

## **PA Criteria**

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| <b>Prior Authorization Group</b>    | ABIRATERONE  |
| <b>Drug Names</b>                   | ABIRATERONE ACETATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Node-positive (N1), non-metastatic (M0) prostate cancer  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | ACITRETIN  |
| <b>Drug Names</b>                   | ACITRETIN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Prevention of non-melanoma skin cancers in high risk individuals, Lichen planus, Keratosis follicularis (Darier Disease)                                     |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Psoriasis: The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to methotrexate or cyclosporine. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | ACTIMMUNE   |
| <b>Drug Names</b>                   | ACTIMMUNE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications |
| <b>Off-label Uses</b>               | Mycosis fungoides, Sezary syndrome.                               |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | ADEMPAS  |
| <b>Drug Names</b>                   | ADEMPAS  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1): PAH was confirmed by right heart catheterization. For PAH new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, AND 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND 3) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. For chronic thromboembolic pulmonary hypertension (CTEPH) (WHO Group 4): 1) Patient has persistent or recurrent CTEPH after pulmonary endarterectomy (PEA), OR 2) Patient has inoperable CTEPH with the diagnosis confirmed by right heart catheterization AND by computed tomography (CT), magnetic resonance imaging (MRI), or pulmonary angiography. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | AIMOVIG  |
| <b>Drug Names</b>                   | AIMOVIG  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For the preventive treatment of migraine, initial: 1) The patient experienced an inadequate treatment response with a 4-week trial of any one of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants OR 2) The patient experienced an intolerance or has a contraindication that would prohibit a 4-week trial of any one of the following: Antiepileptic drugs (AEDs), Beta-adrenergic blocking agents, Antidepressants. For preventive treatment of migraine, continuation: The patient received at least 3 months of treatment with the requested drug, and the patient had a reduction in migraine days per month from baseline.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Initial: 3 months, Continuation: Plan Year   |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ALDURAZYME  |
| <b>Drug Names</b>                   | ALDURAZYME  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For mucopolysaccharidosis I (MPS I): Diagnosis of MPS I was confirmed by an enzyme assay demonstrating a deficiency of alpha-L-iduronidase enzyme activity and/or by genetic testing. Patients with Scheie form (i.e., attenuated MPS I) must have moderate to severe symptoms.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | ALECENSA  |
| <b>Drug Names</b>                   | ALECENSA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent ALK-positive non-small cell lung cancer (NSCLC), brain metastases from ALK-positive NSCLC.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | ALOSETRON   |
| <b>Drug Names</b>                   | ALOSETRON HYDROCHLORIDE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For severe diarrhea-predominant irritable bowel syndrome (IBS): 1) The requested drug is being prescribed for a biological female or a person that self-identifies as a female, 2) chronic IBS symptoms lasting at least 6 months, 3) gastrointestinal tract abnormalities have been ruled out, AND 4) inadequate response to one conventional therapy (e.g., antispasmodics, antidepressants, antidiarrheals). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ALPHA1-PROTEINASE INHIBITOR  |
| <b>Drug Names</b>                   | ARALAST NP, PROLASTIN-C, ZEMAIRA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For alpha1-proteinase inhibitor deficiency: Patient must have 1) clinically evident emphysema and 2) pretreatment serum alpha1-proteinase inhibitor level less than 11 micromol/L (80 mg/dL by radial immunodiffusion or 50 mg/dL by nephelometry).  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ALUNBRIG   |
| <b>Drug Names</b>                   | ALUNBRIG   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent ALK-positive non-small cell lung cancer (NSCLC), brain metastases from ALK-positive NSCLC, Inflammatory myofibroblastic tumor (IMT) with ALK translocation.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | AMBRISANTAN  |
| <b>Drug Names</b>                   | AMBRISANTAN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1): Diagnosis was confirmed by right heart catheterization. For PAH new starts only: 1) Pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) Pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) Pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | AMPHETAMINES  |
| <b>Drug Names</b>                   | AMPHETAMINE/DEXTROAMPHETA   |
| <b>PA Indication Indicator</b>      | All Medically-accepted Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The patient has a diagnosis of Attention-Deficit Hyperactivity Disorder (ADHD) or Attention Deficit Disorder (ADD) OR 2) The patient has a diagnosis of narcolepsy confirmed by a sleep study.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | ARCALYST  |
| <b>Drug Names</b>                   | ARCALYST  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Prevention of gout flares in patients initiating or continuing urate-lowering therapy.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For prevention of gout flares in patients initiating or continuing urate-lowering therapy (e.g., allopurinol) (new starts): 1) two or more gout flares within the previous 12 months, AND 2) inadequate response, intolerance or contraindication to maximum tolerated doses of a non-steroidal anti-inflammatory drug (NSAID) and colchicine, AND 3) concurrent use with urate-lowering therapy. For prevention of gout flares in patients initiating or continuing urate-lowering therapy (e.g., allopurinol) (continuation): 1) patient must have achieved or maintained a clinical benefit (i.e., a fewer number of gout attacks or fewer flare days) compared to baseline, AND 2) continued use of urate-lowering therapy concurrently with the requested drug. For recurrent pericarditis: patient must have had an inadequate response, intolerance or contraindication to maximum tolerated doses of an NSAID and colchicine. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | For prevention of gout flares: 4 months. Other: Plan Year   |
| <b>Other Criteria</b>               | -   |

