUM (PORCINE) 1000 UNIT/ML SOLUTION, HEPARIN SODIUM (PORCINE) 10000 UNIT/ML SOLUTION, HEPARIN SODIUM (PORCINE) 20000 UNIT/ML SOLUTION, HEPARIN SODIUM (PORCINE) 5000 UNIT/ML SOLUTION, HEPATAMINE, HEPLISAV-B, IBANDRONATE SODIUM 3 MG/3ML SOLUTION, INTRALIPID, IPRATROPIUM BROMIDE 0.02 % SOLUTION, IPRATROPIUM-ALBUTEROL, KEPIVANCE 6.25 MG RECON SOLN, LACOSAMIDE 200 MG/20ML SOLUTION, MELPHALAN, METHOTREXATE SODIUM 1 GM RECON SOLN, METHOTREXATE SODIUM 1000 MG/40ML SOLUTION, METHOTREXATE SODIUM 250 MG/10ML SOLUTION, METHOTREXATE SODIUM 50 MG/2ML SOLUTION, METHOTREXATE SODIUM (PF), METHYLPREDNISOLONE SODIUM SUCC 125 MG RECON SOLN, MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL HCL, MYCOPHENOLATE SODIUM, NAGLAZYME, NUTRILIPID, ONDANSETRON, ONDANSETRON HCL 24 MG TAB, ONDANSETRON HCL 4 MG TAB, ONDANSETRON HCL 4 MG/5ML SOLUTION, ONDANSETRON HCL 8 MG TAB, PARICALCITOL, PENTAMIDINE ISETHIONATE, PREHEVBRIO, PREMASOL, PROLASTIN-C, PULMOZYME, RECOMBIVAX HB, RIBAVIRIN 6 GM RECON SOLN, SANDIMMUNE 100 MG/ML SOLUTION, SIROLIMUS, SMOFLIPID, SUNLENCA 463.5 MG/1.5ML SOLUTION, SYNRIPO, TACROLIMUS 0.5 MG CAP, TACROLIMUS 1 MG CAP, TACROLIMUS 5 MG CAP, THIOTEPA, TPN ELECTROLYTES, TRELSTAR MIXJECT, TWINRIX, VENTAVIS, VORICONAZOLE 200 MG RECON SOLN, ZOLEDRONIC ACID

**DETAILS**

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

## PASIREOTIDE (SIGNIFOR)

---

### MEDICATION(S)

SIGNIFOR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Cushings disease, initial use: pituitary surgery is not an option or has not been curative.  
Ongoing use: patient responded to initial treatment confirmed by a decrease in the mean 24-hour urinary free cortisol (UFC).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Initial: 3 months

Reauth: 1 year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## PAZOPANIB HCL (VOTRIENT)

---

### **MEDICATION(S)**

PAZOPANIB HCL, VOTRIENT

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## PEGINTERFERON ALFA-2A (PEGASYS)

---

### MEDICATION(S)

PEGASYS, PEGASYS PROCLICK

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

myelofibrosis, polycythemia vera, essential thrombocythemia, systemic mastocytosis

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Chronic hepatitis C viral infection: criteria will be applied consistent with current AASLD-IDSA guidance.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Hepatitis B: 48 weeks

Hepatitis C: based on AASLD-IDSA guidance

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## PEGVISOMANT (SOMAVERT)

---

### MEDICATION(S)

SOMAVERT

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient failed or is not a candidate for radiation or surgery AND failed treatment or had a side effect with octreotide or Somatuline.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Endocrinologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## PEMIGATINIB (PEMAZYRE)

---

### MEDICATION(S)

PEMAZYRE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **PENCICLOVIR (DENA VIR)**

---

### **MEDICATION(S)**

PENCICLOVIR

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## PENICILLAMINE (DEPEN)

---

### MEDICATION(S)

PENICILLAMINE 250 MG TAB

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Rheumatoid Arthritis: patient tried two of the following: methotrexate, sulfasalazine, hydroxychloroquine, or leflunomide, OR has a medical reason why methotrexate, hydroxychloroquine, sulfasalazine, and leflunomide cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## PEXIDARTINIB (TURALIO)

---

### **MEDICATION(S)**

TURALIO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **PIMAVANSERIN (NUPLAZID)**

---

### **MEDICATION(S)**

NUPLAZID

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used for dementia-related psychosis.

### **REQUIRED MEDICAL INFORMATION**

Evaluation by psychiatrist confirms Parkinson's disease psychosis (PDP) – symptoms of hallucinations (seeing, hearing, or experiencing things that others don't) and delusions (believing things that aren't true) due to Parkinson's disease.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Neurologist or Psychiatrist

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## PIRFENIDONE (ESBRIET)

---

### MEDICATION(S)

PIRFENIDONE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another IPF drug.

### REQUIRED MEDICAL INFORMATION

Idiopathic Pulmonary Fibrosis (IPF), initial use: patient has the following pulmonary function tests: forced vital capacity (FVC) equal or over 50%, and diffusing capacity of carbon monoxide (DLCO) equal or over 30%. Ongoing use: patient has not received a lung transplant.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Pulmonologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **PIRTOBRUTINIB (JAYPIRCA)**

---

### **MEDICATION(S)**

JAYPIRCA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## POMALIDOMIDE (POMALYST)

---

### MEDICATION(S)

POMALYST

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## PONATINIB (ICLUSIG)

---

### MEDICATION(S)

ICLUSIG

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

First line therapy for ALL: medical reason why imatinib cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## POSACONAZOLE (NOXAFIL)

---

### MEDICATION(S)

POSACONAZOLE 100 MG TAB DR, POSACONAZOLE 40 MG/ML SUSPENSION

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Esophageal candidiasis treatment, fusariosis, histoplasmosis, phaeohyphomycosis, Allergic Bronchopulmonary Aspergillosis (ABPA), refractory treatment of pulmonary aspergillosis, chronic (cavitary or necrotizing).

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Prevention of aspergillus or candida infection when there is high risk for developing these type of infections (e.g. weakened defense system due to cancer drug therapy, HIV, GVHD)

Aspergillosis, cryptococcosis, fusariosis, histoplasmosis, phaeohyphomycosis within the body that is confirmed by a positive culture test.

Treatment of candida infection of the esophagus, throat, mouth (esophageal or oropharyngeal candidiasis) after trial of fluconazole or there is a medical reason not to use fluconazole.

Treatment of candida infection within the body that is confirmed by a positive culture and failure of fluconazole or other anti-fungal shown by culture results to treat the infection.

ABPA: current use of chronic oral corticosteroid use after trial of itraconazole or there is a medical reason not to use itraconazole.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

oral or esophageal candidiasis: one month all other conditions: Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## PRALSETINIB (GAVRETO)

---

### MEDICATION(S)

GAVRETO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **PYRIMETHAMINE (DARAPRIM)**

---

### **MEDICATION(S)**

PYRIMETHAMINE

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Toxoplasmosis prevention, Toxoplasmosis chronic maintenance (secondary prophylaxis), Pneumocystis jiroveci (formerly Pneumocystis carinii) Pneumonia (PCP) prevention, Cystoisospora belli (formerly Isospora Belli) treatment or secondary prevention.

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Primary prevention of toxoplasmosis: treatment failure or side effect with trimethoprim-sulfamethoxazole (TMP-SMX) or has a medical reason for not using TMP-SMX and patient is immunocompromised (i.e. cancer, HIV+, post-transplantation).

Chronic maintenance (secondary prophylaxis) of toxoplasmosis: follows initial treatment in HIV-infected patients.

Prevention of Pneumocystis jiroveci (formerly Pneumocystis carinii) Pneumonia (PCP): treatment failure or side effect with trimethoprim-sulfamethoxazole (TMP-SMX) or has a medical reason for not using TMP-SMX AND patient is HIV+.

Treatment of cystoisospora belli (formerly Isospora Belli): Patient is HIV+ AND treatment failure or side effect with trimethoprim-sulfamethoxazole (TMP-SMX) or has a medical reason for not using TMP-SMX.

Chronic maintenance (secondary prophylaxis) of cystoisospora belli (formerly Isospora Belli): follows initial treatment in HIV-infected patients AND treatment failure or side effect with trimethoprim-sulfamethoxazole (TMP-SMX) or has a medical reason for not using TMP-SMX

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Toxoplasmosis: infectious disease specialist, ophthalmologist, or gynecologist. Pneumocystis jiroveci (formerly Pneumocystis carinii) Pneumonia (PCP) prevention and cystoisospora belli (formerly Isospora Belli) treatment or secondary prevention: infectious disease specialist.

**COVERAGE DURATION**

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## QUIZARTINIB (VANFLYTA)

---

### MEDICATION(S)

VANFLYTA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Induction: 2 months. Consolidation: 4 months. Maintenance therapy: 36 months. (see other criteria)

### OTHER CRITERIA

Treatment course consists of:

- a. Up to two cycles for use with standard cytarabine and anthracycline for induction, and
- b. Up to four cycles for use with cytarabine for consolidation, and
- c. Up to 36 cycles as a single agent for maintenance after consolidation therapy or until disease progression.

### PART B PREREQUISITE

N/A



## REGORAFENIB (STIVARGA)

---

### MEDICATION(S)

STIVARGA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## RELUGOLIX (ORGOVYX)

---

### **MEDICATION(S)**

ORGOVYX

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **RIBOCICLIB (KISQALI)**

---

### **MEDICATION(S)**

KISQALI (200 MG DOSE), KISQALI (400 MG DOSE), KISQALI (600 MG DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **RIBOCICLIB SUCCINATE/LETROZOLE (KISQALI FEMARA)**

---

### **MEDICATION(S)**

KISQALI FEMARA (400 MG DOSE), KISQALI FEMARA (600 MG DOSE), KISQALI FEMARA(200 MG DOSE)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **RIFAXIMIN (XIFAXAN)**

---

### **MEDICATION(S)**

XIFAXAN

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Clostridium difficile associated diarrhea (CDAD), Crohn's Disease, Small bowel bacterial overgrowth syndrome/Small intestinal bacterial overgrowth (SIBO)

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Travelers diarrhea: patient has tried azithromycin or a fluoroquinolone like ciprofloxacin or has a medical reason why ciprofloxacin and azithromycin cannot be used.

Hepatic Encephalopathy: patient has tried lactulose.

Irritable bowel syndrome with diarrhea (IBS-D): patient has tried an anti-diarrheal drug (diphenoxylate/atropine, loperamide) OR has a medical reason not to use (contraindication) anti-diarrheal therapies.

Clostridium difficile associated diarrhea (CDAD): patient has tried vancomycin.

Crohn's Disease: patient has tried metronidazole or ciprofloxacin or has a medical reason why metronidazole and ciprofloxacin cannot be used.

SIBO: Confirmation by a current positive breath test (e.g., lactulose hydrogen or glucose hydrogen) AND patient has tried and failed or had side effects with two of the following antibiotics: metronidazole (Flagyl), and ciprofloxacin (Cipro), amoxicillin-clavulanic acid (Augmentin), doxycycline, tetracycline, and trimethoprim-sulfamethoxazole (Bactrim, Septra) or there is a medical reason why all the other antibiotics cannot be tried first.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

One course (see other criteria)

**OTHER CRITERIA**

Travelers diarrhea: 3 days. Hepatic encephalopathy: plan year. IBS-D: 2 weeks. CDAD: 20 days. Crohn's Disease: 12 weeks. SIBO: 14 days.

**PART B PREREQUISITE**

N/A

## RILONACEPT (ARCALYST)

---

### MEDICATION(S)

ARCALYST

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Recurrent Pericarditis: trial of colchicine in combination with oral non-steroidal anti-inflammatory drug (NSAID) or contraindication to colchicine in combination with oral NSAID  
OR patient did not respond to corticosteroids or is on corticosteroids.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Recurrent Pericarditis: Cardiologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **RIOCIGUAT (ADEMPAS)**

---

### **MEDICATION(S)**

ADEMPAS

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Confirmation of Pulmonary Arterial Hypertension (WHO Group I) by right heart catheterization test AND patient has tried an endothelin-receptor antagonist (e.g. Tracleer) and a phosphodiesterase type 5 (PDE-5) inhibitor (e.g. sildenafil).

Confirmation of Chronic Thromboembolic Pulmonary Hypertension (CTEPH) by a right heart catheterization or V/Q scan AND patient has been treated with surgery or cannot be treated surgery.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **RIPRETINIB (QINLOCK)**

---

### **MEDICATION(S)**

QINLOCK

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **RISANKIZUMAB-RZAA (SKYRIZI)**

---

### **MEDICATION(S)**

SKYRIZI 150 MG/ML SOLN PRSYR, SKYRIZI 180 MG/1.2ML SOLN CART, SKYRIZI 360 MG/2.4ML SOLN CART, SKYRIZI (150 MG DOSE), SKYRIZI PEN

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another targeted immunotherapy drug.

### **REQUIRED MEDICAL INFORMATION**

Plaque Psoriasis (PsO), initial use: treatment failure or side effect with one DMARD or has a medical reason why methotrexate, cyclosporine, and acitretin cannot be used AND moderate to severe disease confirmed by Psoriasis Area and Severity Index (PASI) score of 10 or more OR Body Surface Area (BSA) of at least 3% OR sensitive areas are involved OR disease affects daily living. PsO, ongoing use: PASI or BSA improved with use of Skyrizi.  
Psoriatic arthritis (PsA): treatment failure or side effect with one DMARD drug or medical reason why methotrexate, leflunomide, and sulfasalazine cannot be used.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

PsO: Dermatologist or Rheumatologist. PsA: Rheumatologist

### **COVERAGE DURATION**

PsO initial use: 16 weeks. PsO ongoing use: plan year. PsA, Crohn's: plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **RISANKIZUMAB-RZAA IV (SKYRIZI IV)**

---

### **MEDICATION(S)**

SKYRIZI 600 MG/10ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another targeted immunotherapy drug.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

18 years of age or older.

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

one time induction course (8 weeks)

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## RISPERIDONE (PERSERIS ER)

---

### MEDICATION(S)

PERSERIS

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Older adults (65 years and older) with dementia-related psychosis

### REQUIRED MEDICAL INFORMATION

Treatment failure with at least one oral atypical antipsychotic (e.g. risperidone, ziprasidone) and medical reason why injectable risperidone (Risperdal Consta) cannot be used.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Psychiatrist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Excluded under Part D if covered by Part B.

### PART B PREREQUISITE

N/A

## ROFLUMILAST (DALIRESP)

---

### **MEDICATION(S)**

ROFLUMILAST

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## ROPEGINTERFERON ALFA-2B (BESREMI)

---

### MEDICATION(S)

BESREMI

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Polycythemia Vera (PV): treatment failure or side effect with hydroxyurea OR medical reason for not using hydroxyurea

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## RUCAPARIB CAMSYLATE (RUBRACA)

---

### **MEDICATION(S)**

RUBRACA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## RUXOLITINIB (JAKAFI)

---

### MEDICATION(S)

JAKAFI

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Myelofibrosis (MF): Being used along with another agent for myelofibrosis

### REQUIRED MEDICAL INFORMATION

MF: patient has enlarged spleen and platelet count is equal to or more than 50,000 cells/mcl.

Polycythemia Vera (PV): treatment failure or side effect with hydroxyurea OR medical reason for not using hydroxyurea and hematocrit of at least 40%.

Graft vs Host Disease (GvHD): treatment failure or side effect with systemic corticosteroids (e.g. prednisone, methylprednisolone).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## SACROSIDASE (SUCRAID)

---

### MEDICATION(S)

SUCRAID

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Initial use: patient has been evaluated to rule out other causes of disaccharidase deficiency.

Ongoing use: patient's gastrointestinal symptoms (i.e. abdominal pain, diarrhea, watery stools, bloating, flatulence) have been reduced.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Gastroenterologist, Geneticist, or Metabolic Specialist.

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## SAPROPTERIN DIHYDROCHLORIDE (KUVAN)

---

### MEDICATION(S)

JAVYGTOR, SAPROPTERIN DIHYDROCHLORIDE

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Phenylketonuria (PKU), initial: chart notes confirm PKU and baseline (just prior to therapy) blood phenylalanine (Phe) levels are given. PKU, ongoing use: phenylalanine level improved from baseline

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Initial: 3 months dose increases: 3 months, ongoing use: plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## SELEGILINE TRANSDERMAL (EMSAM)

---

### MEDICATION(S)

EMSAM

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Treatment failure or side effect with at least one preferred drug that treats depression (e.g. bupropion, maprotiline, citalopram, paroxetine, sertraline, venlafaxine, duloxetine).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **SELINEXOR (XPOVIO)**

---

### **MEDICATION(S)**

XPOVIO (100 MG ONCE WEEKLY), XPOVIO (40 MG ONCE WEEKLY), XPOVIO (40 MG TWICE WEEKLY), XPOVIO (60 MG ONCE WEEKLY), XPOVIO (60 MG TWICE WEEKLY), XPOVIO (80 MG ONCE WEEKLY), XPOVIO (80 MG TWICE WEEKLY)

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **SELPERCATINIB (RETEVMO)**

---

### **MEDICATION(S)**

RETEVMO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **SELUMETINIB (KOSELUGO)**

---

### **MEDICATION(S)**

KOSELUGO

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## SEMAGLUTIDE (RYBELSUS)

---

### MEDICATION(S)

RYBELSUS

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another GLP-1 agent

### REQUIRED MEDICAL INFORMATION

Type 2 diabetes: Confirmed by one of the following tests: Hemoglobin A1C of 6.5% or higher, or fasting blood sugar (glucose) of at least 126 mg/dl , or blood glucose of greater than or equal to 200 mg/dl either by Oral Glucose Tolerance Test or Plasma (blood) test AND one of the following: trial and failure or side effect with metformin or medical reason why metformin cannot be used, or A1C of 7.5% or greater and being used with another diabetes drug.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Dose and duration is not more than the FDA labeled maximum.

### PART B PREREQUISITE

N/A

## SEMAGLUTIDE SQ (OZEMPIC)

---

### MEDICATION(S)

OZEMPIC (0.25 OR 0.5 MG/DOSE), OZEMPIC (1 MG/DOSE), OZEMPIC (2 MG/DOSE)

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another GLP-1 agent

### REQUIRED MEDICAL INFORMATION

Type 2 diabetes: Confirmed by one of the following tests: Hemoglobin A1C of 6.5% or higher, or fasting blood sugar (glucose) of at least 126 mg/dl , or blood glucose of greater than or equal to 200 mg/dl either by Oral Glucose Tolerance Test or Plasma (blood) test AND one of the following: trial and failure or side effect with metformin or medical reason why metformin cannot be used, or A1C of 7.5% or greater and being used with another diabetes drug, or patient has heart (cardiovascular) disease.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

Dose and duration is not more than the FDA labeled maximum.

### PART B PREREQUISITE

N/A



## SILDENAFIL (REVATIO)

---

### MEDICATION(S)

SILDENAFIL CITRATE 10 MG/ML RECON SUSP, SILDENAFIL CITRATE 20 MG TAB

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Raynauds phenomenon

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

PAH: confirmation of WHO Group I by right heart catheterization test.