**Prior Authorization Group** ARMODAFINIL  
**Drug Names** ARMODAFINIL  
**PA Indication Indicator** All FDA-approved Indications  
**Off-label Uses** -  
**Exclusion Criteria** -  
**Required Medical Information** 1) The patient has a diagnosis of narcolepsy and the diagnosis is confirmed by sleep lab evaluation OR 2) The patient has a diagnosis of Shift Work Disorder (SWD) OR 3) The patient has a diagnosis of obstructive sleep apnea (OSA) and the diagnosis is confirmed by polysomnography.

**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

**Prior Authorization Group** AUSTEDO  
**Drug Names** AUSTEDO  
**PA Indication Indicator** All FDA-approved Indications, Some Medically-accepted Indications  
**Off-label Uses** Tourette's syndrome  
**Exclusion Criteria** -  
**Required Medical Information** -

**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

**Prior Authorization Group** AUVELITY  
**Drug Names** AUVELITY  
**PA Indication Indicator** All FDA-approved Indications  
**Off-label Uses** -  
**Exclusion Criteria** -  
**Required Medical Information** For Major Depressive Disorder (MDD): The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to two of the following: serotonin and norepinephrine reuptake inhibitors (SNRIs), selective serotonin reuptake inhibitors (SSRIs), mirtazapine, bupropion.

**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

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| <b>Prior Authorization Group</b>    | AYVAKIT   |
| <b>Drug Names</b>                   | AYVAKIT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Myeloid and lymphoid neoplasms with eosinophilia, gastrointestinal stromal tumor (GIST) for unresectable, recurrent, or metastatic disease without platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For myeloid and lymphoid neoplasms with eosinophilia, the patient meets all of the following criteria: 1) The disease is FIP1L1- PDGFRA rearrangement-positive, AND 2) The disease harbors a PDGFRA D842A mutation, AND 3) The disease is resistant to imatinib. For GIST, the patient meets either of the following criteria: 1) The disease harbors PDGFRA exon 18 mutation, including PDGFRA D842V mutations, OR 2) The requested drug will be used after failure on at least two Food and Drug Administration (FDA)-approved therapies in unresectable, recurrent, or metastatic disease without PDGFRA exon 18 mutation. For advanced systemic mastocytosis (AdvSM): 1) The patient has a diagnosis of advanced systemic mastocytosis including aggressive systemic mastocytosis (ASM), systemic mastocytosis with associated hematological neoplasm (SM-AHN), and mast cell leukemia (MCL) AND 2) The patient has a platelet count of greater than or equal to 50,000/microliter (mcL). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

**Prior Authorization Group**

**Drug Names**

B VS. D

ABELCET, ACETYLCYSTEINE, ACYCLOVIR SODIUM, ALBUTEROL SULFATE, AMPHOTERICIN B, AMPHOTERICIN B LIPOSOME, APREPITANT, AZACITIDINE, AZATHIOPRINE, BENDEKA, BUDESONIDE, CALCITONIN-SALMON, CALCITRIOL, CARBOPLATIN, CINACALCET HYDROCHLORIDE, CISPLATIN, CLINIMIX 4.25%/DEXTROSE 1, CLINIMIX 4.25%/DEXTROSE 5, CLINIMIX 5%/DEXTROSE 15%, CLINIMIX 5%/DEXTROSE 20%, CLINIMIX 6/5, CLINIMIX 8/10, CLINIMIX 8/14, CLINISOL SF 15%, CLINOLIPID, CROMOLYN SODIUM, CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE MONOHYDR, CYCLOSPORINE, CYCLOSPORINE MODIFIED, CYTARABINE AQUEOUS, DEXTROSE 50%, DEXTROSE 70%, DIPHTHERIA/TETANUS TOXOID, DOCETAXEL, DOXORUBICIN HCL, DOXORUBICIN HYDROCHLORIDE, DRONABINOL, ELLENCE, ENGERIX-B, ETOPOSIDE, EVEROLIMUS, FLUOROURACIL, FREAMINE III, FULVESTRANT, GAMASTAN, GANCICLOVIR, GEMCITABINE HCL, GEMCITABINE HYDROCHLORIDE, GENGRAF, GRANISETRON HYDROCHLORIDE, HEPARIN SODIUM, HUMULIN R U-500 (CONCENTR, IBANDRONATE SODIUM, IMOVAX RABIES (H.D.C.V.), INTRALIPID, INTRON A, IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE/ALBUT, IRINOTECAN, IRINOTECAN HYDROCHLORIDE, KADCYLA, LEUCOVORIN CALCIUM, LEVALBUTEROL, LEVALBUTEROL HCL, LEVOCARNITINE, LIDOCAINE HCL, LIDOCAINE HYDROCHLORIDE, METHOTREXATE, METHOTREXATE SODIUM, METHYLPREDNISOLONE, METHYLPREDNISOLONE ACETAT, METHYLPREDNISOLONE SODIUM, MORPHINE SULFATE, MORPHINE SULFATE/SODIUM C, MYCOPHENOLATE MOFETIL, MYCOPHENOLIC ACID DR, NULOJIX, NUTRILIPID, ONDANSETRON HCL, ONDANSETRON HYDROCHLORIDE, ONDANSETRON ODT, OXALIPLATIN, PACLITAXEL, PACLITAXEL PROTEIN-BOUND, PAMIDRONATE DISODIUM, PARAPLATIN, PARICALCITOL, PEMETREXED, PENTAMIDINE ISETHIONATE, PLENAMINE, PREDNISOLONE, PREDNISOLONE SODIUM PHOSP, PREDNISONE, PREDNISONE INTENSOL, PREHEVBRIO, PREMASOL, PROCALAMINE, PROGRAF, PROSOL, RABAVERT, RECOMBIVAX HB, SANDIMMUNE, SIROLIMUS, TACROLIMUS, TDVAX, TENIVAC, TOPOSAR, TPN ELECTROLYTES, TRAVASOL, TROPHAMINE, VINCRISTINE SULFATE, VINOELBINE TARTRATE, XATMEP, ZOLEDRONIC ACID

**PA Indication Indicator**

All Medically-accepted Indications

**Off-label Uses**

-

**Exclusion Criteria**

-

**Required Medical Information**

-

**Age Restrictions**

-

**Prescriber Restrictions**

-

**Coverage Duration**

N/A

**Other Criteria**

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



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| <b>Prior Authorization Group</b>    | BAFIERTAM   |
| <b>Drug Names</b>                   | BAFIERTAM   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | BALVERSA  |
| <b>Drug Names</b>                   | BALVERSA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent primary carcinoma of the urethra, recurrent or persistent urothelial carcinoma of the bladder.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For urothelial carcinoma: Disease has susceptible fibroblast growth factor receptor 3 (FGFR3) or fibroblast growth factor receptor 2 (FGFR2) genetic alterations AND the requested drug will be used as subsequent therapy for any of the following: a) locally advanced or metastatic urothelial carcinoma, b) recurrent primary carcinoma of the urethra, c) stage II urothelial carcinoma of the bladder if tumor is present following reassessment of tumor status 2-3 months after primary treatment with bladder preserving concurrent chemoradiotherapy, d) urothelial carcinoma of the bladder with metastatic or local recurrence post cystectomy, or e) urothelial carcinoma of the bladder with muscle invasive local recurrence or persistent disease in a preserved bladder. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | BANZEL  |
| <b>Drug Names</b>                   | RUFINAMIDE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | 1 year of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

|                                     |  |
|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | BENLYSTA   |
| <b>Drug Names</b>                   | BENLYSTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | For patients new to therapy: severe active central nervous system lupus.   |
| <b>Required Medical Information</b> | The requested drug will not be used in combination with other biologic therapies. For systemic lupus erythematosus (SLE): 1) Patient is currently receiving a stable standard therapy regimen (e.g., corticosteroid or antimalarial) for SLE OR 2) patient is not currently receiving stable standard therapy regimen for SLE because the patient experienced an intolerance or has a contraindication to standard therapy regimens. For lupus nephritis: 1) Patient is currently receiving a stable standard therapy regimen (e.g., corticosteroid) for lupus nephritis OR 2) patient is not currently receiving a stable standard therapy regimen for lupus nephritis because the patient experienced an intolerance or has a contraindication to standard therapy regimens. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | BERINERT   |
| <b>Drug Names</b>                   | BERINERT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Short-term preprocedural prophylaxis for hereditary angioedema (HAE) attacks   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hereditary angioedema (HAE): The requested drug is being used for the treatment of acute angioedema attacks. Patient has HAE with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has HAE with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, EITHER 1) Patient tested positive for an F12, angiotensin-converting enzyme 1 (ACE1), plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosaminase 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation OR 2) Patient has a family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy for at least one month.   |
| <b>Age Restrictions</b>             | 5 years of age or older  |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an immunologist, allergist, or rheumatologist  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|------------------------------|
| <b>Prior Authorization Group</b>    | BESREMI                      |
| <b>Drug Names</b>                   | BESREMI                      |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