Raynaud's phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## SILDENAFIL (VIAGRA)

---

### MEDICATION(S)

SILDENAFIL CITRATE 100 MG TAB, SILDENAFIL CITRATE 25 MG TAB, SILDENAFIL CITRATE 50 MG TAB

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Penile rehabilitation, Raynaud's phenomenon

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

ED: due to a drug or medical condition that has been reported in the medical literature to cause ED.

Penile rehabilitation: prostate removed by surgery (radical prostatectomy) within the past 6 months.

Raynaud's phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Penile rehab: 6 months

ED and Raynaud's: plan year

### OTHER CRITERIA

ED: limited to 6 tablets per month

### PART B PREREQUISITE

N/A

## **SODIUM OXYBATE (XYREM)**

---

### **MEDICATION(S)**

XYREM

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Xyrem is being used with sedative hypnotic drugs or other CNS depressant drugs.

### **REQUIRED MEDICAL INFORMATION**

Narcolepsy is confirmed by sleep study and patient has brief losses of muscle tone (cataplexy). Excessive daytime sleepiness due to narcolepsy: patient has tried methylphenidate, amphetamine, or dextroamphetamine, and did not respond or had a side effect to modafinil or armodafinil, OR has a medical reason not to use methylphenidates, amphetamines, dextroamphetamines, modafinil and armodafinil.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **SODIUM PHENYLBUTYRATE (BUPHENYL)**

---

### **MEDICATION(S)**

SODIUM PHENYLBUTYRATE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Chart documentation for inherited Urea Cycle enzyme deficiency.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **SOFOSBUVIR/VELPATASVIR (EPCLUSA)**

---

### **MEDICATION(S)**

EPCLUSA, SOFOSBUVIR-VELPATASVIR

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current AASLD/IDSA guidelines.

### **REQUIRED MEDICAL INFORMATION**

Required medical information will be aligned with current AASLD/IDSA guidelines.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Hepatologist, Gastroenterologist, or Infectious Disease.

### **COVERAGE DURATION**

Length of therapy will be based on current AASLD/IDSA guidelines and FDA labeling.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR (VOSEVI)**

---

### **MEDICATION(S)**

VOSEVI

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current AASLD/IDSA guidelines.

### **REQUIRED MEDICAL INFORMATION**

Required medical information will be aligned with current AASLD/IDSA guidelines.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Hepatologist, Gastroenterologist, or Infectious Disease.

### **COVERAGE DURATION**

Length of therapy will be based on current AASLD/IDSA guidelines and FDA labeling.

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## SOMATROPIN (NORDITROPIN)

---

### MEDICATION(S)

NORDITROPIN FLEXPRO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Growth hormone deficiency (GHD) with pituitary disease: ADULTS - evidence of pituitary disease and failed one standard growth hormone stimulation test within one year of starting growth hormone. PEDIATRICS - evidence of pituitary disease, growth velocity decline, and patient failed one standard growth hormone stimulation test. GHD without pituitary disease: ADULTS - there is at least one documented pituitary hormone defect, IGF-I is below mean of reference range (below 50th percentile), and patient has failed one growth hormone stimulation test OR there are three or more documented pituitary hormone defects and IGF-1 is outside of reference range for sex/age. PEDIATRICS - height is less than 3rd percentile for age/sex, height velocity is less than 10th percentile of normal for age/sex tracked over at least one year, and patient either failed two standard growth hormone stimulation tests or failed one standard growth hormone stimulation test and has low IGF-1. GHD continuing from childhood, initial: evidence of pituitary disease OR patient failed one standard growth hormone stimulation test after the age of 18 and within one year of starting growth hormone therapy. GHD continuing from childhood, ongoing use: prescriber states patient responded to therapy. Small for Gestational Age (SGA): patients length at birth or birth weight are two or more standard deviations below the mean (less than the 3rd percentile) for gestational age (adjusted for prematurity) and patients height is two or more standard deviations below the mean. Ongoing use in SGA or pediatric GHD: growth velocity improved while on GH. Ongoing use for pediatrics with growth failure due to chronic kidney disease: patient did not have a kidney transplant within the past year. Ongoing use for Turners or Prader-Willi syndrome: prescriber has determined that benefits outweigh risk and continuation is necessary.

### AGE RESTRICTION

N/A

**PRESCRIBER RESTRICTION**

Adult GHD: Endocrinologist. Turners Syndrome, Prader-Willi Syndrome, pediatric GHD, SGA: Endocrinologist.

**COVERAGE DURATION**

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A



## SONIDEGIB (ODOMZO)

---

### MEDICATION(S)

ODOMZO

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Initial: 6 months

Ongoing use: plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **SORAFENIB (NEXAVAR)**

---

### **MEDICATION(S)**

SORAFENIB TOSYLATE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## SOTORASIB (LUMAKRAS)

---

### MEDICATION(S)

LUMAKRAS

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## STIRIPENTOL (DIACOMIT)

---

### **MEDICATION(S)**

DIACOMIT

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Seizures due to Dravet syndrome: being used with clobazam.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## SUNITINIB MALATE (SUTENT)

---

### MEDICATION(S)

SUNITINIB MALATE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TACROLIMUS (ENVARUSUS XR)

---

### **MEDICATION(S)**

ENVARUSUS XR

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has tried immediate-release (IR) tacrolimus or has a medical reason for not using IR tacrolimus.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under part D if covered by part B.

### **PART B PREREQUISITE**

N/A

## **TACROLIMUS FOR ORAL SUSPENSION (PROGRAF GRANULES)**

---

### **MEDICATION(S)**

PROGRAF 0.2 MG PACKET, PROGRAF 1 MG PACKET

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Patient has a medical reason for not using tacrolimus capsules.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## TADALAFIL (ADCIRCA)

---

### MEDICATION(S)

ALYQ, TADALAFIL (PAH)

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Raynaud's phenomenon

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

PAH: confirmation of WHO Group I by right heart catheterization test. Raynaud's phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## **TADALAFIL (CIALIS ED)**

---

### **MEDICATION(S)**

TADALAFIL 10 MG TAB, TADALAFIL 20 MG TAB

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Raynaud's phenomenon, Penile rehabilitation

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Raynauds phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine) and sildenafil (Viagra). Benign Prostatic Hyperplasia (BPH): treatment failure or side effect with both finasteride and tamsulosin.

ED: due to a drug or medical condition that has been reported in the medical literature to cause ED.

Penile rehabilitation: prostate removed by surgery (radical prostatectomy) within the past 6 months.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Penile rehab: 6 months

ED and Raynaud's: plan year

### **OTHER CRITERIA**

ED: limited to 6 tablets per month

### **PART B PREREQUISITE**

N/A

## TADALAFIL (CIALIS)

---

### MEDICATION(S)

TADALAFIL 2.5 MG TAB, TADALAFIL 5 MG TAB

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Raynauds phenomenon

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Raynauds phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine) and sildenafil (Viagra). Benign Prostatic Hyperplasia (BPH): treatment failure or side effect with both finasteride and tamsulosin.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TAFAMIDIS (VYNDAMAX)

---

### MEDICATION(S)

VYNDAMAX

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with a gene silencer like Tegsedi or Onpattro.

### REQUIRED MEDICAL INFORMATION

Heart disease (cardiomyopathy) is due to transthyretin-mediated amyloidosis (ATTR) confirmed by clinical features, genetic testing, and biopsy or immunochemistry.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Cardiologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TAFAMIDIS MEGLUMINE (VYNDAQEL)

---

### MEDICATION(S)

VYNDAQEL

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with a gene silencer like Tegsedi or Onpattro.

### REQUIRED MEDICAL INFORMATION

Heart disease (cardiomyopathy) is due to transthyretin-mediated amyloidosis (ATTR) confirmed by clinical features, genetic testing, and biopsy or immunochemistry.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Cardiologist

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TALAZOPARIB TOSYLATE (TALZENNA)

---

### **MEDICATION(S)**

TALZENNA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TASIMELTEON (HETLIOZ)

---

### MEDICATION(S)

TASIMELTEON

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Non 24 Sleep Wake Cycle, initial use: patient not able to maintain a stable 24-hour sleep-wake pattern synchronized to 24-hr light/dark cycle. Non 24 Sleep Wake Cycle, ongoing use: patients total sleep time at night is longer and has less day time sleep since starting tasimelteon.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Non 24 Sleep Wake Cycle: sleep specialist, neurologist

### COVERAGE DURATION

Non 24 Sleep Wake Cycle, initial use: 3mos. Ongoing use: plan year Smith-Magenis Syndrome: plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TAZEMETOSTAT (TAZVERIK)

---

### MEDICATION(S)

TAZVERIK

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **TEDUGLUTIDE (GATTEX)**

---

### **MEDICATION(S)**

GATTEX

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Initial use: patient has been dependent on parenteral nutrition for at least 3 months

Ongoing use: patient is still receiving parenteral nutrition (e.g. TPN or PPN) and has had a reduction in weekly parenteral nutrition volume since starting Gattex.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

6 months

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## TELOTRISTAT ETHYL (XERMELO)

---

### MEDICATION(S)

XERMELO

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Treatment failure of octreotide (Sandostatin) or lanreotide (Somatuline) AND being used in combination with octreotide (Sandostatin) or lanreotide (Somatuline).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TEPOTINIB (TEPMETKO)

---

### MEDICATION(S)

TEPMETKO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TERIFLUNOMIDE (AUBAGIO)

---

### **MEDICATION(S)**

TERIFLUNOMIDE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other disease-modifying therapies for relapsing Multiple Sclerosis.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TERIPARATIDE (FORTEO)

---

### MEDICATION(S)

FORTEO

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with other osteoporosis drugs.

### REQUIRED MEDICAL INFORMATION

Osteoporosis: one of the following: patient has a history of a broken bone not due to trauma (non-traumatic fracture) or T-score between -1.0 and -2.5 and is at high risk for fracture or T-score lower than -2.5 AND trial of a bisphosphonate (e.g. alendronate, ibandronate, or zoledronic acid) or Prolia, OR side effect to bisphosphonate therapy or Prolia therapy that supports discontinuation, OR Patient is at very high risk of fracture by meeting at least one of the following: non-traumatic fracture while on bisphosphonate therapy or Prolia, patient has experienced a recent fracture (within the past 12 months) or history of multiple fractures, or T-score less than -3.0, or patient is at high risk for falls, or 10-year hip fracture probability greater than 4.5% based on FRAX score, or 10-year major osteoporosis-related fracture probability greater than 30% based on FRAX score AND medical reason why Tymlos cannot be used. Glucocorticoid-induced osteoporosis: initiating or continuing long-term glucocorticoid treatment (e.g. prednisone, dexamethasone) and either has history of a non-traumatic fracture or is at high risk for fracture.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## TESAMORELIN ACETATE (EGRIFTA)

---

### MEDICATION(S)

EGRIFTA SV, EGRIFTA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Initial use: for men: having a waist circumference greater than 37.4 inches (97 cm) and a waist to hip ratio greater than or equal to 0.94 or for women: having a waist circumference greater than 37 inches (94 cm) and a waist to hip ratio greater than or equal to 0.88. Ongoing use: patient has had or maintained improvement in waist circumference.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Endocrinologist or HIV Specialist

### COVERAGE DURATION

Initial use: 3 months Ongoing use: 6 months

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## **TETRABENAZINE (XENAZINE)**

---

### **MEDICATION(S)**

TETRABENAZINE

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TEZACAFTOR-IVACAFTOR (SYMDEKO)

---

### **MEDICATION(S)**

SYMDEKO

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another CFTR potentiator drug (e.g. Orkambi).

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## THALIDOMIDE (THALOMID)

---

### **MEDICATION(S)**

THALOMID

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **TIOPRONIN (THIOLA)**

---

### **MEDICATION(S)**

THIOLA EC, TIOPRONIN

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **TIVOZANIB (FOTIVDA)**

---

### **MEDICATION(S)**

FOTIVDA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TOBRAMYCIN (TOBI PODHALER)

---

### MEDICATION(S)

TOBI PODHALER

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

bronchiectasis

### EXCLUSION CRITERIA

Being used for acute treatment of an infection.

### REQUIRED MEDICAL INFORMATION

Patient has cystic fibrosis or bronchiectasis and copy of sputum culture positive for Pseudomonas Aeruginosa.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TOBRAMYCIN INHALATION AGENTS

---

### **MEDICATION(S)**

TOBRAMYCIN 300 MG/4ML NEBU SOLN, TOBRAMYCIN 300 MG/5ML NEBU SOLN

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

bronchiectasis

### **EXCLUSION CRITERIA**

Being used for acute treatment of an infection.

### **REQUIRED MEDICAL INFORMATION**

Patient has cystic fibrosis or a bronchiectasis and copy of sputum culture is positive for Pseudomonas Aeruginosa.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## TOFACITINIB (XELJANZ, XELJANZ XR)

---

### MEDICATION(S)

XELJANZ, XELJANZ XR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Rheumatoid Arthritis and Polyarticular Juvenile Idiopathic Arthritis (pJIA): treatment failure or side effect with methotrexate OR medical reason why methotrexate cannot be used AND trial of a TNF inhibitor (i.e., Humira). Psoriatic arthritis (PsA): treatment failure or side effect with one DMARD drug or medical reason why methotrexate, leflunomide, and sulfasalazine cannot be used AND trial of a TNF inhibitor (i.e., Humira). Spondyloarthritis (SpA): trial of a TNF inhibitor (i.e., Humira). For oral solution for all indications: patient is unable to swallow a tablet.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

RA, PsA, pJIA, SpA: Rheumatologist.

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TOPICAL TESTOSTERONE PRODUCTS

---

### MEDICATION(S)

TESTOSTERONE 1.62 % GEL, TESTOSTERONE 10 MG/ACT (2%) GEL, TESTOSTERONE 12.5 MG/ACT (1%) GEL, TESTOSTERONE 20.25 MG/1.25GM (1.62%) GEL, TESTOSTERONE 20.25 MG/ACT (1.62%) GEL, TESTOSTERONE 25 MG/2.5GM (1%) GEL, TESTOSTERONE 30 MG/ACT SOLUTION, TESTOSTERONE 40.5 MG/2.5GM (1.62%) GEL, TESTOSTERONE 50 MG/5GM (1%) GEL

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

transgender, gender dysphoria

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TOPIRAMATE EXTENDED RELEASE (QUDEXY XR)

---

### MEDICATION(S)

TOPIRAMATE ER 100 MG CP24 SPRNK, TOPIRAMATE ER 150 MG CP24 SPRNK, TOPIRAMATE ER 200 MG CP24 SPRNK, TOPIRAMATE ER 25 MG CP24 SPRNK, TOPIRAMATE ER 50 MG CP24 SPRNK

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Patient has tried immediate-release topiramate or has a medical reason why patient cannot use immediate-release topiramate.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## TOPIRAMATE ORAL SOLUTION (EPRONTIA)

---

### **MEDICATION(S)**

EPRONTIA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Medical reason why patient cannot use topiramate tablet or sprinkle capsules

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TRAMADOL ER (CONZIP, RYZOLT) – NARCOTIC SAFETY INITIATIVE

---

### **MEDICATION(S)**

TRAMADOL HCL (ER BIPHASIC), TRAMADOL HCL ER (BIPHASIC)

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other long-acting narcotic drugs.

### **REQUIRED MEDICAL INFORMATION**

Cancer pain: dose has been consolidated to the least number of higher strength forms.

Non-cancer pain, initial: cause of pain cannot be removed or treated with other treatment options, and pain occurs daily and has lasted for at least 3 months, and pain is severe enough to need daily around-the-clock long-term narcotic use, and total daily dose across all narcotic drugs is less than 90 MME, and dose has been consolidated to the least number of higher strength forms and trial of tramadol immediate-release and intolerable side effect to the preferred tramadol extended release (generic Ultram ER), and chart notes document pain history including baseline pain intensity score and functional interference score, a plan for monitoring side effects and misuse, and a plan to taper down narcotics.

Non-cancer pain, reauth: total daily dose across all narcotic drugs is less than 90 MME per day, and dose has been consolidated to the least number of higher strength forms, and chart notes document current pain intensity score, functional interference score, any side effects and/or misuse with current pain treatment regimen, and plan to taper narcotic use.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Cancer Pain: Oncologist or Pain Specialist.

### **COVERAGE DURATION**

Cancer pain: plan year

Non-cancer pain: initial 30 days, 1st reauth 3mos, ongoing reauths plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## TRAMADOL ER (ULTRAM ER) – NARCOTIC SAFETY INITIATIVE

---

### **MEDICATION(S)**

TRAMADOL HCL ER 100 MG TAB ER 24H, TRAMADOL HCL ER 200 MG TAB ER 24H,  
TRAMADOL HCL ER 300 MG TAB ER 24H

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with other long-acting narcotic drugs.

### **REQUIRED MEDICAL INFORMATION**

Cancer pain: dose has been consolidated to the least number of higher strength forms.

Non-cancer pain, initial: cause of pain cannot be removed or treated with other treatment options, and pain occurs daily and has lasted for at least 3 months, and pain is severe enough to need daily around-the-clock long-term narcotic use, and total daily dose across all narcotic drugs is less than 90 MME, and dose has been consolidated to the least number of higher strength forms and trial of short-acting tramadol, and chart notes document pain history including baseline pain intensity score and functional interference score, a plan for monitoring side effects and misuse, and a plan to taper down narcotics.

Non-cancer pain, reauth: total daily dose across all narcotic drugs is less than 90 MME per day, and dose has been consolidated to the least number of higher strength forms, and chart notes document current pain intensity score, functional interference score, any side effects and/or misuse with current pain treatment regimen, and plan to taper narcotic use.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

Cancer pain: Oncologist or Pain Specialist.

### **COVERAGE DURATION**

Cancer pain: plan year

Non-cancer pain: initial 30 days, 1st reauth 3mos, ongoing reauths plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## TRAMETINIB (MEKINIST)

---

### MEDICATION(S)

MEKINIST

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TRANSMUCOSAL FENTANYL PRODUCTS

---

### MEDICATION(S)

FENTANYL CITRATE 100 MCG TAB, FENTANYL CITRATE 1200 MCG LOZ HANDLE, FENTANYL CITRATE 1600 MCG LOZ HANDLE, FENTANYL CITRATE 200 MCG LOZ HANDLE, FENTANYL CITRATE 200 MCG TAB, FENTANYL CITRATE 400 MCG LOZ HANDLE, FENTANYL CITRATE 400 MCG TAB, FENTANYL CITRATE 600 MCG LOZ HANDLE, FENTANYL CITRATE 600 MCG TAB, FENTANYL CITRATE 800 MCG LOZ HANDLE, FENTANYL CITRATE 800 MCG TAB

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Documentation of pain due to cancer.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Oncologist or Pain Management Specialist

### COVERAGE DURATION

6 months

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TRETINOIN PRODUCTS (AVITA, RETIN-A, ATRALIN)

---

### MEDICATION(S)

AVITA, TRETINOIN 0.01 % GEL, TRETINOIN 0.025 % CREAM, TRETINOIN 0.025 % GEL, TRETINOIN 0.05 % CREAM, TRETINOIN 0.05 % GEL, TRETINOIN 0.1 % CREAM

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

40 years of age or older. No prior authorization required for less than 40 years old.