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|-------------------------------------|------------------------------|
| <b>Prior Authorization Group</b>    | BETASERON                    |
| <b>Drug Names</b>                   | BETASERON                    |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | BEXAROTENE  |
| <b>Drug Names</b>                   | BEXAROTENE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Mycosis fungoides, Sezary syndrome, CD30-positive primary cutaneous anaplastic large cell lymphoma, CD30-positive lymphomatoid papulosis. |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | BOSENTAN   |
| <b>Drug Names</b>                   | BOSENTAN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1): Diagnosis was confirmed by right heart catheterization. For PAH new starts only: 1) Pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) Pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) Pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | BOSULIF  |
| <b>Drug Names</b>                   | BOSULIF  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in the chronic phase or blast phase  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML) or acute lymphoblastic leukemia (ALL), including patients who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene, and 2) If patient experienced resistance to an alternative tyrosine kinase inhibitor, patient is negative for all of the following mutations: T315I, G250E, V299L, and F317L. For CML, including patients newly diagnosed with CML and patients who have received a hematopoietic stem cell transplant: patient has experienced resistance or intolerance to imatinib or dasatinib. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | BRAFTOVI   |
| <b>Drug Names</b>                   | BRAFTOVI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Adjuvant systemic therapy for cutaneous melanoma   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For colorectal cancer: The patient must meet both of the following criteria: 1) Tumor is positive for BRAF V600E mutation, 2) The requested drug will be used for either of the following: a) as subsequent therapy for advanced or metastatic disease, or b) as primary treatment for unresectable metachronous metastases. For cutaneous melanoma: The patient must meet all of the following criteria: 1) Tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), 2) The requested drug will be used in combination with binimetinib, and 3) The requested drug will be used for either of the following: a) unresectable or metastatic disease, or b) adjuvant systemic therapy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | BRIVIACT   |
| <b>Drug Names</b>                   | BRIVIACT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | 1) The patient has experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) The patient has experienced an inadequate treatment response, intolerance, or contraindication to any of the following: Aptiom, Vimpat, Xcopri, Spritam.  |
| <b>Age Restrictions</b>             | 1 month of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | BRIVIACT INJ   |
| <b>Drug Names</b>                   | BRIVIACT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | 1) The patient has experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) The patient has experienced an inadequate treatment response, intolerance, or contraindication to any of the following: Aptiom, Vimpat, Xcopri, Spritam.  |
| <b>Age Restrictions</b>             | 1 month of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | BRUKINSA   |
| <b>Drug Names</b>                   | BRUKINSA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Chronic lymphocytic leukemia (CLL), small lymphocytic lymphoma (SLL)   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For marginal zone lymphoma: 1) the requested drug is being used for the treatment of relapsed or refractory disease AND the patient has received at least one anti-CD20-based regimen OR 2) the requested drug is being used for the treatment of refractory or progressive disease AND the patient has experienced an intolerance or has a contraindication to ibrutinib. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | BUDESONIDE CAP   |
| <b>Drug Names</b>                   | BUDESONIDE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Treatment and maintenance of microscopic colitis in adults   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For the maintenance of microscopic colitis: patient has had a clinical relapse after cessation of treatment (induction) therapy.   |
| <b>Age Restrictions</b>             | Crohn's, treatment: 8 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Microscopic colitis, maintenance: 12 months, all other indications: 3 months   |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | BUPRENORPHINE  |
| <b>Drug Names</b>                   | BUPRENORPHINE HCL  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug is being prescribed for the treatment of opioid use disorder AND patient meets one of the following: 1) The patient is pregnant or breastfeeding, and the requested drug is being prescribed for induction therapy and/or subsequent maintenance therapy for treatment of opioid use disorder OR 2) The requested drug is being prescribed for induction therapy for transition from opioid use to treatment of opioid use disorder OR 3) The requested drug is being prescribed for maintenance therapy for treatment of opioid use disorder in a patient who is intolerant to naloxone.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | CABOMETYX  |
| <b>Drug Names</b>                   | CABOMETYX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Non-small cell lung cancer, Ewing sarcoma, osteosarcoma, gastrointestinal stromal tumor  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For renal cell carcinoma: The disease is advanced, relapsed, or stage IV. For non-small cell lung cancer: 1) the disease is rearranged during transfection (RET) positive AND 2) the disease is recurrent, advanced, or metastatic. For hepatocellular carcinoma: the requested drug will be used as subsequent treatment. For gastrointestinal stromal tumor (GIST): the disease is unresectable, recurrent, or metastatic AND the patient has failed on an FDA-approved therapy (e.g., imatinib, sunitinib, regorafenib, ripretinib). For Ewing sarcoma and osteosarcoma: the requested drug will be used as subsequent therapy. For differentiated thyroid cancer (DTC) (follicular, papillary, Hurthle cell): 1) The disease is locally advanced or metastatic disease, 2) the disease has progressed after a vascular endothelial growth factor receptor (VEGFR)- targeted therapy, AND 3) the patient is refractory to radioactive iodine therapy (RAI) or ineligible for RAI. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | CALCIPOTRIENE   |
| <b>Drug Names</b>                   | CALCIPOTRIENE, CALCITRENE, ENSTILAR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For Treatment of Psoriasis: The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to a topical steroid.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | CALQUENCE   |
| <b>Drug Names</b>                   | CALQUENCE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Waldenstrom macroglobulinemia, lymphoplasmacytic lymphoma, gastric mucosa-associated lymphoid tissue (MALT) lymphoma, non-gastric MALT lymphoma (noncutaneous), nodal marginal zone lymphoma, splenic marginal zone lymphoma  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For gastric MALT lymphoma, non-gastric MALT lymphoma, nodal marginal zone lymphoma, and splenic marginal zone lymphoma: 1) the requested drug is being used for the treatment of refractory or progressive disease AND 2) the patient has experienced an intolerance or has a contraindication to ibrutinib. For chronic lymphocytic leukemia or small lymphocytic lymphoma: the patient has experienced an intolerance or has a contraindication to zanubrutinib or ibrutinib. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |



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| <b>Prior Authorization Group</b>    | CAPLYTA   |
| <b>Drug Names</b>                   | CAPLYTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For treatment of schizophrenia: 1) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following generic products: aripiprazole, asenapine, olanzapine, quetiapine, risperidone, ziprasidone, AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following brand products: Latuda, Rexulti, Secuado, Vraylar. For treatment of depressive episodes associated with bipolar I: 1) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following generic products: olanzapine, quetiapine, AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following brand products: Latuda, Vraylar. For treatment of depressive episodes associated with bipolar II: The patient experienced an inadequate treatment response, intolerance, or contraindication to generic quetiapine. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | CAPRELSA  |
| <b>Drug Names</b>                   | CAPRELSA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Differentiated thyroid carcinoma: papillary, follicular, and Hurthle cell.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | CARBAGLU  |
| <b>Drug Names</b>                   | CARGLUMIC ACID  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For N-acetylglutamate synthase (NAGS) deficiency: Diagnosis of NAGS deficiency was confirmed by enzymatic, biochemical, or genetic testing. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | CAYSTON  |
| <b>Drug Names</b>                   | CAYSTON  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For treatment of respiratory symptoms in cystic fibrosis patients: 1) Pseudomonas aeruginosa is present in the patient's airway cultures OR 2) The patient has a history of pseudomonas aeruginosa infection or colonization in the airways. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | CERDELGA  |
| <b>Drug Names</b>                   | CERDELGA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For type 1 Gaucher disease (GD1): 1) The diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing, and 2) The patient's CYP2D6 metabolizer status has been established using an FDA-cleared test, and 3) The patient is a CYP2D6 extensive metabolizer, an intermediate metabolizer, or a poor metabolizer. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | CEREZYME  |
| <b>Drug Names</b>                   | CEREZYME  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Type 2 Gaucher disease, Type 3 Gaucher disease  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For Gaucher disease, the diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | CLOBAZAM  |
| <b>Drug Names</b>                   | CLOBAZAM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | 2 years of age or older   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | CLOMIPRAMINE  |
| <b>Drug Names</b>                   | CLOMIPRAMINE HYDROCHLORID   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Depression, Panic Disorder  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The requested drug is being prescribed for one of the following: a) Obsessive-Compulsive Disorder (OCD), b) Panic Disorder AND 2) The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to any of the following: a) a serotonin and norepinephrine reuptake inhibitor (SNRI), b) a selective serotonin reuptake inhibitor (SSRI) OR 3) The requested drug is being prescribed for Depression AND 4) The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to two of the following: a) serotonin and norepinephrine reuptake inhibitors (SNRIs), b) selective serotonin reuptake inhibitors (SSRIs), c) mirtazapine, d) bupropion |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | CLORAZEPATE  |
| <b>Drug Names</b>                   | CLORAZEPATE DIPOTASSIUM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For all indications: The prescriber must acknowledge the benefit of therapy with this prescribed medication outweighs the potential risks for the patient. (Note: The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) For the management of anxiety disorders: 1) The requested drug is being used concurrently with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the SSRI/SNRI becomes effective for the symptoms of anxiety, OR 2) The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to AT LEAST TWO agents from the following classes: a) selective serotonin reuptake inhibitors (SSRIs), b) serotonin-norepinephrine reuptake inhibitors (SNRIs). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Short-term relief anxiety-1 month, Anxiety Disorders-4 months, All other Diagnoses-Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 65 years of age or older.  |
| <b>Prior Authorization Group</b>    | CLOZAPINE ODT  |
| <b>Drug Names</b>                   | CLOZAPINE ODT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | COMETRIQ   |
| <b>Drug Names</b>                   | COMETRIQ   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Non-small cell lung cancer (NSCLC), differentiated thyroid carcinoma: papillary, follicular, and Hurthle cell.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For NSCLC: The requested medication is used for NSCLC when the patient's disease expresses rearranged during transfection (RET) gene rearrangements.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | COPIKTRA   |
| <b>Drug Names</b>                   | COPIKTRA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL): the patient has relapsed or refractory disease.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | COTELLIC   |
| <b>Drug Names</b>                   | COTELLIC   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma), Erdheim-Chester disease, Langerhans cell histiocytosis, Rosai-Dorfman disease   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For adjuvant treatment of melanoma, and central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma): The patient must meet both of the following criteria: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used in combination with vemurafenib. For unresectable or metastatic melanoma: The patient must meet both of the following criteria: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used in combination with vemurafenib (with or without atezolizumab). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | CYSTADROPS   |
| <b>Drug Names</b>                   | CYSTADROPS   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient meets both of the following: 1) Diagnosis of cystinosis was confirmed by ANY of the following: a) the presence of increased cystine concentration in leukocytes, OR b) genetic testing, OR c) demonstration of corneal cystine crystals by slit lamp examination, AND 2) the patient has corneal cystine crystal accumulation. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | CYSTAGON   |
| <b>Drug Names</b>                   | CYSTAGON   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Diagnosis of nephropathic cystinosis was confirmed by ANY of the following: 1) the presence of increased cystine concentration in leukocytes, OR 2) genetic testing, OR 3) demonstration of corneal cystine crystals by slit lamp examination.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | CYSTARAN   |
| <b>Drug Names</b>                   | CYSTARAN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient meets both of the following: 1) Diagnosis of cystinosis was confirmed by ANY of the following: a) the presence of increased cystine concentration in leukocytes, OR b) genetic testing, OR c) demonstration of corneal cystine crystals by slit lamp examination, AND 2) the patient has corneal cystine crystal accumulation. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | DALFAMPRIDINE  |
| <b>Drug Names</b>                   | DALFAMPRIDINE ER   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For multiple sclerosis, patient must meet the following: For new starts, prior to initiating therapy, patient demonstrates sustained walking impairment. For continuation of therapy: patient must have experienced an improvement in walking speed OR other objective measure of walking ability since starting the requested drug.       |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | DAURISMO   |
| <b>Drug Names</b>                   | DAURISMO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Post induction therapy following response to previous therapy with the same regimen for acute myeloid leukemia (AML). Relapsed/refractory AML as a component of repeating the initial successful induction regimen.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acute myeloid leukemia: 1) the requested drug must be used in combination with cytarabine, 2) the patient is 75 years of age or older OR has comorbidities that preclude intensive chemotherapy, and 3) the requested drug will be used as treatment for induction therapy, post-induction therapy, or relapsed or refractory disease. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | DEFERASIROX  |
| <b>Drug Names</b>                   | DEFERASIROX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic iron overload due to blood transfusions: pretreatment serum ferritin level is greater than 1000 mcg/L.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | DEMSEER   |
| <b>Drug Names</b>                   | METYROSINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to an alpha-adrenergic antagonist.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | DESVENLAFAXINE  |
| <b>Drug Names</b>                   | DESVENLAFAXINE ER   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to TWO of the following: a) serotonin and norepinephrine reuptake inhibitors (SNRIs), b) selective serotonin reuptake inhibitors (SSRIs), c) mirtazapine, d) bupropion |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | DEXMETHYLPHENIDATE  |
| <b>Drug Names</b>                   | DEXMETHYLPHENIDATE HCL, DEXMETHYLPHENIDATE HYDROC   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Cancer-related fatigue  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The patient has a diagnosis of Attention-Deficit Hyperactivity Disorder (ADHD) or Attention Deficit Disorder (ADD) OR 2) The requested drug is being prescribed for the treatment of cancer-related fatigue after other causes of fatigue have been ruled out.                   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |



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| <b>Prior Authorization Group</b>    | DHE NASAL  |
| <b>Drug Names</b>                   | DIHYDROERGOTAMINE MESYLAT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | Coverage will be denied when used in conjunction with potent CYP3A4 inhibitors (e.g., ritonavir, nelfinavir, indinavir, erythromycin, clarithromycin). |
| <b>Required Medical Information</b> | The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one triptan 5-HT1 receptor agonist.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | DIACOMIT   |
| <b>Drug Names</b>                   | DIACOMIT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | 6 months of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | DIAZEPAM   |
| <b>Drug Names</b>                   | DIAZEPAM, DIAZEPAM INTENSOL  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For all indications: The prescriber must acknowledge the benefit of therapy with this prescribed medication outweighs the potential risks for the patient. (Note: The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) For the management of anxiety disorders: 1) The requested drug is being used concurrently with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the SSRI/SNRI becomes effective for the symptoms of anxiety, OR 2) The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to AT LEAST TWO agents from the following classes: a) selective serotonin reuptake inhibitors (SSRIs), b) serotonin-norepinephrine reuptake inhibitors (SNRIs). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Short-term relief anx-1 mo, skeletal muscle spasm-3 mo, Anx Disorders-4 mo, Other Diagnoses-PlanYR   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 65 years of age or older.  |