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A



## TRIAMCINOLONE TOPICAL SPRAY (KENALOG)

---

### MEDICATION(S)

TRIAMCINOLONE ACETONIDE 0.147 MG/GM AERO SOLN

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Trial of triamcinolone cream, ointment, or lotion AND one other topical steroid in the same or higher potency (strength) class.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TRIENTINE HCL (SYPRINE)

---

### MEDICATION(S)

CLOVIQUE, TRIENTINE HCL

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## TRIFLURIDINE/TIPIRACIL HCL (LONSURF)

---

### **MEDICATION(S)**

LONSURF

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## TUCATINIB (TUKYSA)

---

### **MEDICATION(S)**

TUKYSA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## UBROGEPANT (UBRELVY)

---

### MEDICATION(S)

UBRELVY

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Trial of at least two triptans (e.g., sumatriptan, rizatriptan, naratriptan) or has a medical reason (contraindication) for not using triptans.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## UMBRALISIB TOSYLATE (UKONIQ)

---

### **MEDICATION(S)**

UKONIQ

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## UPADACITINIB (RINVOQ)

---

### MEDICATION(S)

RINVOQ

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Rheumatoid Arthritis: treatment failure or side effect with methotrexate or medical reason why methotrexate cannot be used AND trial of a TNF inhibitor (i.e., Humira). Psoriatic arthritis (PsA): treatment failure or side effect with one DMARD drug or medical reason why methotrexate, leflunomide, and sulfasalazine cannot be used AND trial of a TNF inhibitor (i.e., Humira). Atopic Dermatitis: moderate to severe disease confirmed by Investigator's Global Assessment (IGA) score of 3-4, Eczema Area and Severity Index (EASI) score of at least 16, Body surface area of at least 10%, or Severity Scoring of Atopic Dermatitis Index (SCORAD) score of at least 25 AND treatment failure or side effect with a topical calcineurin inhibitor (i.e., tacrolimus ointment) AND any of the following systemic therapies: methotrexate, corticosteroids, azathioprine, mycophenolate mofetil OR has a medical reason why these topical and systemic therapies cannot be used. Ulcerative Colitis (UC): Trial of a TNF inhibitor (i.e., Humira). Spondyloarthritis (SpA): Trial of a TNF inhibitor (i.e., Humira).

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

RA, PsA: Rheumatologist. Atopic Dermatitis: Dermatologist, Allergist, or Immunologist

### COVERAGE DURATION

Plan year

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A



## USTEKINUMAB IV (STELARA IV)

---

### **MEDICATION(S)**

STELARA 130 MG/26ML SOLUTION

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Being used with another targeted immunotherapy drug.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

18 years of age or older.

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

one time induction infusion

### **OTHER CRITERIA**

Excluded under Part D if covered by Part B.

### **PART B PREREQUISITE**

N/A

## USTEKINUMAB SQ (STELARA)

---

### MEDICATION(S)

STELARA 45 MG/0.5ML SOLN PRSYR, STELARA 45 MG/0.5ML SOLUTION, STELARA 90 MG/ML SOLN PRSYR

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Being used with another targeted immunotherapy drug.

### REQUIRED MEDICAL INFORMATION

Plaque Psoriasis (PsO), initial use: treatment failure or side effect with one DMARD or has a medical reason why methotrexate, cyclosporine, and acitretin cannot be used AND moderate to severe disease confirmed by Psoriasis Area and Severity Index (PASI) score of 10 or more OR Body Surface Area (BSA) of at least 3% OR sensitive areas are involved OR disease affects daily living. PsO, ongoing use: PASI or BSA improved with use of Stelara.

Psoriatic arthritis (PsA): treatment failure or side effect with one DMARD drug or medical reason why methotrexate, leflunomide, and sulfasalazine cannot be used.

Crohn's Disease (CD), initial use: SQ formulation will be started after initial IV dose. CD, ongoing use: symptom improvement with use of Stelara.

Ulcerative colitis (UC), initial use: disease is moderate to severe AND SQ formulation will be started after initial IV dose. UC, ongoing use: symptom improvement with use of Stelara.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

PsO: Dermatologist or Rheumatologist. PsA: Rheumatologist.

### COVERAGE DURATION

PsO initial use: 28 weeks. PsO ongoing use, CD, UC, and PsA: plan year.

**OTHER CRITERIA**

N/A

**PART B PREREQUISITE**

N/A

## VALBENAZINE (INGREZZA)

---

### MEDICATION(S)

INGREZZA

### PA INDICATION INDICATOR

1 - All FDA-Approved Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Tardive Dyskinesia: Neurologist or Psychiatrist.

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## VANCOMYCIN ORAL SOLUTION (FIRVANQ)

---

### **MEDICATION(S)**

VANCOMYCIN HCL 250 MG/5ML RECON SOLN, VANCOMYCIN HCL 50 MG/ML RECON SOLN

### **PA INDICATION INDICATOR**

1 - All FDA-Approved Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Clostridium difficile: evidence of current infection.

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

10 days

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **VANDETANIB (CAPRELSA)**

---

### **MEDICATION(S)**

CAPRELSA

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## **VARDENAFIL (LEVITRA, STAXYN)**

---

### **MEDICATION(S)**

VARDENAFIL HCL

### **PA INDICATION INDICATOR**

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### **OFF LABEL USES**

Benign Prostatic Hyperplasia, Raynaud's phenomenon

### **EXCLUSION CRITERIA**

N/A

### **REQUIRED MEDICAL INFORMATION**

Raynaud's phenomenon: treatment failure or side effect with a calcium-channel blocker (e.g. nifedipine) and treatment failure or side effect with sildenafil (Viagra).

BPH: trial of both finasteride and tamsulosin.

ED: due to a drug or medical condition that has been reported in the medical literature to cause ED and treatment failure or side effect with sildenafil (Viagra) and tadalafil (Cialis).

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

ED: limited to 6 tablets per month.

### **PART B PREREQUISITE**

N/A

## **VEMURAFENIB (ZELBORAF)**

---

### **MEDICATION(S)**

ZELBORAF

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A



## **VENETOCLAX (VENCLEXTA)**

---

### **MEDICATION(S)**

VENCLEXTA, VENCLEXTA STARTING PACK

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## VIGADRONE AND VIGABATRIN (SABRIL)

---

### MEDICATION(S)

VIGABATRIN, VIGADRONE

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

For continued use: ongoing diagnosis of infantile spasm is confirmed by EEG OR prescriber provides medical reason for continued use.

Complex partial seizures: patient has tried two preferred partial seizure drugs (e.g. carbamazepine, clonazepam, divalproex, felbamate, lamotrigine, levetiracetam, oxcarbazepine, phenytoin, topiramate, zonisamide)

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

Infantile spasms: Neurologist

### COVERAGE DURATION

Seizures: annual

Infantile spasms: 6 months

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## VISMODEGIB (ERIVEDGE)

---

### **MEDICATION(S)**

ERIVEDGE

### **PA INDICATION INDICATOR**

3 - All Medically-Accepted Indications

### **OFF LABEL USES**

N/A

### **EXCLUSION CRITERIA**

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### **REQUIRED MEDICAL INFORMATION**

N/A

### **AGE RESTRICTION**

N/A

### **PRESCRIBER RESTRICTION**

N/A

### **COVERAGE DURATION**

Plan year

### **OTHER CRITERIA**

N/A

### **PART B PREREQUISITE**

N/A

## VORICONAZOLE ORAL (VFEND)

---

### MEDICATION(S)

VORICONAZOLE 200 MG TAB, VORICONAZOLE 40 MG/ML RECON SUSP, VORICONAZOLE 50 MG TAB

### PA INDICATION INDICATOR

4 - All FDA-Approved Indications, Some Medically-Accepted Indications

### OFF LABEL USES

Prophylaxis of Disseminated Candidiasis, Candida Endophthalmitis, Oropharyngeal Candidiasis, Allergic bronchopulmonary aspergillosis, maintenance treatment of talaromycosis (*Talaromyces marneffe* -formerly *Penicillium marneffe*) in HIV-positive patients, treatment of *Lomentospora* (formerly *Scedosporium*) proliferans infection, treatment of pulmonary aspergillosis, chronic (cavitary or necrotizing), prophylaxis of Invasive Aspergillosis in high-risk patients.

### EXCLUSION CRITERIA

N/A

### REQUIRED MEDICAL INFORMATION

Systemic fungal infection treatment: culture test confirms Aspergillosis, candidemia, deep-tissue candida infection, blastomycosis, *scedosporium apiospermum*, *fusarium* species.  
Candida infection of the esophagus, throat, mouth (esophageal or oropharyngeal candidiasis) after trial of fluconazole or there is a medical reason not to use fluconazole.  
Prophylaxis of asperillosis or candidiasis after a bone marrow or lung transplant.

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

BMT:6mo Lung tx:3mo Esophageal candida:1mo Candidemia/deep-tissue:1mo Other ind in other criteria

**OTHER CRITERIA**

coverage duration: ABPA: 4 month systemic treatment: plan year

**PART B PREREQUISITE**

N/A

## VORINOSTAT (ZOLINZA)

---

### MEDICATION(S)

ZOLINZA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## ZANUBRUTINIB (BRUKINSA)

---

### MEDICATION(S)

BRUKINSA

### PA INDICATION INDICATOR

3 - All Medically-Accepted Indications

### OFF LABEL USES

N/A

### EXCLUSION CRITERIA

Exclusion criteria will be based on current National Comprehensive Cancer Network (NCCN) guidelines and FDA labeling.

### REQUIRED MEDICAL INFORMATION

N/A

### AGE RESTRICTION

N/A

### PRESCRIBER RESTRICTION

N/A

### COVERAGE DURATION

Plan year

### OTHER CRITERIA

N/A

### PART B PREREQUISITE

N/A

## Part B vs D drugs

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

### Medication(s)

MEDICATION NAME	ROUTE	DOSE FORM
Abelcet 5 MG/ML SUSPENSION	IV	SUSPENSION
Abraxane 100 MG RECON SUSP	IV	RECON SUSP
Acetadote 200 MG/ML SOLUTION	IV	SOLUTION
Acetaminophen 10 MG/ML SOLUTION	IV	SOLUTION
Acetylcysteine 10 % SOLUTION	IN	SOLUTION
Acetylcysteine 20 % SOLUTION	IN	SOLUTION
Acetylcysteine 200 MG/ML SOLUTION	IV	SOLUTION
Acyclovir Sodium 50 MG/ML SOLUTION	IV	SOLUTION
Adcetris 50 MG RECON SOLN	IV	RECON SOLN
Adrenalin 1 MG/ML SOLUTION	IJ	SOLUTION
Adrenalin 30 MG/30ML SOLUTION	IJ	SOLUTION
Adriamycin 10 MG RECON SOLN	IV	RECON SOLN
Adriamycin 2 MG/ML SOLUTION	IV	SOLUTION
Adriamycin 50 MG RECON SOLN	IV	RECON SOLN
Adrucil 2.5 GM/50ML SOLUTION	IV	SOLUTION
Adrucil 5 GM/100ML SOLUTION	IV	SOLUTION
Adrucil 500 MG/10ML SOLUTION	IV	SOLUTION
Adstiladrin 300000000000 VP/ML SUSPENSION	IS	SUSPENSION
Akynzeo 235-0.25 MG RECON SOLN	IV	RECON SOLN
Akynzeo 235-0.25 MG/20ML SOLUTION	IV	SOLUTION
Akynzeo 300-0.5 MG CAP	PO	CAP
Albuterol Sulfate (2.5 MG/3ML) 0.083% NEBU SOLN	IN	NEBU SOLN
Albuterol Sulfate (5 MG/ML) 0.5% NEBU SOLN	IN	NEBU SOLN
Albuterol Sulfate 0.63 MG/3ML NEBU SOLN	IN	NEBU SOLN
Albuterol Sulfate 1.25 MG/3ML NEBU SOLN	IN	NEBU SOLN
Albuterol Sulfate 2.5 MG/0.5ML NEBU SOLN	IN	NEBU SOLN
Aldurazyme 2.9 MG/5ML SOLUTION	IV	SOLUTION
Alferon N 5000000 UNIT/ML SOLUTION	IJ	SOLUTION
Alimta 100 MG RECON SOLN	IV	RECON SOLN
Alimta 500 MG RECON SOLN	IV	RECON SOLN
Aliqopa 60 MG RECON SOLN	IV	RECON SOLN
Alkeran 2 MG TAB	PO	TAB
Alkeran 50 MG RECON SOLN	IV	RECON SOLN
Allopurinol Sodium 500 MG RECON SOLN	IV	RECON SOLN
Aloprim 500 MG RECON SOLN	IV	RECON SOLN
Aloxi 0.25 MG/5ML SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
Allymsys 100 MG/4ML SOLUTION	IV	SOLUTION
Allymsys 400 MG/16ML SOLUTION	IV	SOLUTION
AmBisome 50 MG RECON SUSP	IV	RECON SUSP
Amino Acid 5 % SOLUTION	IV	SOLUTION
Aminophylline 25 MG/ML SOLUTION	IV	SOLUTION
AminoProtect 5 % SOLUTION	IV	SOLUTION
Aminosyn II 10 % SOLUTION	IV	SOLUTION
Aminosyn II 15 % SOLUTION	IV	SOLUTION
Aminosyn-PF 10 % SOLUTION	IV	SOLUTION
Aminosyn-PF 7 % SOLUTION	IV	SOLUTION
Aminosyn-PF 7% 7 % SOLUTION	IV	SOLUTION
Amiodarone HCl 150 MG/3ML SOLUTION	IV	SOLUTION
Amiodarone HCl 450 MG/9ML SOLUTION	IV	SOLUTION
Amiodarone HCl 900 MG/18ML SOLUTION	IV	SOLUTION
Amphotericin B 50 MG RECON SOLN	IV	RECON SOLN
Amphotericin B Liposome 50 MG RECON SUSP	IV	RECON SUSP
Amvuttra 25 MG/0.5ML SOLN PRSYR	SC	SOLN PRSYR
Anzemet 100 MG TAB	PO	TAB
Anzemet 50 MG TAB	PO	TAB
Aprepitant 125 MG CAP	PO	CAP
Aprepitant 80 & 125 MG CAP	PO	CAP
Aprepitant 80 & 125 MG MISC	PO	MISC
Aprepitant 80 MG CAP	PO	CAP
Apretude 600 MG/3ML SUSP	IM	SUSP
Aralast NP 1000 MG RECON SOLN	IV	RECON SOLN
Aralast NP 500 MG RECON SOLN	IV	RECON SOLN
Arformoterol Tartrate 15 MCG/2ML NEBU SOLN	IN	NEBU SOLN
Argatroban 250 MG/2.5ML SOLUTION	IV	SOLUTION
Argatroban 50 MG/50ML SOLUTION	IV	SOLUTION
Arranon 5 MG/ML SOLUTION	IV	SOLUTION
Arsenic Trioxide 10 MG/10ML SOLUTION	IV	SOLUTION
Arsenic Trioxide 12 MG/6ML SOLUTION	IV	SOLUTION
Arzerra 100 MG/5ML CONC	IV	CONC
Arzerra 1000 MG/50ML CONC	IV	CONC
Asceniv 5 GM/50ML SOLUTION	IV	SOLUTION
Asparlas 3750 UNIT/5ML SOLUTION	IV	SOLUTION
Astagraf XL 0.5 MG CAP ER 24H	PO	CAP ER 24H
Astagraf XL 1 MG CAP ER 24H	PO	CAP ER 24H
Astagraf XL 5 MG CAP ER 24H	PO	CAP ER 24H
Atgam 50 MG/ML INJECTABLE	IV	INJECTABLE
Ativan 2 MG/ML SOLUTION	IJ	SOLUTION
Ativan 4 MG/ML SOLUTION	IJ	SOLUTION
Avastin 100 MG/4ML SOLUTION	IV	SOLUTION
Avastin 400 MG/16ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Aveed 750 MG/3ML SOLUTION	IM	SOLUTION
Avelox 400 MG/250ML SOLUTION	IV	SOLUTION
Avsola 100 MG RECON SOLN	IV	RECON SOLN
Avycaz 2.5 (2-0.5) GM RECON SOLN	IV	RECON SOLN
AzaCITIDine 100 MG RECON SUSP	IJ	RECON SUSP
Azasan 100 MG TAB	PO	TAB
Azasan 75 MG TAB	PO	TAB
azaTHIOprine 100 MG TAB	PO	TAB
AzaTHIOprine 50 MG TAB	PO	TAB
azaTHIOprine 75 MG TAB	PO	TAB
AzaTHIOprine Sodium 100 MG RECON SOLN	IJ	RECON SOLN
Baclofen 10 MG/20ML SOLUTION	IT	SOLUTION
Baclofen 20000 MCG/20ML SOLUTION	IT	SOLUTION
Baclofen 40 MG/20ML SOLUTION	IT	SOLUTION
Baclofen 50 MCG/ML SOLN PRSYR	IT	SOLN PRSYR
Bavencio 200 MG/10ML SOLUTION	IV	SOLUTION
Beleodaq 500 MG RECON SOLN	IV	RECON SOLN
Belrapzo 100 MG/4ML SOLUTION	IV	SOLUTION
Bendamustine HCl 100 MG RECON SOLN	IV	RECON SOLN
Bendamustine HCl 100 MG/4ML SOLUTION	IV	SOLUTION
Bendamustine HCl 25 MG RECON SOLN	IV	RECON SOLN
Bendeka 100 MG/4ML SOLUTION	IV	SOLUTION
Benlysta 120 MG RECON SOLN	IV	RECON SOLN
Benlysta 400 MG RECON SOLN	IV	RECON SOLN
Bentyl 10 MG/ML SOLUTION	IM	SOLUTION
Beovu 6 MG/0.05ML SOLN PRSYR	IZ	SOLN PRSYR
Beovu 6 MG/0.05ML SOLUTION	IZ	SOLUTION
Besponsa 0.9 MG RECON SOLN	IV	RECON SOLN
Betamethasone Combo 6 (3-3) MG/ML SUSPENSION	IJ	SUSPENSION
Betamethasone Sod Phos & Acet 6 (3-3) MG/ML SUSPENSION	IJ	SUSPENSION
Bethkis 300 MG/4ML NEBU SOLN	IN	NEBU SOLN
BiCNU 100 MG RECON SOLN	IV	RECON SOLN
Blenrep 100 MG RECON SOLN	IV	RECON SOLN
Bleomycin Sulfate 15 UNIT RECON SOLN	IJ	RECON SOLN
Bleomycin Sulfate 30 UNIT RECON SOLN	IJ	RECON SOLN
Blinicyto 35 MCG RECON SOLN	IV	RECON SOLN
Boniva 3 MG/3ML SOLUTION	IV	SOLUTION
Bortezomib 1 MG RECON SOLN	IJ	RECON SOLN
Bortezomib 2.5 MG RECON SOLN	IJ	RECON SOLN
Bortezomib 3.5 MG RECON SOLN	IJ	RECON SOLN
Bortezomib 3.5 MG RECON SOLN	IV	RECON SOLN
Bortezomib 3.5 MG/1.4ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Botox 100 UNIT RECON SOLN	IJ	RECON SOLN
Botox 200 UNIT RECON SOLN	IJ	RECON SOLN
Briumvi 150 MG/6ML SOLUTION	IV	SOLUTION
Brovana 15 MCG/2ML NEBU SOLN	IN	NEBU SOLN
Budesonide 0.25 MG/2ML SUSPENSION	IN	SUSPENSION
Budesonide 0.5 MG/2ML SUSPENSION	IN	SUSPENSION
Budesonide 1 MG/2ML SUSPENSION	IN	SUSPENSION
Buprenex 0.3 MG/ML SOLUTION	IJ	SOLUTION
Buprenorphine HCl 0.3 MG/ML SOLUTION	IJ	SOLUTION
Busulfan 6 MG/ML SOLUTION	IV	SOLUTION
Busulfex 6 MG/ML SOLUTION	IV	SOLUTION
Butorphanol Tartrate 1 MG/ML SOLUTION	IJ	SOLUTION
Butorphanol Tartrate 2 MG/ML SOLUTION	IJ	SOLUTION
Byooviz 0.5 MG/0.05ML SOLUTION	IZ	SOLUTION
Cabenuva 400 & 600 MG/2ML SUSP	IM	SUSP
Cabenuva 600 & 900 MG/3ML SUSP	IM	SUSP
Calcitonin (Salmon) 200 UNIT/ML SOLUTION	IJ	SOLUTION
Calcitriol 0.25 MCG CAP	PO	CAP
Calcitriol 0.5 MCG CAP	PO	CAP
Calcitriol 1 MCG/ML SOLUTION	IV	SOLUTION
Calcitriol 1 MCG/ML SOLUTION	PO	SOLUTION
Calcium Gluconate 10 % SOLUTION	IV	SOLUTION
Caldolor 800 MG/200ML SOLUTION	IV	SOLUTION
Caldolor 800 MG/8ML SOLUTION	IV	SOLUTION
Camcevi 42 MG PRSYR	SC	PRSYR
Campath 30 MG/ML SOLUTION	IV	SOLUTION
Camptosar 100 MG/5ML SOLUTION	IV	SOLUTION
Camptosar 300 MG/15ML SOLUTION	IV	SOLUTION
Camptosar 40 MG/2ML SOLUTION	IV	SOLUTION
CARBOplatin 150 MG/15ML SOLUTION	IV	SOLUTION
CARBOplatin 450 MG/45ML SOLUTION	IV	SOLUTION
CARBOplatin 50 MG/5ML SOLUTION	IV	SOLUTION
CARBOplatin 600 MG/60ML SOLUTION	IV	SOLUTION
Cardene IV 20-0.86 MG/200ML-% SOLUTION	IV	SOLUTION
Cardene IV 20-4.8 MG/200ML-% SOLUTION	IV	SOLUTION
Cardene IV 40-0.83 MG/200ML-% SOLUTION	IV	SOLUTION
Carmustine 100 MG RECON SOLN	IV	RECON SOLN
Carmustine 300 MG RECON SOLN	IV	RECON SOLN
Carmustine 50 MG RECON SOLN	IV	RECON SOLN
Carnitor 200 MG/ML SOLUTION	IV	SOLUTION
ceFAZolin in Sodium Chloride 2-0.9 GM/100ML-% SOLUTION	IV	SOLUTION
CeFAZolin in Sodium Chloride 3-0.9 GM/100ML-% SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
CeFAZolin Sodium-Dextrose 1-4 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CeFAZolin Sodium-Dextrose 1-4 GM/50ML-% SOLUTION	IV	SOLUTION
CeFAZolin Sodium-Dextrose 2-3 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CeFAZolin Sodium-Dextrose 2-4 GM/100ML-% SOLUTION	IV	SOLUTION
ceFAZolin Sodium-Dextrose 2-5 GM/100ML-% SOLUTION	IV	SOLUTION
Cefepime HCl 100 GM RECON SOLN	IV	RECON SOLN
Cefepime-Dextrose 1-5 GM-%(50ML) RECON SOLN	IV	RECON SOLN
Cefepime-Dextrose 2-5 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefoTetan Disodium-Dextrose 1-3.58 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefoTetan Disodium-Dextrose 2-2.08 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefOXitin Sodium-Dextrose 1-4 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefOXitin Sodium-Dextrose 2-2.2 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefTAZidime and Dextrose 1-5 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefTAZidime and Dextrose 2-5 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefTRIAXone Sodium 100 GM RECON SOLN	IJ	RECON SOLN
CefTRIAXone Sodium in Dextrose 20 MG/ML SOLUTION	IV	SOLUTION
CefTRIAXone Sodium in Dextrose 40 MG/ML SOLUTION	IV	SOLUTION
CefTRIAXone Sodium-Dextrose 1-3.74 GM-%(50ML) RECON SOLN	IV	RECON SOLN
CefTRIAXone Sodium-Dextrose 2-2.22 GM-%(50ML) RECON SOLN	IV	RECON SOLN
Celestone Soluspan 6 (3-3) MG/ML SUSPENSION	IJ	SUSPENSION
CellCept 200 MG/ML RECON SUSP	PO	RECON SUSP
CellCept 250 MG CAP	PO	CAP
CellCept 500 MG TAB	PO	TAB
CellCept Intravenous 500 MG RECON SOLN	IV	RECON SOLN
Cerebyx 100 MG PE/2ML SOLUTION	IJ	SOLUTION
Cerebyx 500 MG PE/10ML SOLUTION	IJ	SOLUTION
Chlorothiazide Sodium 500 MG RECON SOLN	IV	RECON SOLN
Chorionic Gonadotropin 10000 UNIT RECON SOLN	IM	RECON SOLN
Cidofovir 75 MG/ML SOLUTION	IV	SOLUTION
Cimerli 0.3 MG/0.05ML SOLUTION	IZ	SOLUTION
Cimerli 0.5 MG/0.05ML SOLUTION	IZ	SOLUTION
Cinacalcet HCl 30 MG TAB	PO	TAB
Cinacalcet HCl 60 MG TAB	PO	TAB

MEDICATION NAME	ROUTE	DOSE FORM
Cinacalcet HCl 90 MG TAB	PO	TAB
Cinqair 100 MG/10ML SOLUTION	IV	SOLUTION
Cinvanti 130 MG/18ML EMULSION	IV	EMULSION
Cipro in D5W 400 MG/200ML SOLUTION	IV	SOLUTION
Ciprofloxacin in D5W 400 MG/200ML SOLUTION	IV	SOLUTION
CISplatin 100 MG/100ML SOLUTION	IV	SOLUTION
CISplatin 200 MG/200ML SOLUTION	IV	SOLUTION
CISplatin 50 MG RECON SOLN	IV	RECON SOLN
CISplatin 50 MG/50ML SOLUTION	IV	SOLUTION
Cladribine 10 MG/10ML SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (2.75/5) 2.75 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (4.25/10) 4.25 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (4.25/5) 4.25 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (5/15) 5 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (5/20) 5 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (8/10) 8 % SOLUTION	IV	SOLUTION
Clinimix E/Dextrose (8/14) 8 % SOLUTION	IV	SOLUTION
Clinimix N14G30E 4.25 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (4.25/10) 4.25 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (4.25/25) 4.25 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (4.25/5) 4.25 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (5/15) 5 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (5/20) 5 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (5/25) 5 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (6/5) 6 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (8/10) 8 % SOLUTION	IV	SOLUTION
Clinimix/Dextrose (8/14) 8 % SOLUTION	IV	SOLUTION
Clinisol SF 15 % SOLUTION	IV	SOLUTION
Clinolipid 20 % EMULSION	IV	EMULSION
Clofarabine 1 MG/ML SOLUTION	IV	SOLUTION
Clolar 1 MG/ML SOLUTION	IV	SOLUTION
CloNIDine HCl (Analgesia) 100 MCG/ML SOLUTION	EP	SOLUTION
Cocaine HCl 40 MG/ML SOLUTION	NA	SOLUTION
Columvi 10 MG/10ML SOLUTION	IV	SOLUTION
Columvi 2.5 MG/2.5ML SOLUTION	IV	SOLUTION
Cosela 300 MG RECON SOLN	IV	RECON SOLN
Cosentyx 125 MG/5ML SOLUTION	IV	SOLUTION
Cosmegen 0.5 MG RECON SOLN	IV	RECON SOLN
Cromolyn Sodium 20 MG/2ML NEBU SOLN	IN	NEBU SOLN
Cupric Chloride 0.4 MG/ML SOLUTION	IV	SOLUTION
Cutaquig 1 GM/6ML SOLUTION	SC	SOLUTION
Cutaquig 1.65 GM/10ML SOLUTION	SC	SOLUTION
Cutaquig 2 GM/12ML SOLUTION	SC	SOLUTION
Cutaquig 3.3 GM/20ML SOLUTION	SC	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Cutaquig 4 GM/24ML SOLUTION	SC	SOLUTION
Cutaquig 8 GM/48ML SOLUTION	SC	SOLUTION
Cuvitru 1 GM/5ML SOLUTION	SC	SOLUTION
Cuvitru 10 GM/50ML SOLUTION	SC	SOLUTION
Cuvitru 2 GM/10ML SOLUTION	SC	SOLUTION
Cuvitru 4 GM/20ML SOLUTION	SC	SOLUTION
Cuvitru 8 GM/40ML SOLUTION	SC	SOLUTION
Cyclophosphamide 1 GM RECON SOLN	IJ	RECON SOLN
Cyclophosphamide 1 GM/5ML SOLUTION	IV	SOLUTION
Cyclophosphamide 2 GM RECON SOLN	IJ	RECON SOLN
Cyclophosphamide 2 GM/10ML SOLUTION	IV	SOLUTION
Cyclophosphamide 25 MG CAP	PO	CAP
Cyclophosphamide 25 MG TAB	PO	TAB
Cyclophosphamide 50 MG CAP	PO	CAP
Cyclophosphamide 50 MG TAB	PO	TAB
Cyclophosphamide 500 MG RECON SOLN	IJ	RECON SOLN
Cyclophosphamide 500 MG/2.5ML SOLUTION	IV	SOLUTION
Cyclophosphamide 500 MG/ML SOLUTION	IV	SOLUTION
CycloSPORINE 100 MG CAP	PO	CAP
CycloSPORINE 25 MG CAP	PO	CAP
CycloSPORINE 50 MG/ML SOLUTION	IV	SOLUTION
CycloSPORINE Modified 100 MG CAP	PO	CAP
CycloSPORINE Modified 100 MG/ML SOLUTION	PO	SOLUTION
CycloSPORINE Modified 25 MG CAP	PO	CAP
CycloSPORINE Modified 50 MG CAP	PO	CAP
Cyklokapron 1000 MG/10ML SOLUTION	IV	SOLUTION
Cyramza 100 MG/10ML SOLUTION	IV	SOLUTION
Cyramza 500 MG/50ML SOLUTION	IV	SOLUTION
Cytarabine (PF) 100 MG/ML SOLUTION	IJ	SOLUTION
Cytarabine (PF) 20 MG/ML SOLUTION	IJ	SOLUTION
Cytarabine 20 MG/ML SOLUTION	IJ	SOLUTION
Cytogam 50 MG/ML INJECTABLE	IV	INJECTABLE
Cytovene 500 MG RECON SOLN	IV	RECON SOLN
Dacarbazine 100 MG RECON SOLN	IV	RECON SOLN
Dacarbazine 200 MG RECON SOLN	IV	RECON SOLN
Dacogen 50 MG RECON SOLN	IV	RECON SOLN
DACTINomycin 0.5 MG RECON SOLN	IV	RECON SOLN
Danyelza 40 MG/10ML SOLUTION	IV	SOLUTION
DAPTOMycin-Sodium Chloride 1000-0.9 MG/100ML-% SOLUTION	IV	SOLUTION
DAPTOMycin-Sodium Chloride 350-0.9 MG/50ML-% SOLUTION	IV	SOLUTION
DAPTOMycin-Sodium Chloride 500-0.9 MG/50ML-% SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
DAPTOMycin-Sodium Chloride 700-0.9 MG/100ML-% SOLUTION	IV	SOLUTION
Darzalex 100 MG/5ML SOLUTION	IV	SOLUTION
Darzalex 400 MG/20ML SOLUTION	IV	SOLUTION
Darzalex Faspro 1800-30000 MG-UT/15ML SOLUTION	SC	SOLUTION
DAUNOrubicin HCl 20 MG/4ML SOLUTION	IV	SOLUTION
DAUNOrubicin HCl 50 MG/10ML SOLUTION	IV	SOLUTION
Decitabine 50 MG RECON SOLN	IV	RECON SOLN
Deferoxamine Mesylate 2 GM RECON SOLN	IJ	RECON SOLN
Deferoxamine Mesylate 500 MG RECON SOLN	IJ	RECON SOLN
Defitelio 200 MG/2.5ML SOLUTION	IV	SOLUTION
Demerol 100 MG/2ML SOLUTION	IJ	SOLUTION
Demerol 100 MG/ML SOLUTION	IJ	SOLUTION
Demerol 25 MG/0.5ML SOLUTION	IJ	SOLUTION
Demerol 25 MG/ML SOLUTION	IJ	SOLUTION
Demerol 50 MG/ML SOLUTION	IJ	SOLUTION
Demerol 75 MG/1.5ML SOLUTION	IJ	SOLUTION
Demerol 75 MG/ML SOLUTION	IJ	SOLUTION
DEPO-Medrol 20 MG/ML SUSPENSION	IJ	SUSPENSION
Desferal 500 MG RECON SOLN	IJ	RECON SOLN
Dexamethasone Sod Phosphate PF 10 MG/ML SOLN PRSYR	IJ	SOLN PRSYR
Dexamethasone Sod Phosphate PF 10 MG/ML SOLUTION	IJ	SOLUTION
Dexamethasone Sodium Phosphate 10 MG/ML SOLUTION	IJ	SOLUTION
Dexamethasone Sodium Phosphate 100 MG/10ML SOLUTION	IJ	SOLUTION
Dexrazoxane 250 MG RECON SOLN	IV	RECON SOLN
Dexrazoxane HCl 250 MG RECON SOLN	IV	RECON SOLN
Dexrazoxane HCl 500 MG RECON SOLN	IV	RECON SOLN
Dextrose 20 % SOLUTION	IV	SOLUTION
Dextrose 250 MG/ML SOLUTION	IV	SOLUTION
Dextrose 30 % SOLUTION	IV	SOLUTION
Dextrose 40 % SOLUTION	IV	SOLUTION
Dextrose 5%/Electrolyte #48 SOLUTION	IV	SOLUTION
Dextrose 50 % SOLUTION	IV	SOLUTION
Dextrose 70 % SOLUTION	IV	SOLUTION
Dexycu 9 % SUSPENSION	IO	SUSPENSION
diazePAM 10 MG/2ML SOLUTION	IJ	SOLUTION
DiazePAM 5 MG/ML SOLUTION	IJ	SOLUTION
Dicyclomine HCl 10 MG/ML SOLUTION	IM	SOLUTION
Digoxin 0.25 MG/ML SOLUTION	IJ	SOLUTION
Dilaudid 0.2 MG/ML SOLUTION	IJ	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Dilaudid 1 MG/ML SOLUTION	IJ	SOLUTION
Dilaudid 2 MG/ML SOLUTION	IJ	SOLUTION
DiITIAZem HCl 100 MG RECON SOLN	IV	RECON SOLN
DiITIAZem HCl 125 MG/25ML SOLUTION	IV	SOLUTION
DiITIAZem HCl 25 MG/5ML SOLUTION	IV	SOLUTION
DiITIAZem HCl 50 MG/10ML SOLUTION	IV	SOLUTION
DOBUTamine HCl 12.5 MG/ML SOLUTION	IV	SOLUTION
DOBUTamine HCl 250 MG/20ML SOLUTION	IV	SOLUTION
DOBUTamine in D5W 1-5 MG/ML-% SOLUTION	IV	SOLUTION
DOBUTamine in D5W 2 MG/ML SOLUTION	IV	SOLUTION
DOBUTamine in D5W 4-5 MG/ML-% SOLUTION	IV	SOLUTION
DOCEtaxel 160 MG/16ML SOLUTION	IV	SOLUTION
DOCEtaxel 160 MG/8ML CONC	IV	CONC
DOCEtaxel 20 MG/2ML SOLUTION	IV	SOLUTION
DOCEtaxel 20 MG/ML CONC	IV	CONC
DOCEtaxel 200 MG/10ML CONC	IV	CONC
DOCEtaxel 80 MG/4ML CONC	IV	CONC
DOCEtaxel 80 MG/8ML SOLUTION	IV	SOLUTION
DOPamine HCl 40 MG/ML SOLUTION	IV	SOLUTION
DOPamine in D5W 0.8-5 MG/ML-% SOLUTION	IV	SOLUTION
DOPamine in D5W 1.6-5 MG/ML-% SOLUTION	IV	SOLUTION
DOPamine in D5W 3.2-5 MG/ML-% SOLUTION	IV	SOLUTION
Doxercalciferol 0.5 MCG CAP	PO	CAP
Doxercalciferol 1 MCG CAP	PO	CAP
Doxercalciferol 2.5 MCG CAP	PO	CAP
Doxercalciferol 4 MCG/2ML SOLUTION	IV	SOLUTION
Doxil 2 MG/ML INJECTABLE	IV	INJECTABLE
DOXOrubicin HCl 10 MG RECON SOLN	IV	RECON SOLN
DOXOrubicin HCl 2 MG/ML SOLUTION	IV	SOLUTION
DOXOrubicin HCl 50 MG RECON SOLN	IV	RECON SOLN
DOXOrubicin HCl Liposomal 2 MG/ML INJECTABLE	IV	INJECTABLE
Duopa 4.63-20 MG/ML SUSPENSION	EN	SUSPENSION
Duraclon 100 MCG/ML SOLUTION	EP	SOLUTION
Duramorph 0.5 MG/ML SOLUTION	IJ	SOLUTION
Duramorph 1 MG/ML SOLUTION	IJ	SOLUTION
Durysta 10 MCG IMPLANT	IO	IMPLANT
Dysport 300 UNIT RECON SOLN	IM	RECON SOLN
Dysport 500 UNIT RECON SOLN	IM	RECON SOLN
Elahere 100 MG/20ML SOLUTION	IV	SOLUTION
Elaprase 6 MG/3ML SOLUTION	IV	SOLUTION
Elcys 50 MG/ML SOLUTION	IV	SOLUTION
ElELYso 200 UNIT RECON SOLN	IV	RECON SOLN
Elfabrio 20 MG/10ML SOLUTION	IV	SOLUTION
Elitek 1.5 MG RECON SOLN	IV	RECON SOLN