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| <b>Prior Authorization Group</b>    | DOPTELET  |
| <b>Drug Names</b>                   | DOPTELET  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For thrombocytopenia in patients with chronic liver disease: Untransfused platelet count prior to a scheduled procedure is less than 50,000/mcL. For chronic immune thrombocytopenia (ITP): 1) For new starts: a) Patient has had an inadequate response or is intolerant to prior therapy such as corticosteroids or immunoglobulins, AND b) Untransfused platelet count at any point prior to the initiation of the requested medication is less than 30,000/mcL OR 30,000 to 50,000/mcL with symptomatic bleeding or risk factor(s) for bleeding (e.g., undergoing a medical or dental procedure where blood loss is anticipated, comorbidities such as peptic ulcer disease and hypertension, anticoagulation therapy, profession or lifestyle that predisposes patient to trauma). 2) For continuation of therapy, platelet count response to the requested drug: a) Current platelet count is less than or equal to 200,000/mcL OR b) Current platelet count is greater than 200,000/mcL and less than or equal to 400,000/mcL and dosing will be adjusted to a platelet count sufficient to avoid clinically important bleeding. |
| <b>Age Restrictions</b>             | 18 years of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Chronic liver disease: 1 month, ITP initial: 6 months, ITP reauthorization: Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | DRIZALMA   |
| <b>Drug Names</b>                   | DRIZALMA SPRINKLE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Cancer pain, chemotherapy-induced neuropathic pain   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | 1) The patient has tried duloxetine capsules OR 2) The patient is unable to take duloxetine capsules for any reason (e.g., difficulty swallowing capsules, requires nasogastric administration). |
| <b>Age Restrictions</b>             | Generalized Anxiety Disorder - 7 years of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | DUPIXENT   |
| <b>Drug Names</b>                   | DUPIXENT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For atopic dermatitis (AD), initial therapy: 1) Patient has moderate-to-severe disease, 2) Patient has had an inadequate treatment response to either a topical corticosteroid or a topical calcineurin inhibitor, OR topical corticosteroids and topical calcineurin inhibitors are not advisable for the patient. For AD, continuation of therapy: the patient achieved or maintained positive clinical response. For moderate-to-severe asthma, initial therapy: Patient meets either of the following: 1) patient is oral corticosteroid dependent and asthma remains inadequately controlled despite current treatment with both of the following medications: a) high-dose inhaled corticosteroid and b) an additional controller (long acting beta2-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline) unless patient has an intolerance or contraindication to such therapies, OR 2) patient has a baseline blood eosinophil count of at least 150 cells per microliter and their asthma remains inadequately controlled despite current treatment with both of the following medications: a) medium-to-high-dose inhaled corticosteroid and b) additional controller (long acting beta2-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline) unless patient has an intolerance or contraindication to such therapies. For moderate-to-severe asthma, continuation of therapy: asthma control has improved on treatment with the requested drug. For chronic rhinosinusitis with nasal polyposis (CRSwNP): 1) the requested drug is used as add-on maintenance treatment, AND 2) the patient has experienced an inadequate treatment response to Xhance (fluticasone). |
| <b>Age Restrictions</b>             | Atopic Dermatitis: 6 months of age or older, Asthma: 6 years of age or older, Chronic Rhinosinusitis with Nasal Polyposis and Prurigo Nodularis: 18 years of age or older, Eosinophilic Esophagitis: 12 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | AD, initial: 4 months, PN, initial: 6 months, All other: Plan Year   |
| <b>Other Criteria</b>               | For eosinophilic esophagitis (EoE), initial therapy: 1) diagnosis has been confirmed by esophageal biopsy, 2) patient weighs at least 40 kilograms, 3) patient experienced an inadequate treatment response, intolerance, or patient has a contraindication to a topical corticosteroid (e.g., fluticasone propionate or budesonide). For EoE, continuation of therapy: the patient achieved or maintained a positive clinical response. For prurigo nodularis (PN), initial therapy: Patient has had an inadequate treatment response to a topical corticosteroid OR topical corticosteroids are not advisable for the patient. For PN, continuation of therapy: The patient achieved or maintained a positive clinical response.   |