MEDICATION NAME	ROUTE	DOSE FORM
Elitek 7.5 MG RECON SOLN	IV	RECON SOLN
Ellence 200 MG/100ML SOLUTION	IV	SOLUTION
Ellence 50 MG/25ML SOLUTION	IV	SOLUTION
Elliotts B SOLUTION	IT	SOLUTION
Elrexio 44 MG/1.1ML SOLUTION	SC	SOLUTION
Elrexio 76 MG/1.9ML SOLUTION	SC	SOLUTION
Emend 125 MG CAP	PO	CAP
Emend 125 MG/5ML RECON SUSP	PO	RECON SUSP
Emend 150 MG RECON SOLN	IV	RECON SOLN
Emend 80 MG CAP	PO	CAP
Emend Tri-Pack 80 & 125 MG CAP	PO	CAP
Empaveli 1080 MG/20ML SOLUTION	SC	SOLUTION
Empliciti 300 MG RECON SOLN	IV	RECON SOLN
Empliciti 400 MG RECON SOLN	IV	RECON SOLN
Enalaprilat 1.25 MG/ML INJECTABLE	IV	INJECTABLE
Engerix-B 10 MCG/0.5ML SUSP PRSYR	IJ	SUSP PRSYR
Engerix-B 20 MCG/ML SUSP PRSYR	IJ	SUSP PRSYR
Engerix-B 20 MCG/ML SUSPENSION	IJ	SUSPENSION
Enhertu 100 MG RECON SOLN	IV	RECON SOLN
Enjaymo 1100 MG/22ML SOLUTION	IV	SOLUTION
Entyvio 300 MG RECON SOLN	IV	RECON SOLN
EPINEPHrine (Anaphylaxis) 1 MG/ML SOLUTION	IJ	SOLUTION
EPINEPHrine (Anaphylaxis) 30 MG/30ML SOLUTION	IJ	SOLUTION
EpiRUBicin HCl 200 MG/100ML SOLUTION	IV	SOLUTION
EpiRUBicin HCl 50 MG/25ML SOLUTION	IV	SOLUTION
Epkinly 4 MG/0.8ML SOLUTION	SC	SOLUTION
Epkinly 48 MG/0.8ML SOLUTION	SC	SOLUTION
Epogen 10000 UNIT/ML SOLUTION	IJ	SOLUTION
Epogen 2000 UNIT/ML SOLUTION	IJ	SOLUTION
Epogen 20000 UNIT/ML SOLUTION	IJ	SOLUTION
Epogen 3000 UNIT/ML SOLUTION	IJ	SOLUTION
Epogen 4000 UNIT/ML SOLUTION	IJ	SOLUTION
Epoprostenol Sodium 0.5 MG RECON SOLN	IV	RECON SOLN
Epoprostenol Sodium 1.5 MG RECON SOLN	IV	RECON SOLN
Erbix 100 MG/50ML SOLUTION	IV	SOLUTION
Erbix 200 MG/100ML SOLUTION	IV	SOLUTION
Erwinase 10000 UNIT RECON SOLN	IJ	RECON SOLN
Erwinaze 10000 UNIT RECON SOLN	IJ	RECON SOLN
Ethacrynate Sodium 50 MG RECON SOLN	IV	RECON SOLN
Ethylol 500 MG RECON SOLN	IV	RECON SOLN
Etopophos 100 MG RECON SOLN	IV	RECON SOLN
Etoposide 1 GM/50ML SOLUTION	IV	SOLUTION
Etoposide 100 MG/5ML SOLUTION	IV	SOLUTION
Etoposide 500 MG/25ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Evenity 105 MG/1.17ML SOLN PRSYR	SC	SOLN PRSYR
Evkeeza 1200 MG/8ML SOLUTION	IV	SOLUTION
Evkeeza 345 MG/2.3ML SOLUTION	IV	SOLUTION
Evomela 50 MG RECON SOLN	IV	RECON SOLN
Eylea 2 MG/0.05ML SOLN PRSYR	IZ	SOLN PRSYR
Eylea 2 MG/0.05ML SOLUTION	IZ	SOLUTION
Eylea HD 8 MG/0.07ML SOLUTION	IZ	SOLUTION
Fabrazyme 35 MG RECON SOLN	IV	RECON SOLN
Fabrazyme 5 MG RECON SOLN	IV	RECON SOLN
Fensolvi (6 Month) 45 MG KIT	SC	KIT
FentaNYL Citrate (PF) 100 MCG/2ML SOLN CART	IJ	SOLN CART
FentaNYL Citrate (PF) 100 MCG/2ML SOLUTION	IJ	SOLUTION
FentaNYL Citrate (PF) 1000 MCG/20ML SOLUTION	IJ	SOLUTION
FentaNYL Citrate (PF) 250 MCG/5ML SOLUTION	IJ	SOLUTION
FentaNYL Citrate (PF) 2500 MCG/50ML SOLUTION	IJ	SOLUTION
fentaNYL Citrate (PF) 50 MCG/ML SOLUTION	IJ	SOLUTION
FentaNYL Citrate (PF) 500 MCG/10ML SOLUTION	IJ	SOLUTION
Flolan 0.5 MG RECON SOLN	IV	RECON SOLN
Flolan 1.5 MG RECON SOLN	IV	RECON SOLN
Floxuridine 0.5 GM RECON SOLN	IJ	RECON SOLN
Fludarabine Phosphate 25 MG/ML SOLUTION	IV	SOLUTION
Fludarabine Phosphate 50 MG RECON SOLN	IV	RECON SOLN
Fludarabine Phosphate 50 MG/2ML SOLUTION	IV	SOLUTION
Fluorouracil 1 GM/20ML SOLUTION	IV	SOLUTION
Fluorouracil 2.5 GM/50ML SOLUTION	IV	SOLUTION
Fluorouracil 5 GM/100ML SOLUTION	IV	SOLUTION
Fluorouracil 500 MG/10ML SOLUTION	IV	SOLUTION
Folotyn 20 MG/ML SOLUTION	IV	SOLUTION
Folotyn 40 MG/2ML SOLUTION	IV	SOLUTION
Formoterol Fumarate 20 MCG/2ML NEBU SOLN	IN	NEBU SOLN
Fosaprepitant Dimeglumine 150 MG RECON SOLN	IV	RECON SOLN
Fosphenytoin Sodium 100 MG PE/2ML SOLUTION	IJ	SOLUTION
Fosphenytoin Sodium 500 MG PE/10ML SOLUTION	IJ	SOLUTION
FreAmine HBC 6.9 % SOLUTION	IV	SOLUTION
FreAmine III 10 % SOLUTION	IV	SOLUTION
Fulphila 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Fusilev 50 MG RECON SOLN	IV	RECON SOLN
Fyarro 100 MG RECON SUSP	IV	RECON SUSP
Fylnetra 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Gablofen 10000 MCG/20ML SOLN PRSYR	IT	SOLN PRSYR
Gablofen 10000 MCG/20ML SOLUTION	IT	SOLUTION
Gablofen 20000 MCG/20ML SOLN PRSYR	IT	SOLN PRSYR
Gablofen 20000 MCG/20ML SOLUTION	IT	SOLUTION
Gablofen 40000 MCG/20ML SOLN PRSYR	IT	SOLN PRSYR

MEDICATION NAME	ROUTE	DOSE FORM
Gablofen 40000 MCG/20ML SOLUTION	IT	SOLUTION
Gablofen 50 MCG/ML SOLN PRSYR	IT	SOLN PRSYR
Gamifant 10 MG/2ML SOLUTION	IV	SOLUTION
Gamifant 100 MG/20ML SOLUTION	IV	SOLUTION
Gamifant 50 MG/10ML SOLUTION	IV	SOLUTION
Ganciclovir 500 MG/250ML SOLUTION	IV	SOLUTION
Ganciclovir Sodium 500 MG RECON SOLN	IV	RECON SOLN
Ganciclovir Sodium 500 MG/10ML SOLUTION	IV	SOLUTION
Gazyva 1000 MG/40ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 1 GM RECON SOLN	IV	RECON SOLN
Gemcitabine HCl 1 GM/10ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 1 GM/26.3ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 1.5 GM/15ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 2 GM RECON SOLN	IV	RECON SOLN
Gemcitabine HCl 2 GM/20ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 2 GM/52.6ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 200 MG RECON SOLN	IV	RECON SOLN
Gemcitabine HCl 200 MG/2ML SOLUTION	IV	SOLUTION
Gemcitabine HCl 200 MG/5.26ML SOLUTION	IV	SOLUTION
Gengraf 100 MG CAP	PO	CAP
Gengraf 100 MG/ML SOLUTION	PO	SOLUTION
Gengraf 25 MG CAP	PO	CAP
Gentamicin in Saline 0.8-0.9 MG/ML-% SOLUTION	IV	SOLUTION
Gentamicin in Saline 1-0.9 MG/ML-% SOLUTION	IV	SOLUTION
Gentamicin in Saline 1.2-0.9 MG/ML-% SOLUTION	IV	SOLUTION
Gentamicin in Saline 1.6-0.9 MG/ML-% SOLUTION	IV	SOLUTION
Gentamicin in Saline 2-0.9 MG/ML-% SOLUTION	IV	SOLUTION
Givlaari 189 MG/ML SOLUTION	SC	SOLUTION
Glassia 1000 MG/50ML SOLUTION	IV	SOLUTION
Glycophos 1 MMOLE/ML SOLUTION	IV	SOLUTION
Goprelto 40 MG/ML SOLUTION	NA	SOLUTION
Granisetron HCl 1 MG TAB	PO	TAB
Granisetron HCl 1 MG/ML SOLUTION	IV	SOLUTION
Granisetron HCl 4 MG/4ML SOLUTION	IV	SOLUTION
Granix 300 MCG/0.5ML SOLN PRSYR	SC	SOLN PRSYR
Granix 300 MCG/ML SOLUTION	SC	SOLUTION
Granix 480 MCG/0.8ML SOLN PRSYR	SC	SOLN PRSYR
Granix 480 MCG/1.6ML SOLUTION	SC	SOLUTION
Halaven 1 MG/2ML SOLUTION	IV	SOLUTION
Hectorol 2 MCG/ML SOLUTION	IV	SOLUTION
Hectorol 4 MCG/2ML SOLUTION	IV	SOLUTION
HepaGam B 312 UNIT/ML SOLUTION	IJ	SOLUTION
Heparin (Porcine) in NaCl 1000-0.9 UT/500ML-% SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Heparin (Porcine) in NaCl 12500-0.45 UT/250ML-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 2000-0.9 UNIT/L-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 2500-0.9 UT/500ML-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 25000-0.45 UT/250ML-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 25000-0.45 UT/500ML-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 30000-0.9 UNIT/L-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 4000-0.9 UNIT/L-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 500-0.9 UT/500ML-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 5000-0.9 UNIT/L-% SOLUTION	IV	SOLUTION
Heparin (Porcine) in NaCl 5000-0.9 UT/500ML-% SOLUTION	IV	SOLUTION
Heparin Sod (Porcine) in D5W 100 UNIT/ML SOLUTION	IV	SOLUTION
Heparin Sod (Porcine) in D5W 25000-5 UT/500ML-% SOLUTION	IV	SOLUTION
Heparin Sod (Porcine) in D5W 40-5 UNIT/ML-% SOLUTION	IV	SOLUTION
Heparin Sodium (Porcine) 1000 UNIT/ML SOLUTION	IJ	SOLUTION
Heparin Sodium (Porcine) 10000 UNIT/ML SOLUTION	IJ	SOLUTION
Heparin Sodium (Porcine) 20000 UNIT/ML SOLUTION	IJ	SOLUTION
Heparin Sodium (Porcine) 5000 UNIT/0.5ML SOLN PRSYR	IJ	SOLN PRSYR
Heparin Sodium (Porcine) 5000 UNIT/ML SOLUTION	IJ	SOLUTION
Heparin Sodium (Porcine) PF 5000 UNIT/0.5ML SOLUTION	IJ	SOLUTION
Heparin Sodium (Porcine) PF 5000 UNIT/ML SOLUTION	IJ	SOLUTION
Hepatamine 8 % SOLUTION	IV	SOLUTION
Heplisav-B 20 MCG/0.5ML SOLN PRSYR	IM	SOLN PRSYR
Herceptin 150 MG RECON SOLN	IV	RECON SOLN
Herceptin Hylecta 600-10000 MG-UNT/5ML SOLUTION	SC	SOLUTION
Herzuma 150 MG RECON SOLN	IV	RECON SOLN
Herzuma 420 MG RECON SOLN	IV	RECON SOLN
Hycamtin 4 MG RECON SOLN	IV	RECON SOLN
HYDROmorphone HCl 0.2 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl 1 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl 2 MG/ML SOLUTION	IJ	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
HYDROmorphone HCl 4 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 1 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 10 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 2 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 4 MG/ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 50 MG/5ML SOLUTION	IJ	SOLUTION
HYDROmorphone HCl PF 500 MG/50ML SOLUTION	IJ	SOLUTION
HyperHEP B 110 UNIT/0.5ML SOLN PRSYR	IM	SOLN PRSYR
HyperHEP B 220 UNIT/ML SOLN PRSYR	IM	SOLN PRSYR
HyperHEP B 220 UNIT/ML SOLUTION	IM	SOLUTION
Hyperlyte-CR CONC	IV	CONC
HyperRAB 1500 UNIT/5ML SOLUTION	IJ	SOLUTION
HyperRAB 300 UNIT/ML SOLUTION	IJ	SOLUTION
HyperRAB 900 UNIT/3ML SOLUTION	IJ	SOLUTION
HyperRAB S/D 1500 UNIT/10ML SOLUTION	IJ	SOLUTION
HyperRAB S/D 300 UNIT/2ML SOLUTION	IJ	SOLUTION
HyperRHO S/D 1500 UNIT SOLN PRSYR	IM	SOLN PRSYR
HyperRHO S/D 250 UNIT SOLN PRSYR	IM	SOLN PRSYR
Hyqvia 10 GM/100ML KIT	SC	KIT
Hyqvia 2.5 GM/25ML KIT	SC	KIT
Hyqvia 20 GM/200ML KIT	SC	KIT
Hyqvia 30 GM/300ML KIT	SC	KIT
Hyqvia 5 GM/50ML KIT	SC	KIT
Ibandronate Sodium 3 MG/3ML SOLUTION	IV	SOLUTION
Idamycin PFS 10 MG/10ML SOLUTION	IV	SOLUTION
Idamycin PFS 20 MG/20ML SOLUTION	IV	SOLUTION
Idamycin PFS 5 MG/5ML SOLUTION	IV	SOLUTION
IDArubicin HCl 10 MG/10ML SOLUTION	IV	SOLUTION
IDArubicin HCl 20 MG/20ML SOLUTION	IV	SOLUTION
IDArubicin HCl 5 MG/5ML SOLUTION	IV	SOLUTION
Ifex 1 GM RECON SOLN	IV	RECON SOLN
Ifex 3 GM RECON SOLN	IV	RECON SOLN
Ifosfamide 1 GM RECON SOLN	IV	RECON SOLN
Ifosfamide 1 GM/20ML SOLUTION	IV	SOLUTION
Ifosfamide 3 GM RECON SOLN	IV	RECON SOLN
Ifosfamide 3 GM/60ML SOLUTION	IV	SOLUTION
Imfinzi 120 MG/2.4ML SOLUTION	IV	SOLUTION
Imfinzi 500 MG/10ML SOLUTION	IV	SOLUTION
Imjudo 25 MG/1.25ML SOLUTION	IV	SOLUTION
Imjudo 300 MG/15ML SOLUTION	IV	SOLUTION
Imlygic 1000000 UNIT/ML SUSPENSION	LS	SUSPENSION
Imlygic 100000000 UNIT/ML SUSPENSION	LS	SUSPENSION
Imogam Rabies-HT 300 UNIT/2ML SOLUTION	IJ	SOLUTION
Imuran 50 MG TAB	PO	TAB

MEDICATION NAME	ROUTE	DOSE FORM
Inflectra 100 MG RECON SOLN	IV	RECON SOLN
inFLIXimab 100 MG RECON SOLN	IV	RECON SOLN
Infugem 1200-0.9 MG/120ML-% SOLUTION	IV	SOLUTION
Infugem 1300-0.9 MG/130ML-% SOLUTION	IV	SOLUTION
Infugem 1400-0.9 MG/140ML-% SOLUTION	IV	SOLUTION
Infugem 1500-0.9 MG/150ML-% SOLUTION	IV	SOLUTION
Infugem 1600-0.9 MG/160ML-% SOLUTION	IV	SOLUTION
Infugem 1700-0.9 MG/170ML-% SOLUTION	IV	SOLUTION
Infugem 1800-0.9 MG/180ML-% SOLUTION	IV	SOLUTION
Infugem 1900-0.9 MG/190ML-% SOLUTION	IV	SOLUTION
Infugem 2000-0.9 MG/200ML-% SOLUTION	IV	SOLUTION
Infugem 2200-0.9 MG/220ML-% SOLUTION	IV	SOLUTION
Infumorph 200 200 MG/20ML (10 MG/ML) SOLUTION	IJ	SOLUTION
Infumorph 500 500 MG/20ML (25 MG/ML) SOLUTION	IJ	SOLUTION
Intralipid 20 % EMULSION	IV	EMULSION
Intralipid 30 % EMULSION	IV	EMULSION
Ionosol-MB in D5W SOLUTION	IV	SOLUTION
Ipratropium Bromide 0.02 % SOLUTION	IN	SOLUTION
Ipratropium-Albuterol 0.5-2.5 (3) MG/3ML SOLUTION	IN	SOLUTION
Irinotecan HCl 100 MG/5ML SOLUTION	IV	SOLUTION
Irinotecan HCl 300 MG/15ML SOLUTION	IV	SOLUTION
Irinotecan HCl 40 MG/2ML SOLUTION	IV	SOLUTION
Irinotecan HCl 500 MG/25ML SOLUTION	IV	SOLUTION
Isolyte-P in D5W SOLUTION	IV	SOLUTION
Isolyte-S SOLUTION	IV	SOLUTION
Isolyte-S pH 7.4 SOLUTION	IV	SOLUTION
Istodax 10 MG RECON SOLN	IV	RECON SOLN
Ixempra Kit 15 MG RECON SOLN	IV	RECON SOLN
Ixempra Kit 45 MG RECON SOLN	IV	RECON SOLN
Jelmyto 80 (2 x 40) MG RECON SOLN	UL	RECON SOLN
Jemperli 500 MG/10ML SOLUTION	IV	SOLUTION
Jevtana 60 MG/1.5ML SOLUTION	IV	SOLUTION
Kabiven 3.3-9.8-3.9-0.7 % EMULSION	IV	EMULSION
Kadcyla 100 MG RECON SOLN	IV	RECON SOLN
Kadcyla 160 MG RECON SOLN	IV	RECON SOLN
Kanjinti 150 MG RECON SOLN	IV	RECON SOLN
Kanjinti 420 MG RECON SOLN	IV	RECON SOLN
Kanuma 20 MG/10ML SOLUTION	IV	SOLUTION
KCl (0.149%) in NaCl 20-0.45 MEQ/L-% SOLUTION	IV	SOLUTION
KCl (in NaCl 0.9%) 10 MEQ/100ML SOLUTION	IV	SOLUTION
KCl (in NaCl 0.9%) 20 MEQ/250ML SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
KCl (in NaCl 0.9%) 40 MEQ/250ML SOLUTION	IV	SOLUTION
KCl (in NaCl 0.9%) 40 MEQ/500ML SOLUTION	IV	SOLUTION
KCl in D5W Lactated Ringers 40 MEQ/L SOLUTION	IV	SOLUTION
KCl in Dextrose-NaCl 10-5-0.45 MEQ/L-%-% SOLUTION	IV	SOLUTION
KCl in Dextrose-NaCl 30-5-0.45 MEQ/L-%-% SOLUTION	IV	SOLUTION
KCl in Dextrose-NaCl 40-5-0.45 MEQ/L-%-% SOLUTION	IV	SOLUTION
Kedrab 1500 UNIT/10ML SOLUTION	IJ	SOLUTION
Kedrab 300 UNIT/2ML SOLUTION	IJ	SOLUTION
Kenalog 10 MG/ML SUSPENSION	IJ	SUSPENSION
Kenalog 40 MG/ML SUSPENSION	IJ	SUSPENSION
Kenalog-80 80 MG/ML SUSPENSION	IJ	SUSPENSION
Kepivance 6.25 MG RECON SOLN	IV	RECON SOLN
Keppra 500 MG/5ML SOLUTION	IV	SOLUTION
Ketorolac Tromethamine 15 MG/ML SOLUTION	IJ	SOLUTION
Ketorolac Tromethamine 30 MG/ML SOLUTION	IJ	SOLUTION
Ketorolac Tromethamine 60 MG/2ML SOLUTION	IM	SOLUTION
Keytruda 100 MG/4ML SOLUTION	IV	SOLUTION
Khapzory 175 MG RECON SOLN	IV	RECON SOLN
Khapzory 300 MG RECON SOLN	IV	RECON SOLN
Kimmtrak 100 MCG/0.5ML SOLUTION	IV	SOLUTION
Kimyrsa 1200 MG RECON SOLN	IV	RECON SOLN
Kitabis Pak 300 MG/5ML NEBU SOLN	IN	NEBU SOLN
Krystexxa 8 MG/ML SOLUTION	IV	SOLUTION
Kyleena 19.5 MG IUD	IU	IUD
Kyprolis 10 MG RECON SOLN	IV	RECON SOLN
Kyprolis 30 MG RECON SOLN	IV	RECON SOLN
Kyprolis 60 MG RECON SOLN	IV	RECON SOLN
Labetalol HCl 5 MG/ML SOLUTION	IV	SOLUTION
Lacosamide 200 MG/20ML SOLUTION	IV	SOLUTION
Lamzedo 10 MG RECON SOLN	IV	RECON SOLN
Lanoxin 0.25 MG/ML SOLUTION	IJ	SOLUTION
Lanoxin Pediatric 0.1 MG/ML SOLUTION	IJ	SOLUTION
Lartruvo 190 MG/19ML SOLUTION	IV	SOLUTION
Lartruvo 500 MG/50ML SOLUTION	IV	SOLUTION
Lemtrada 12 MG/1.2ML SOLUTION	IV	SOLUTION
Leqvio 284 MG/1.5ML SOLN PRSYR	SC	SOLN PRSYR
LevETIRAcetam 500 MG/5ML SOLUTION	IV	SOLUTION
LevETIRAcetam in NaCl 1000 MG/100ML SOLUTION	IV	SOLUTION
LevETIRAcetam in NaCl 1500 MG/100ML SOLUTION	IV	SOLUTION
levETIRAcetam in NaCl 250 MG/50ML SOLUTION	IV	SOLUTION
LevETIRAcetam in NaCl 500 MG/100ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
levOCARNitine 200 MG/ML SOLUTION	IV	SOLUTION
LevoFLOXacin in D5W 250 MG/50ML SOLUTION	IV	SOLUTION
LEVOleucovorin Calcium 50 MG RECON SOLN	IV	RECON SOLN
LEVOleucovorin Calcium PF 175 MG/17.5ML SOLUTION	IV	SOLUTION
LEVOleucovorin Calcium PF 250 MG/25ML SOLUTION	IV	SOLUTION
Levothyroxine Sodium 100 MCG RECON SOLN	IV	RECON SOLN
Levothyroxine Sodium 100 MCG/5ML SOLUTION	IV	SOLUTION
Levothyroxine Sodium 100 MCG/ML SOLUTION	IV	SOLUTION
Levothyroxine Sodium 200 MCG RECON SOLN	IV	RECON SOLN
Levothyroxine Sodium 200 MCG/5ML SOLUTION	IV	SOLUTION
Levothyroxine Sodium 500 MCG RECON SOLN	IV	RECON SOLN
Levothyroxine Sodium 500 MCG/5ML SOLUTION	IV	SOLUTION
Libtayo 350 MG/7ML SOLUTION	IV	SOLUTION
Lidocaine HCl (Cardiac) 100 MG/5ML SOLN PRSYR	IV	SOLN PRSYR
Lidocaine HCl (Cardiac) 50 MG/5ML SOLN PRSYR	IV	SOLN PRSYR
Lidocaine HCl (Cardiac) PF 100 MG/5ML SOLN PRSYR	IV	SOLN PRSYR
Lidocaine HCl (Cardiac) PF 100 MG/5ML SOLUTION	IV	SOLUTION
Lidocaine HCl (Cardiac) PF 50 MG/5ML SOLN PRSYR	IV	SOLN PRSYR
Lidocaine in D5W 4-5 MG/ML-% SOLUTION	IV	SOLUTION
Lidocaine-Epinephrine 0.5 %-1:200000 SOLUTION	IJ	SOLUTION
Lidocaine-Epinephrine 1 %-1:100000 SOLUTION	IJ	SOLUTION
Lidocaine-Epinephrine 1.5 %-1:200000 SOLUTION	IJ	SOLUTION
Lidocaine-Epinephrine 2 %-1:100000 SOLUTION	IJ	SOLUTION
Lidocaine-Epinephrine 2 %-1:200000 SOLUTION	IJ	SOLUTION
Lidocaine-Epinephrine 2 %-1:50000 SOLUTION	IJ	SOLUTION
Liletta (52 MG) 20.1 MCG/DAY IUD	IU	IUD
Lioresal 0.05 MG/ML SOLUTION	IT	SOLUTION
Lioresal 10 MG/20ML SOLUTION	IT	SOLUTION
Lioresal 10 MG/5ML SOLUTION	IT	SOLUTION
Lioresal 40 MG/20ML SOLUTION	IT	SOLUTION
Liothyronine Sodium 10 MCG/ML SOLUTION	IV	SOLUTION
LORazepam 2 MG/ML SOLUTION	IJ	SOLUTION
LORazepam 4 MG/ML SOLUTION	IJ	SOLUTION
Lucentis 0.3 MG/0.05ML SOLN PRSYR	IZ	SOLN PRSYR
Lucentis 0.3 MG/0.05ML SOLUTION	IZ	SOLUTION
Lucentis 0.5 MG/0.05ML SOLN PRSYR	IZ	SOLN PRSYR
Lucentis 0.5 MG/0.05ML SOLUTION	IZ	SOLUTION
Lumizyme 50 MG RECON SOLN	IV	RECON SOLN
Lumoxiti 1 MG RECON SOLN	IV	RECON SOLN
Lunsumio 1 MG/ML SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
Lunsumio 30 MG/30ML SOLUTION	IV	SOLUTION
Lupron Depot-Ped (6-Month) 45 MG KIT	IM	KIT
Magnesium Sulfate 2 GM/50ML SOLUTION	IV	SOLUTION
Magnesium Sulfate 20 GM/500ML SOLUTION	IV	SOLUTION
Magnesium Sulfate 4 GM/100ML SOLUTION	IV	SOLUTION
Magnesium Sulfate 4 GM/50ML SOLUTION	IV	SOLUTION
Magnesium Sulfate 40 GM/1000ML SOLUTION	IV	SOLUTION
Magnesium Sulfate in D5W 1-5 GM/100ML-% SOLUTION	IV	SOLUTION
Magnesium Sulfate-NaCl 2-0.9 GM/50ML-% SOLUTION	IV	SOLUTION
Magnesium Sulfate-NaCl 20-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Magnesium Sulfate-NaCl 3-0.9 GM/150ML-% SOLUTION	IV	SOLUTION
Manganese Chloride 0.1 MG/ML SOLUTION	IV	SOLUTION
Manganese Sulfate 0.1 MG/ML SOLUTION	IV	SOLUTION
Mannitol 20 % SOLUTION	IV	SOLUTION
Mannitol 25 % SOLUTION	IV	SOLUTION
Margenza 250 MG/10ML SOLUTION	IV	SOLUTION
Marinol 10 MG CAP	PO	CAP
Marinol 2.5 MG CAP	PO	CAP
Marinol 5 MG CAP	PO	CAP
Marqibo 5 MG/31ML SUSPENSION	IV	SUSPENSION
Melphalan 2 MG TAB	PO	TAB
Melphalan HCl 50 MG RECON SOLN	IV	RECON SOLN
Meperidine HCl 100 MG/ML SOLUTION	IJ	SOLUTION
Meperidine HCl 25 MG/ML SOLUTION	IJ	SOLUTION
Meperidine HCl 50 MG/ML SOLUTION	IJ	SOLUTION
Mepsevii 10 MG/5ML SOLUTION	IV	SOLUTION
Methocarbamol 1000 MG/10ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium (PF) 1 GM/40ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium (PF) 250 MG/10ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium (PF) 50 MG/2ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium 1 GM RECON SOLN	IJ	RECON SOLN
Methotrexate Sodium 1000 MG/40ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium 250 MG/10ML SOLUTION	IJ	SOLUTION
Methotrexate Sodium 50 MG/2ML SOLUTION	IJ	SOLUTION
MethylPREDNISolone Acetate 50 MG/ML SUSPENSION	IJ	SUSPENSION
MethylPREDNISolone Sodium Succ 1000 MG RECON SOLN	IJ	RECON SOLN
MethylPREDNISolone Sodium Succ 125 MG RECON SOLN	IJ	RECON SOLN

MEDICATION NAME	ROUTE	DOSE FORM
methyLPREDNISolone Sodium Succ 500 MG RECON SOLN	IJ	RECON SOLN
Metoprolol Tartrate 5 MG/5ML SOLUTION	IV	SOLUTION
Miacalcin 200 UNIT/ML SOLUTION	IJ	SOLUTION
MICRhoGAM Ultra-Filtered Plus 250 UNIT SOLN PRSYR	IM	SOLN PRSYR
Milrinone Lactate 10 MG/10ML SOLUTION	IV	SOLUTION
Milrinone Lactate 20 MG/20ML SOLUTION	IV	SOLUTION
Milrinone Lactate 50 MG/50ML SOLUTION	IV	SOLUTION
Milrinone Lactate in Dextrose 20-5 MG/100ML-% SOLUTION	IV	SOLUTION
Milrinone Lactate in Dextrose 40-5 MG/200ML-% SOLUTION	IV	SOLUTION
Minocin 100 MG RECON SOLN	IV	RECON SOLN
Mirena (52 MG) 20 MCG/DAY IUD	IU	IUD
Mitigo 200 MG/20ML (10 MG/ML) SOLUTION	IJ	SOLUTION
Mitigo 500 MG/20ML (25 MG/ML) SOLUTION	IJ	SOLUTION
MitoMYcin 20 MG RECON SOLN	IV	RECON SOLN
MitoMYcin 20 MG/40ML SOLN PRSYR	IS	SOLN PRSYR
MitoMYcin 40 MG RECON SOLN	IV	RECON SOLN
MitoMYcin 5 MG RECON SOLN	IV	RECON SOLN
MitoXANTRONE HCl 20 MG/10ML CONC	IV	CONC
MitoXANTRONE HCl 25 MG/12.5ML CONC	IV	CONC
MitoXANTRONE HCl 30 MG/15ML CONC	IV	CONC
Monjuvi 200 MG RECON SOLN	IV	RECON SOLN
Morphine Sulfate (PF) 0.5 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 1 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 1 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate (PF) 10 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 10 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate (PF) 2 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 2 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate (PF) 4 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 4 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate (PF) 5 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 8 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate (PF) 8 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 0.5 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 1 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate 1 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 10 MG/0.7ML DEVICE	IM	DEVICE
Morphine Sulfate 10 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 150 MG/30ML SOLUTION	IV	SOLUTION
Morphine Sulfate 2 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate 4 MG/ML SOLUTION	IJ	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Morphine Sulfate 4 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 5 MG/ML SOLUTION	IJ	SOLUTION
Morphine Sulfate 50 MG/ML SOLUTION	IV	SOLUTION
Morphine Sulfate 8 MG/ML SOLUTION	IV	SOLUTION
Moxifloxacin HCl 400 MG/250ML SOLUTION	IV	SOLUTION
Moxifloxacin HCl in NaCl 400 MG/250ML SOLUTION	IV	SOLUTION
Multiple Electro Type 1 pH 5.5 SOLUTION	IV	SOLUTION
Multiple Electro Type 1 pH 7.4 SOLUTION	IV	SOLUTION
Mutamycin 20 MG RECON SOLN	IV	RECON SOLN
Mutamycin 40 MG RECON SOLN	IV	RECON SOLN
Mutamycin 5 MG RECON SOLN	IV	RECON SOLN
Mvasi 100 MG/4ML SOLUTION	IV	SOLUTION
Mvasi 400 MG/16ML SOLUTION	IV	SOLUTION
Mycophenolate Mofetil 200 MG/ML RECON SUSP	PO	RECON SUSP
Mycophenolate Mofetil 250 MG CAP	PO	CAP
Mycophenolate Mofetil 500 MG RECON SOLN	IV	RECON SOLN
Mycophenolate Mofetil 500 MG TAB	PO	TAB
Mycophenolate Mofetil HCl 500 MG RECON SOLN	IV	RECON SOLN
Mycophenolate Sodium 180 MG TAB DR	PO	TAB DR
Mycophenolate Sodium 360 MG TAB DR	PO	TAB DR
Myfortic 180 MG TAB DR	PO	TAB DR
Myfortic 360 MG TAB DR	PO	TAB DR
Mylotarg 4.5 MG RECON SOLN	IV	RECON SOLN
Myobloc 10000 UNIT/2ML SOLUTION	IM	SOLUTION
Myobloc 2500 UNIT/0.5ML SOLUTION	IM	SOLUTION
Myobloc 5000 UNIT/ML SOLUTION	IM	SOLUTION
Nabi-HB 312 UNIT/ML SOLUTION	IM	SOLUTION
Nafcillin Sodium in Dextrose 1 GM/50ML SOLUTION	IV	SOLUTION
Nafcillin Sodium in Dextrose 2 GM/100ML SOLUTION	IV	SOLUTION
Naglazyme 1 MG/ML SOLUTION	IV	SOLUTION
Nalbuphine HCl 10 MG/ML SOLUTION	IJ	SOLUTION
Nalbuphine HCl 20 MG/ML SOLUTION	IJ	SOLUTION
Navelbine 10 MG/ML SOLUTION	IV	SOLUTION
Navelbine 50 MG/5ML SOLUTION	IV	SOLUTION
Nebupent 300 MG RECON SOLN	IN	RECON SOLN
Nelarabine 5 MG/ML SOLUTION	IV	SOLUTION
Neoral 100 MG CAP	PO	CAP
Neoral 100 MG/ML SOLUTION	PO	SOLUTION
Neoral 25 MG CAP	PO	CAP
NephrAmine 5.4 % SOLUTION	IV	SOLUTION
Neulasta 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Neulasta Onpro 6 MG/0.6ML PREF SY KT	SC	PREF SY KT
Neupogen 300 MCG/0.5ML SOLN PRSYR	IJ	SOLN PRSYR

MEDICATION NAME	ROUTE	DOSE FORM
Neupogen 300 MCG/ML SOLUTION	IJ	SOLUTION
Neupogen 480 MCG/0.8ML SOLN PRSYR	IJ	SOLN PRSYR
Neupogen 480 MCG/1.6ML SOLUTION	IJ	SOLUTION
Nexplanon 68 MG IMPLANT	SC	IMPLANT
Nexterone 150-4.21 MG/100ML-% SOLUTION	IV	SOLUTION
Nexterone 360-4.14 MG/200ML-% SOLUTION	IV	SOLUTION
Nexvazyme 100 MG RECON SOLN	IV	RECON SOLN
NiCARDipine HCl 2.5 MG/ML SOLUTION	IV	SOLUTION
niCARDipine HCl in NaCl 20-0.9 MG/200ML-% SOLUTION	IV	SOLUTION
niCARDipine HCl in NaCl 40-0.9 MG/200ML-% SOLUTION	IV	SOLUTION
Nipent 10 MG RECON SOLN	IV	RECON SOLN
Nitroglycerin 5 MG/ML SOLUTION	IV	SOLUTION
Nitroglycerin in D5W 100-5 MCG/ML-% SOLUTION	IV	SOLUTION
Nitroglycerin in D5W 200-5 MCG/ML-% SOLUTION	IV	SOLUTION
Nitroglycerin in D5W 400-5 MCG/ML-% SOLUTION	IV	SOLUTION
Nivestym 300 MCG/0.5ML SOLN PRSYR	IJ	SOLN PRSYR
Nivestym 300 MCG/ML SOLUTION	IJ	SOLUTION
Nivestym 480 MCG/0.8ML SOLN PRSYR	IJ	SOLN PRSYR
Nivestym 480 MCG/1.6ML SOLUTION	IJ	SOLUTION
Normosol-R SOLUTION	IV	SOLUTION
Normosol-R in D5W SOLUTION	IV	SOLUTION
Normosol-R pH 7.4 SOLUTION	IV	SOLUTION
Novarel 10000 UNIT RECON SOLN	IM	RECON SOLN
Novarel 5000 UNIT RECON SOLN	IM	RECON SOLN
Noxafil 300 MG/16.7ML SOLUTION	IV	SOLUTION
Nplate 125 MCG RECON SOLN	SC	RECON SOLN
Nplate 250 MCG RECON SOLN	SC	RECON SOLN
Nplate 500 MCG RECON SOLN	SC	RECON SOLN
Nulibry 9.5 MG RECON SOLN	IV	RECON SOLN
Nulojix 250 MG RECON SOLN	IV	RECON SOLN
Numbrino 40 MG/ML SOLUTION	NA	SOLUTION
Nutrilipid 20 % EMULSION	IV	EMULSION
Nuzyra 100 MG RECON SOLN	IV	RECON SOLN
Nyvepria 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Ocrevus 300 MG/10ML SOLUTION	IV	SOLUTION
Octagam 1 GM/20ML SOLUTION	IV	SOLUTION
Octagam 10 GM/100ML SOLUTION	IV	SOLUTION
Octagam 10 GM/200ML SOLUTION	IV	SOLUTION
Octagam 2 GM/20ML SOLUTION	IV	SOLUTION
Octagam 2.5 GM/50ML SOLUTION	IV	SOLUTION
Octagam 20 GM/200ML SOLUTION	IV	SOLUTION
Octagam 25 GM/500ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Octagam 30 GM/300ML SOLUTION	IV	SOLUTION
Octagam 5 GM/100ML SOLUTION	IV	SOLUTION
Octagam 5 GM/50ML SOLUTION	IV	SOLUTION
Ofirmev 10 MG/ML SOLUTION	IV	SOLUTION
Ogivri 150 MG RECON SOLN	IV	RECON SOLN
Ogivri 420 MG RECON SOLN	IV	RECON SOLN
Omegaven 10 GM/100ML EMULSION	IV	EMULSION
Omegaven 5 GM/50ML EMULSION	IV	EMULSION
Oncaspar 750 UNIT/ML SOLUTION	IJ	SOLUTION
Ondansetron 4 MG TAB DISP	PO	TAB DISP
Ondansetron 8 MG TAB DISP	PO	TAB DISP
Ondansetron HCl 24 MG TAB	PO	TAB
Ondansetron HCl 4 MG TAB	PO	TAB
Ondansetron HCl 4 MG/2ML SOLN PRSYR	IJ	SOLN PRSYR
Ondansetron HCl 4 MG/2ML SOLUTION	IJ	SOLUTION
Ondansetron HCl 4 MG/5ML SOLUTION	PO	SOLUTION
Ondansetron HCl 40 MG/20ML SOLUTION	IJ	SOLUTION
Ondansetron HCl 8 MG TAB	PO	TAB
Onivyde 43 MG/10ML INJECTABLE	IV	INJECTABLE
Onpattro 10 MG/5ML SOLUTION	IV	SOLUTION
Ontruzant 150 MG RECON SOLN	IV	RECON SOLN
Ontruzant 420 MG RECON SOLN	IV	RECON SOLN
Opdivo 100 MG/10ML SOLUTION	IV	SOLUTION
Opdivo 120 MG/12ML SOLUTION	IV	SOLUTION
Opdivo 240 MG/24ML SOLUTION	IV	SOLUTION
Opdivo 40 MG/4ML SOLUTION	IV	SOLUTION
Opdualag 240-80 MG/20ML SOLUTION	IV	SOLUTION
Orphenadrine Citrate 30 MG/ML SOLUTION	IJ	SOLUTION
Osmitrol 10 % SOLUTION	IV	SOLUTION
Osmitrol 15 % SOLUTION	IV	SOLUTION
Osmitrol 20 % SOLUTION	IV	SOLUTION
Osmitrol 5 % SOLUTION	IV	SOLUTION
Oxacillin Sodium 1 GM RECON SOLN	IJ	RECON SOLN
Oxacillin Sodium 10 GM RECON SOLN	IV	RECON SOLN
Oxacillin Sodium 2 GM RECON SOLN	IJ	RECON SOLN
Oxacillin Sodium in Dextrose 1 GM/50ML SOLUTION	IV	SOLUTION
Oxacillin Sodium in Dextrose 2 GM/50ML SOLUTION	IV	SOLUTION
Oxaliplatin 100 MG RECON SOLN	IV	RECON SOLN
Oxaliplatin 100 MG/20ML SOLUTION	IV	SOLUTION
Oxaliplatin 200 MG/40ML SOLUTION	IV	SOLUTION
Oxaliplatin 50 MG RECON SOLN	IV	RECON SOLN
Oxaliplatin 50 MG/10ML SOLUTION	IV	SOLUTION
Oxlumo 94.5 MG/0.5ML SOLUTION	SC	SOLUTION
PACLitaxel 100 MG/16.7ML CONC	IV	CONC

MEDICATION NAME	ROUTE	DOSE FORM
PACLitaxel 150 MG/25ML CONC	IV	CONC
PACLitaxel 30 MG/5ML CONC	IV	CONC
PACLitaxel 300 MG/50ML CONC	IV	CONC
PACLitaxel Protein-Bound Part 100 MG RECON SUSP	IV	RECON SUSP
Padcev 20 MG RECON SOLN	IV	RECON SOLN
Padcev 30 MG RECON SOLN	IV	RECON SOLN
Palonosetron HCl 0.25 MG/2ML SOLUTION	IV	SOLUTION
Palonosetron HCl 0.25 MG/5ML SOLN PRSYR	IV	SOLN PRSYR
Palonosetron HCl 0.25 MG/5ML SOLUTION	IV	SOLUTION
Pamidronate Disodium 30 MG RECON SOLN	IV	RECON SOLN
Pamidronate Disodium 30 MG/10ML SOLUTION	IV	SOLUTION
Pamidronate Disodium 6 MG/ML SOLUTION	IV	SOLUTION
Pamidronate Disodium 90 MG RECON SOLN	IV	RECON SOLN
Pamidronate Disodium 90 MG/10ML SOLUTION	IV	SOLUTION
Panzyga 1 GM/10ML SOLUTION	IV	SOLUTION
Panzyga 10 GM/100ML SOLUTION	IV	SOLUTION
Panzyga 2.5 GM/25ML SOLUTION	IV	SOLUTION
Panzyga 20 GM/200ML SOLUTION	IV	SOLUTION
Panzyga 30 GM/300ML SOLUTION	IV	SOLUTION
Panzyga 5 GM/50ML SOLUTION	IV	SOLUTION
Paragard Intrauterine Copper IUD	IU	IUD
Paraplatin 1000 MG/100ML SOLUTION	IV	SOLUTION
Paraplatin 150 MG/15ML SOLUTION	IV	SOLUTION
Paraplatin 450 MG/45ML SOLUTION	IV	SOLUTION
Paraplatin 50 MG/5ML SOLUTION	IV	SOLUTION
Paraplatin 600 MG/60ML SOLUTION	IV	SOLUTION
Paricalcitol 1 MCG CAP	PO	CAP
Paricalcitol 2 MCG CAP	PO	CAP
Paricalcitol 2 MCG/ML SOLUTION	IV	SOLUTION
Paricalcitol 4 MCG CAP	PO	CAP
Paricalcitol 5 MCG/ML SOLUTION	IV	SOLUTION
Pedmark 12.5 % SOLUTION	IV	SOLUTION
PEMEtrexed 1 GM/40ML SOLUTION	IV	SOLUTION
PEMEtrexed 100 MG/4ML SOLUTION	IV	SOLUTION
PEMEtrexed 500 MG/20ML SOLUTION	IV	SOLUTION
PEMEtrexed Disodium 1 GM/40ML SOLUTION	IV	SOLUTION
PEMEtrexed Disodium 100 MG RECON SOLN	IV	RECON SOLN
PEMEtrexed Disodium 100 MG/4ML SOLUTION	IV	SOLUTION
PEMEtrexed Disodium 1000 MG RECON SOLN	IV	RECON SOLN
PEMEtrexed Disodium 500 MG RECON SOLN	IV	RECON SOLN
PEMEtrexed Disodium 500 MG/20ML SOLUTION	IV	SOLUTION
PEMEtrexed Disodium 750 MG RECON SOLN	IV	RECON SOLN
PEMEtrexed Disodium 850 MG/34ML SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
PEMEtrexed Ditromethamine 100 MG RECON SOLN	IV	RECON SOLN
PEMEtrexed Ditromethamine 500 MG RECON SOLN	IV	RECON SOLN
Pemfexy 500 MG/20ML SOLUTION	IV	SOLUTION
Penicillin G Pot in Dextrose 20000 UNIT/ML SOLUTION	IV	SOLUTION
Penicillin G Pot in Dextrose 40000 UNIT/ML SOLUTION	IV	SOLUTION
Penicillin G Pot in Dextrose 60000 UNIT/ML SOLUTION	IV	SOLUTION
Penicillin G Procaine 600000 UNIT/ML SUSPENSION	IM	SUSPENSION
Pentamidine Isethionate 300 MG RECON SOLN	IN	RECON SOLN
Pepaxto 20 MG RECON SOLN	IV	RECON SOLN
Perforomist 20 MCG/2ML NEBU SOLN	IN	NEBU SOLN
Perikabiven 2.4-6.8-3.5-0.5 % EMULSION	IV	EMULSION
Perjeta 420 MG/14ML SOLUTION	IV	SOLUTION
Phenytoin Sodium 50 MG/ML SOLUTION	IJ	SOLUTION
Phesgo 60-60-2000 MG-MG-U/ML SOLUTION	SC	SOLUTION
Phesgo 80-40-2000 MG-MG-U/ML SOLUTION	SC	SOLUTION
Plasma-Lyte 148 SOLUTION	IV	SOLUTION
Plasma-Lyte A SOLUTION	IV	SOLUTION
Plenamine 15 % SOLUTION	IV	SOLUTION
Polivy 140 MG RECON SOLN	IV	RECON SOLN
Polivy 30 MG RECON SOLN	IV	RECON SOLN
Pombiliti 105 MG RECON SOLN	IV	RECON SOLN
Portrazza 800 MG/50ML SOLUTION	IV	SOLUTION
Posaconazole 300 MG/16.7ML SOLUTION	IV	SOLUTION
Potassium Acetate 2 MEQ/ML SOLUTION	IV	SOLUTION
Potassium Chloride 10 MEQ/50ML SOLUTION	IV	SOLUTION
Potassium Chloride 20 MEQ/50ML SOLUTION	IV	SOLUTION
Potassium Chloride in D5W 20 MEQ/250ML SOLUTION	IV	SOLUTION
Potassium Chloride in Dextrose 10-5 MEQ/L-% SOLUTION	IV	SOLUTION
Potassium Chloride in NaCl 20 MEQ/250ML SOLUTION	IV	SOLUTION
Potassium Chloride in NaCl 20-0.45 MEQ/L-% SOLUTION	IV	SOLUTION
Potassium Phosphates 15 MMOLE/5ML SOLUTION	IV	SOLUTION
Potassium Phosphates 150 MMOLE/50ML SOLUTION	IV	SOLUTION
Potassium Phosphates(66 mEq K) 45 MMOLE/15ML SOLUTION	IV	SOLUTION
Potassium Phosphates(71 mEq K) 45 MMOLE/15ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Potassium Phosphates-NaCl 15 MMOL/250ML SOLUTION	IV	SOLUTION
Poteligeo 20 MG/5ML SOLUTION	IV	SOLUTION
PRALAtrexate 20 MG/ML SOLUTION	IV	SOLUTION
PRALAtrexate 40 MG/2ML SOLUTION	IV	SOLUTION
Pregnyl 10000 UNIT RECON SOLN	IM	RECON SOLN
PreHevbrio 10 MCG/ML SUSPENSION	IM	SUSPENSION
Premasol 10 % SOLUTION	IV	SOLUTION
Prevymis 240 MG/12ML SOLUTION	IV	SOLUTION
Prevymis 480 MG/24ML SOLUTION	IV	SOLUTION
Prialt 100 MCG/ML SOLUTION	IT	SOLUTION
Prialt 500 MCG/20ML SOLUTION	IT	SOLUTION
Prialt 500 MCG/5ML SOLUTION	IT	SOLUTION
Procainamide HCl 100 MG/ML SOLUTION	IJ	SOLUTION
Procainamide HCl 500 MG/ML SOLUTION	IJ	SOLUTION
Procalamine 3 % SOLUTION	IV	SOLUTION
Prochlorperazine Edisylate 10 MG/2ML SOLUTION	IJ	SOLUTION
Prochlorperazine Edisylate 50 MG/10ML SOLUTION	IJ	SOLUTION
Procrit 10000 UNIT/ML SOLUTION	IJ	SOLUTION
Procrit 2000 UNIT/ML SOLUTION	IJ	SOLUTION
Procrit 20000 UNIT/ML SOLUTION	IJ	SOLUTION
Procrit 3000 UNIT/ML SOLUTION	IJ	SOLUTION
Procrit 4000 UNIT/ML SOLUTION	IJ	SOLUTION
Procrit 40000 UNIT/ML SOLUTION	IJ	SOLUTION
Prograf 0.5 MG CAP	PO	CAP
Prograf 1 MG CAP	PO	CAP
Prograf 5 MG CAP	PO	CAP
Prograf 5 MG/ML SOLUTION	IV	SOLUTION
Prolastin-C 1000 MG RECON SOLN	IV	RECON SOLN
Prolastin-C 1000 MG/20ML SOLUTION	IV	SOLUTION
Proleukin 22000000 UNIT RECON SOLN	IV	RECON SOLN
Propranolol HCl 1 MG/ML SOLUTION	IV	SOLUTION
Prosol 20 % SOLUTION	IV	SOLUTION
Pulmicort 0.25 MG/2ML SUSPENSION	IN	SUSPENSION
Pulmicort 0.5 MG/2ML SUSPENSION	IN	SUSPENSION
Pulmicort 1 MG/2ML SUSPENSION	IN	SUSPENSION
Pulmozyme 2.5 MG/2.5ML SOLUTION	IN	SOLUTION
Radicava 30 MG/100ML SOLUTION	IV	SOLUTION
Rapamune 0.5 MG TAB	PO	TAB
Rapamune 1 MG TAB	PO	TAB
Rapamune 1 MG/ML SOLUTION	PO	SOLUTION
Rapamune 2 MG TAB	PO	TAB
Reblozyl 25 MG RECON SOLN	SC	RECON SOLN
Reblozyl 75 MG RECON SOLN	SC	RECON SOLN



MEDICATION NAME	ROUTE	DOSE FORM
Recarbrio 1.25 GM RECON SOLN	IV	RECON SOLN
Reclast 5 MG/100ML SOLUTION	IV	SOLUTION
Recombivax HB 10 MCG/ML SUSP PRSYR	IJ	SUSP PRSYR
Recombivax HB 10 MCG/ML SUSPENSION	IJ	SUSPENSION
Recombivax HB 40 MCG/ML SUSPENSION	IJ	SUSPENSION
Recombivax HB 5 MCG/0.5ML SUSP PRSYR	IJ	SUSP PRSYR
Recombivax HB 5 MCG/0.5ML SUSPENSION	IJ	SUSPENSION
Regonol 10 MG/2ML SOLUTION	IV	SOLUTION
Releuko 300 MCG/0.5ML SOLN PRSYR	SC	SOLN PRSYR
Releuko 300 MCG/ML SOLUTION	IJ	SOLUTION
Releuko 480 MCG/0.8ML SOLN PRSYR	SC	SOLN PRSYR
Releuko 480 MCG/1.6ML SOLUTION	IJ	SOLUTION
Remicade 100 MG RECON SOLN	IV	RECON SOLN
Remodulin 100 MG/20ML SOLUTION	IJ	SOLUTION
Remodulin 20 MG/20ML SOLUTION	IJ	SOLUTION
Remodulin 200 MG/20ML SOLUTION	IJ	SOLUTION
Remodulin 50 MG/20ML SOLUTION	IJ	SOLUTION
Renflexis 100 MG RECON SOLN	IV	RECON SOLN
Revatio 10 MG/12.5ML SOLUTION	IV	SOLUTION
Rezzayo 200 MG RECON SOLN	IV	RECON SOLN
RhoGAM Ultra-Filtered Plus 1500 UNIT SOLN PRSYR	IM	SOLN PRSYR
Rhophylac 1500 UNIT/2ML SOLN PRSYR	IJ	SOLN PRSYR
Riabni 100 MG/10ML SOLUTION	IV	SOLUTION
Riabni 500 MG/50ML SOLUTION	IV	SOLUTION
Ribavirin 6 GM RECON SOLN	IN	RECON SOLN
Rituxan 100 MG/10ML SOLUTION	IV	SOLUTION
Rituxan 500 MG/50ML SOLUTION	IV	SOLUTION
Rituxan Hycela 1400-23400 MG -UT/11.7ML SOLUTION	SC	SOLUTION
Rituxan Hycela 1600-26800 MG -UT/13.4ML SOLUTION	SC	SOLUTION
Robaxin 1000 MG/10ML SOLUTION	IJ	SOLUTION
Rocaltrol 0.25 MCG CAP	PO	CAP
Rocaltrol 0.5 MCG CAP	PO	CAP
Rocaltrol 1 MCG/ML SOLUTION	PO	SOLUTION
Rolvedon 13.2 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
RomiDEPsin 10 MG RECON SOLN	IV	RECON SOLN
romiDEPsin 27.5 MG/5.5ML SOLUTION	IV	SOLUTION
Ruxience 100 MG/10ML SOLUTION	IV	SOLUTION
Ruxience 500 MG/50ML SOLUTION	IV	SOLUTION
Rybrevant 350 MG/7ML SOLUTION	IV	SOLUTION
Rykindo 25 MG SRER	IM	
Rykindo 37.5 MG SRER	IM	
Rykindo 50 MG SRER	IM	

MEDICATION NAME	ROUTE	DOSE FORM
Rylaze 10 MG/0.5ML SOLUTION	IM	SOLUTION
Rystiggo 280 MG/2ML SOLUTION	SC	SOLUTION
SandIMMUNE 100 MG CAP	PO	CAP
SandIMMUNE 100 MG/ML SOLUTION	PO	SOLUTION
SandIMMUNE 25 MG CAP	PO	CAP
SandIMMUNE 50 MG/ML SOLUTION	IV	SOLUTION
Saphnelo 300 MG/2ML SOLUTION	IV	SOLUTION
Sarclisa 100 MG/5ML SOLUTION	IV	SOLUTION
Sarclisa 500 MG/25ML SOLUTION	IV	SOLUTION
Sensipar 30 MG TAB	PO	TAB
Sensipar 60 MG TAB	PO	TAB
Sensipar 90 MG TAB	PO	TAB
Sezaby 100 MG RECON SOLN	IV	RECON SOLN
Sildenafil Citrate 10 MG/12.5ML SOLUTION	IV	SOLUTION
Simponi Aria 50 MG/4ML SOLUTION	IV	SOLUTION
Simulect 10 MG RECON SOLN	IV	RECON SOLN
Simulect 20 MG RECON SOLN	IV	RECON SOLN
Sirolimus 0.5 MG TAB	PO	TAB
Sirolimus 1 MG TAB	PO	TAB
Sirolimus 1 MG/ML SOLUTION	PO	SOLUTION
Sirolimus 2 MG TAB	PO	TAB
Sivextro 200 MG RECON SOLN	IV	RECON SOLN
Skylla 13.5 MG IUD	IU	IUD
SMOFlipid 20 % EMULSION	IV	EMULSION
Sodium Acetate 2 MEQ/ML SOLUTION	IV	SOLUTION
Sodium Acetate 4 MEQ/ML SOLUTION	IV	SOLUTION
Sodium Bicarbonate 4.2 % SOLUTION	IV	SOLUTION
Sodium Bicarbonate 7.5 % SOLUTION	IV	SOLUTION
Sodium Bicarbonate 8.4 % SOLUTION	IV	SOLUTION
Sodium Chloride 4 MEQ/ML SOLUTION	IV	SOLUTION
Sodium Diuril 500 MG RECON SOLN	IV	RECON SOLN
Sodium Edecrin 50 MG RECON SOLN	IV	RECON SOLN
Sodium Phosphate-NaCl 10 MMOL/100ML SOLUTION	IV	SOLUTION
Sodium Phosphate-NaCl 15 MMOL/100ML SOLUTION	IV	SOLUTION
Sodium Phosphate-NaCl 15 MMOL/250ML SOLUTION	IV	SOLUTION
Sodium Phosphate-NaCl 30 MMOL/250ML SOLUTION	IV	SOLUTION
Sodium Phosphates 15 MMOLE/5ML SOLUTION	IV	SOLUTION
Sodium Phosphates 150 MMOLE/50ML SOLUTION	IV	SOLUTION
Sodium Phosphates 45 MMOLE/15ML SOLUTION	IV	SOLUTION
Sodium Phosphates-Dextrose 15 MMOL/100ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Sodium Phosphates-Dextrose 15 MMOL/250ML SOLUTION	IV	SOLUTION
Soliris 300 MG/30ML SOLUTION	IV	SOLUTION
Solu-CORTEF 1000 MG RECON SOLN	IJ	RECON SOLN
Solu-CORTEF 250 MG RECON SOLN	IJ	RECON SOLN
Solu-CORTEF 500 MG RECON SOLN	IJ	RECON SOLN
SOLU-Medrol (PF) 1000 MG RECON SOLN	IJ	RECON SOLN
SOLU-Medrol (PF) 125 MG RECON SOLN	IJ	RECON SOLN
SOLU-Medrol (PF) 40 MG RECON SOLN	IJ	RECON SOLN
SOLU-Medrol (PF) 500 MG RECON SOLN	IJ	RECON SOLN
SOLU-medrol 1000 MG RECON SOLN	IJ	RECON SOLN
SOLU-medrol 2 GM RECON SOLN	IJ	RECON SOLN
SOLU-medrol 500 MG RECON SOLN	IJ	RECON SOLN
Somatuline Depot 120 MG/0.5ML SOLUTION	SC	SOLUTION
Spevigo 450 MG/7.5ML SOLUTION	IV	SOLUTION
Spravato (56 MG Dose) 28 MG/DEVICE SOLN THPK	NA	SOLN THPK
Spravato (84 MG Dose) 28 MG/DEVICE SOLN THPK	NA	SOLN THPK
Stimufend 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Sublocade 100 MG/0.5ML SOLN PRSYR	SC	SOLN PRSYR
Sublocade 300 MG/1.5ML SOLN PRSYR	SC	SOLN PRSYR
Sunlenca 463.5 MG/1.5ML SOLUTION	SC	SOLUTION
Sustol 10 MG/0.4ML PRSYR	SC	PRSYR
Susvimo (Implant 1st Fill) 10 MG/0.1ML SOLUTION	IZ	SOLUTION
Susvimo (Implant Refill) 10 MG/0.1ML SOLUTION	IZ	SOLUTION
Syfovre 15 MG/0.1ML SOLUTION	IZ	SOLUTION
Sylvant 100 MG RECON SOLN	IV	RECON SOLN
Sylvant 400 MG RECON SOLN	IV	RECON SOLN
Synagis 100 MG/ML SOLUTION	IM	SOLUTION
Synagis 50 MG/0.5ML SOLUTION	IM	SOLUTION
Syndros 5 MG/ML SOLUTION	PO	SOLUTION
Synribo 3.5 MG RECON SOLN	SC	RECON SOLN
Synthamin 17 10 % SOLUTION	IV	SOLUTION
Tacrolimus 0.5 MG CAP	PO	CAP
Tacrolimus 1 MG CAP	PO	CAP
Tacrolimus 5 MG CAP	PO	CAP
Talvey 3 MG/1.5ML SOLUTION	SC	SOLUTION
Talvey 40 MG/ML SOLUTION	SC	SOLUTION
Taxotere 20 MG/ML CONC	IV	CONC
Taxotere 80 MG/4ML CONC	IV	CONC
Tazicef 1 GM/50ML SOLUTION	IV	SOLUTION
Tecentriq 1200 MG/20ML SOLUTION	IV	SOLUTION
Tecentriq 840 MG/14ML SOLUTION	IV	SOLUTION
Temodar 100 MG RECON SOLN	IV	RECON SOLN
Temsirolimus 25 MG/ML SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Teniposide 10 MG/ML SOLUTION	IV	SOLUTION
Tepadina 100 MG RECON SOLN	IJ	RECON SOLN
Tepadina 15 MG RECON SOLN	IJ	RECON SOLN
Tepezza 500 MG RECON SOLN	IV	RECON SOLN
Tezspire 210 MG/1.91ML SOLN A-INJ	SC	SOLN A-INJ
Tezspire 210 MG/1.91ML SOLN PRSYR	SC	SOLN PRSYR
Theophylline in D5W 0.8-5 MG/ML-% SOLUTION	IV	SOLUTION
Thiotepa 100 MG RECON SOLN	IJ	RECON SOLN
Thiotepa 15 MG RECON SOLN	IJ	RECON SOLN
Thymoglobulin 25 MG RECON SOLN	IV	RECON SOLN
Tice BCG 50 MG RECON SUSP	IS	RECON SUSP
Tivdak 40 MG RECON SOLN	IV	RECON SOLN
Tobi 300 MG/5ML NEBU SOLN	IN	NEBU SOLN
Tobramycin 300 MG/5ML NEBU SOLN	IN	NEBU SOLN
Toposar 1 GM/50ML SOLUTION	IV	SOLUTION
Toposar 100 MG/5ML SOLUTION	IV	SOLUTION
Toposar 500 MG/25ML SOLUTION	IV	SOLUTION
Topotecan HCl 4 MG RECON SOLN	IV	RECON SOLN
Topotecan HCl 4 MG/4ML SOLUTION	IV	SOLUTION
Torisel 25 MG/ML SOLUTION	IV	SOLUTION
Totect 500 MG RECON SOLN	IV	RECON SOLN
TPN Electrolytes CONC	IV	CONC
Tranexamic Acid 1000 MG/10ML SOLUTION	IV	SOLUTION
Travasol 10 % SOLUTION	IV	SOLUTION
Trazimera 150 MG RECON SOLN	IV	RECON SOLN
Trazimera 420 MG RECON SOLN	IV	RECON SOLN
Treanda 100 MG RECON SOLN	IV	RECON SOLN
Treanda 25 MG RECON SOLN	IV	RECON SOLN
Trelstar Mixject 11.25 MG RECON SUSP	IM	RECON SUSP
Trelstar Mixject 22.5 MG RECON SUSP	IM	RECON SUSP
Trelstar Mixject 3.75 MG RECON SUSP	IM	RECON SUSP
Treprostinil 100 MG/20ML SOLUTION	IJ	SOLUTION
Treprostinil 20 MG/20ML SOLUTION	IJ	SOLUTION
Treprostinil 200 MG/20ML SOLUTION	IJ	SOLUTION
Treprostinil 50 MG/20ML SOLUTION	IJ	SOLUTION
Triamcinolone Acetonide 40 MG/ML SUSPENSION	IJ	SUSPENSION
Triamcinolone Acetonide 50 MG/ML SUSPENSION	IJ	SUSPENSION
Triesence 40 MG/ML SUSPENSION	IO	SUSPENSION
Triostat 10 MCG/ML SOLUTION	IV	SOLUTION
Triptodur 22.5 MG SRER	IM	
Trisenox 12 MG/6ML SOLUTION	IV	SOLUTION
Trodelvy 180 MG RECON SOLN	IV	RECON SOLN
TrophAmine 10 % SOLUTION	IV	SOLUTION
Trophamine 6 % SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Truxima 100 MG/10ML SOLUTION	IV	SOLUTION
Truxima 500 MG/50ML SOLUTION	IV	SOLUTION
Twinrix 720-20 ELU-MCG/ML SUSP PRSYR	IM	SUSP PRSYR
Tyvaso 0.6 MG/ML SOLUTION	IN	SOLUTION
Tyvaso Refill 0.6 MG/ML SOLUTION	IN	SOLUTION
Tyvaso Starter 0.6 MG/ML SOLUTION	IN	SOLUTION
Udenyca 6 MG/0.6ML SOLN A-INJ	SC	SOLN A-INJ
Udenyca 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Ultomiris 1100 MG/11ML SOLUTION	IV	SOLUTION
Ultomiris 300 MG/30ML SOLUTION	IV	SOLUTION
Ultomiris 300 MG/3ML SOLUTION	IV	SOLUTION
Unituxin 17.5 MG/5ML SOLUTION	IV	SOLUTION
Uplizna 100 MG/10ML SOLUTION	IV	SOLUTION
Uptravi 1800 MCG RECON SOLN	IV	RECON SOLN
Uvadex 20 MCG/ML SOLUTION	EC	SOLUTION
Uzedy 100 MG/0.28ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 125 MG/0.35ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 150 MG/0.42ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 200 MG/0.56ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 250 MG/0.7ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 50 MG/0.14ML SUSP PRSYR	SC	SUSP PRSYR
Uzedy 75 MG/0.21ML SUSP PRSYR	SC	SUSP PRSYR
Vabomere 2 (1-1) GM RECON SOLN	IV	RECON SOLN
Vabysmo 6 MG/0.05ML SOLUTION	IZ	SOLUTION
Valrubicin 40 MG/ML SOLUTION	IS	SOLUTION
Valstar 40 MG/ML SOLUTION	IS	SOLUTION
Vancomycin HCl 1000 MG/10ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1000 MG/200ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1250 MG/12.5ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1250 MG/250ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1500 MG/15ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1500 MG/300ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1750 MG/17.5ML SOLUTION	IV	SOLUTION
Vancomycin HCl 1750 MG/350ML SOLUTION	IV	SOLUTION
Vancomycin HCl 2000 MG/20ML SOLUTION	IV	SOLUTION
Vancomycin HCl 2000 MG/400ML SOLUTION	IV	SOLUTION
Vancomycin HCl 5 GM RECON SOLN	IV	RECON SOLN
Vancomycin HCl 500 MG/100ML SOLUTION	IV	SOLUTION
Vancomycin HCl 750 MG/150ML SOLUTION	IV	SOLUTION
Vancomycin HCl 750 MG/7.5ML SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1-5 GM/100ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1-5 GM/200ML-% SOLUTION	IV	SOLUTION

MEDICATION NAME	ROUTE	DOSE FORM
Vancomycin HCl in Dextrose 1-5 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1.25-5 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1.5-5 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1.5-5 GM/300ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1.5-5 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 1.75-5 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 2-5 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 500-5 MG/100ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in Dextrose 750-5 MG/150ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1-0.9 GM/200ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.25-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.5-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.5-0.9 GM/300ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.5-0.9 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.75-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.75-0.9 GM/300ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 1.75-0.9 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 2-0.9 GM/250ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 2-0.9 GM/500ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 500-0.9 MG/100ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 750-0.9 MG/150ML-% SOLUTION	IV	SOLUTION
Vancomycin HCl in NaCl 750-0.9 MG/250ML-% SOLUTION	IV	SOLUTION
Vantas 50 MG KIT	SC	KIT
Varubi (180 MG Dose) 2 x 90 MG TAB THPK	PO	TAB THPK
Vasopressin 20 UNIT/ML SOLUTION	IV	SOLUTION
Vasostrict 0.2 UNIT/ML SOLUTION	IV	SOLUTION



MEDICATION NAME	ROUTE	DOSE FORM
Vasostrict 0.4 UNIT/ML SOLUTION	IV	SOLUTION
Vasostrict 20 UNIT/ML SOLUTION	IV	SOLUTION
Vectibix 100 MG/5ML SOLUTION	IV	SOLUTION
Vectibix 400 MG/20ML SOLUTION	IV	SOLUTION
Vegzelma 100 MG/4ML SOLUTION	IV	SOLUTION
Vegzelma 400 MG/16ML SOLUTION	IV	SOLUTION
Velcade 3.5 MG RECON SOLN	IJ	RECON SOLN
Veletri 0.5 MG RECON SOLN	IV	RECON SOLN
Veletri 1.5 MG RECON SOLN	IV	RECON SOLN
Ventavis 10 MCG/ML SOLUTION	IN	SOLUTION
Ventavis 20 MCG/ML SOLUTION	IN	SOLUTION
Veopoz 400 MG/2ML SOLUTION	IJ	SOLUTION
Verapamil HCl 2.5 MG/ML SOLUTION	IV	SOLUTION
Vfend IV 200 MG RECON SOLN	IV	RECON SOLN
Vibativ 750 MG RECON SOLN	IV	RECON SOLN
Vidaza 100 MG RECON SUSP	IJ	RECON SUSP
Vimizim 5 MG/5ML SOLUTION	IV	SOLUTION
Vimpat 200 MG/20ML SOLUTION	IV	SOLUTION
VinBLAStine Sulfate 1 MG/ML SOLUTION	IV	SOLUTION
Vincasar PFS 1 MG/ML SOLUTION	IV	SOLUTION
VinCRISStine Sulfate 1 MG/ML SOLUTION	IV	SOLUTION
Vinorelbine Tartrate 10 MG/ML SOLUTION	IV	SOLUTION
Vinorelbine Tartrate 50 MG/5ML SOLUTION	IV	SOLUTION
Virazole 6 GM RECON SOLN	IN	RECON SOLN
Vivimusta 100 MG/4ML SOLUTION	IV	SOLUTION
Voriconazole 200 MG RECON SOLN	IV	RECON SOLN
Vpriv 400 UNIT RECON SOLN	IV	RECON SOLN
Vyepti 100 MG/ML SOLUTION	IV	SOLUTION
Vyvgart 400 MG/20ML SOLUTION	IV	SOLUTION
Vyvgart Hytrulo 180-2000 MG-UNIT/ML SOLUTION	SC	SOLUTION
Vyxeos 44-100 MG RECON SUSP	IV	RECON SUSP
WinRho SDF 1500 UNIT/1.3ML SOLUTION	IJ	SOLUTION
WinRho SDF 15000 UNIT/13ML SOLUTION	IJ	SOLUTION
WinRho SDF 2500 UNIT/2.2ML SOLUTION	IJ	SOLUTION
WinRho SDF 5000 UNIT/4.4ML SOLUTION	IJ	SOLUTION
Xembify 1 GM/5ML SOLUTION	SC	SOLUTION
Xembify 10 GM/50ML SOLUTION	SC	SOLUTION
Xembify 2 GM/10ML SOLUTION	SC	SOLUTION
Xembify 4 GM/20ML SOLUTION	SC	SOLUTION
Xenleta 150 MG/15ML SOLUTION	IV	SOLUTION
Xenpozyme 20 MG RECON SOLN	IV	RECON SOLN
Xenpozyme 4 MG RECON SOLN	IV	RECON SOLN
Xeomin 100 UNIT RECON SOLN	IM	RECON SOLN
Xeomin 200 UNIT RECON SOLN	IM	RECON SOLN

MEDICATION NAME	ROUTE	DOSE FORM
Xeomin 50 UNIT RECON SOLN	IM	RECON SOLN
Xerava 100 MG RECON SOLN	IV	RECON SOLN
Xerava 50 MG RECON SOLN	IV	RECON SOLN
Xipere 40 MG/ML SUSPENSION	IO	SUSPENSION
Xopenex 0.31 MG/3ML NEBU SOLN	IN	NEBU SOLN
Xopenex 0.63 MG/3ML NEBU SOLN	IN	NEBU SOLN
Xopenex 1.25 MG/3ML NEBU SOLN	IN	NEBU SOLN
Xopenex Concentrate 1.25 MG/0.5ML NEBU SOLN	IN	NEBU SOLN
Xylocaine Dental 2 %-1:100000 SOLUTION	IJ	SOLUTION
Xylocaine Dental 2 %-1:50000 SOLUTION	IJ	SOLUTION
Xylocaine-MPF/Epinephrine 1 %-1:200000 SOLUTION	IJ	SOLUTION
Xylocaine-MPF/Epinephrine 1.5 %-1:200000 SOLUTION	IJ	SOLUTION
Xylocaine-MPF/Epinephrine 2 %-1:200000 SOLUTION	IJ	SOLUTION
Xylocaine/Epinephrine 0.5 %-1:200000 SOLUTION	IJ	SOLUTION
Xylocaine/Epinephrine 1 %-1:100000 SOLUTION	IJ	SOLUTION
Xylocaine/Epinephrine 2 %-1:100000 SOLUTION	IJ	SOLUTION
Yervoy 200 MG/40ML SOLUTION	IV	SOLUTION
Yervoy 50 MG/10ML SOLUTION	IV	SOLUTION
Yondelis 1 MG RECON SOLN	IV	RECON SOLN
Yupelri 175 MCG/3ML SOLUTION	IN	SOLUTION
Zaltrap 100 MG/4ML SOLUTION	IV	SOLUTION
Zaltrap 200 MG/8ML SOLUTION	IV	SOLUTION
Zanosar 1 GM RECON SOLN	IV	RECON SOLN
Zemaira 1000 MG RECON SOLN	IV	RECON SOLN
Zemdri 500 MG/10ML SOLUTION	IV	SOLUTION
Zemplar 1 MCG CAP	PO	CAP
Zemplar 2 MCG CAP	PO	CAP
Zemplar 2 MCG/ML SOLUTION	IV	SOLUTION
Zemplar 5 MCG/ML SOLUTION	IV	SOLUTION
Zepzelca 4 MG RECON SOLN	IV	RECON SOLN
Zerbaxa 1.5 (1-0.5) GM RECON SOLN	IV	RECON SOLN
Zevalin Y-90 3.2 MG/2ML KIT	IV	KIT
Ziextenzo 6 MG/0.6ML SOLN PRSYR	SC	SOLN PRSYR
Zilretta 32 MG SRER	IX	
Zinecard 250 MG RECON SOLN	IV	RECON SOLN
Zinecard 500 MG RECON SOLN	IV	RECON SOLN
Zinplava 1000 MG/40ML SOLUTION	IV	SOLUTION
Zirabev 100 MG/4ML SOLUTION	IV	SOLUTION
Zirabev 400 MG/16ML SOLUTION	IV	SOLUTION
Zofran 4 MG TAB	PO	TAB
Zofran 8 MG TAB	PO	TAB



MEDICATION NAME	ROUTE	DOSE FORM
Zoladex 10.8 MG IMPLANT	SC	IMPLANT
Zoladex 3.6 MG IMPLANT	SC	IMPLANT
Zoledronic Acid 4 MG/100ML SOLUTION	IV	SOLUTION
Zoledronic Acid 4 MG/5ML CONC	IV	CONC
Zoledronic Acid 5 MG/100ML SOLUTION	IV	SOLUTION
Zortress 0.25 MG TAB	PO	TAB
Zortress 0.5 MG TAB	PO	TAB
Zortress 0.75 MG TAB	PO	TAB
Zortress 1 MG TAB	PO	TAB
Zulresso 100 MG/20ML SOLUTION	IV	SOLUTION
Zuplenz 4 MG FILM	PO	FILM
Zuplenz 8 MG FILM	PO	FILM
Zynlonta 10 MG RECON SOLN	IV	RECON SOLN
Zynyz 500 MG/20ML SOLUTION	IV	SOLUTION