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| <b>Prior Authorization Group</b>    | ELIGARD  |
| <b>Drug Names</b>                   | ELIGARD  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent androgen receptor positive salivary gland tumors   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | EMSAM  |
| <b>Drug Names</b>                   | EMSAM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | 1) Patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to TWO of the following: a) serotonin and norepinephrine reuptake inhibitors (SNRIs), b) selective serotonin reuptake inhibitors (SSRIs), c) mirtazapine, d) bupropion OR 2) Patient is unable to swallow oral formulations. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ENBREL   |
| <b>Drug Names</b>                   | ENBREL, ENBREL MINI, ENBREL SURECLICK  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Hidradenitis suppurativa   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For moderately to severely active rheumatoid arthritis (new starts only): 1) Inadequate response, intolerance or contraindication to methotrexate (MTX) OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis (new starts only): 1) Inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial OR 2) Intolerance or contraindication to NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis AND 2) Patient meets any of the following: a) Patient has experienced an inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, OR b) Pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, OR c) Patient has severe psoriasis that warrants a biologic as first-line therapy (i.e. at least 10% of the BSA or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected). For hidradenitis suppurativa (new starts only): patient has severe, refractory disease. |

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| <b>Age Restrictions</b>        | -         |
| <b>Prescriber Restrictions</b> | -         |
| <b>Coverage Duration</b>       | Plan Year |
| <b>Other Criteria</b>          | -         |

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| <b>Prior Authorization Group</b>    | ENDARI                       |
| <b>Drug Names</b>                   | ENDARI                       |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | 5 years of age or older      |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

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| <b>Prior Authorization Group</b>    | EPCLUSA  |
| <b>Drug Names</b>                   | EPCLUSA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases (AASLD) treatment guidelines. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Criteria will be applied consistent with current AASLD-IDSA guidance.  |
| <b>Other Criteria</b>               | -  |
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| <b>Prior Authorization Group</b>    | EPIDIOLEX  |
| <b>Drug Names</b>                   | EPIDIOLEX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | 1 year of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | EPRONTIA   |
| <b>Drug Names</b>                   | EPRONTIA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For treatment of partial-onset seizures: 1)The patient has experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) the patient has experienced an inadequate treatment response, intolerance, or contraindication to any of the following: Aptiom, Vimpat, Xcopri, Spritam. For monotherapy treatment of primary generalized tonic-clonic seizures: 1) The patient has experienced an inadequate treatment response or intolerance to topiramate tablets or capsules, OR 2) The patient has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules). For adjunctive treatment of primary generalized tonic-clonic seizures: 1) The patient has experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) If the patient is 4 years of age or older, the patient has experienced an inadequate treatment response, intolerance, or contraindication to Spritam or Vimpat. For the preventative treatment of migraines: 1) The patient has experienced an inadequate treatment response or intolerance to topiramate tablets or capsules, OR 2) The patient has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules). |
| <b>Age Restrictions</b>             | Epilepsy: 2 years of age or older, Migraine: 12 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ERGOTAMINE   |
| <b>Drug Names</b>                   | ERGOTAMINE TARTRATE/CAFFE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | Coverage will be denied when used in conjunction with potent CYP3A4 inhibitors (e.g., ritonavir, nelfinavir, indinavir, erythromycin, clarithromycin).   |
| <b>Required Medical Information</b> | The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least ONE triptan 5-HT1 agonist.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | ERIVEDGE   |
| <b>Drug Names</b>                   | ERIVEDGE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Adult medulloblastoma  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Adult medulloblastoma: patient has received chemotherapy previously AND has tumor(s) with mutations in the sonic hedgehog pathway  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ERLEADA  |
| <b>Drug Names</b>                   | ERLEADA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ERLOTINIB  |
| <b>Drug Names</b>                   | ERLOTINIB HYDROCHLORIDE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent non-small cell lung cancer (NSCLC), recurrent chordoma, relapsed or stage IV renal cell carcinoma (RCC), brain metastases from non-small cell lung cancer (NSCLC), recurrent pancreatic cancer.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For NSCLC (including brain metastases from NSCLC): 1) the disease is recurrent, advanced, or metastatic and 2) the patient has sensitizing EGFR mutation-positive disease. For pancreatic cancer: the disease is locally advanced, unresectable, recurrent, or metastatic. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ESBRIET   |
| <b>Drug Names</b>                   | ESBRIET, PIRFENIDONE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For idiopathic pulmonary fibrosis (Initial Review Only): 1) a high-resolution computed tomography (HRCT) study of the chest or a lung biopsy reveals the usual interstitial pneumonia (UIP) pattern, OR 2) HRCT study of the chest reveals a result other than the UIP pattern (e.g., probable UIP, indeterminate for UIP) and the diagnosis is supported either by a lung biopsy or by a multidisciplinary discussion between at least a radiologist and pulmonologist who are experienced in idiopathic pulmonary fibrosis if a lung biopsy has not been conducted. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | EVEROLIMUS  |
| <b>Drug Names</b>                   | EVEROLIMUS  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Classic Hodgkin lymphoma, thymomas and thymic carcinomas, previously treated Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma, soft tissue sarcoma (perivascular epithelioid cell tumors (PEComa) and lymphangiomyomatosis subtypes), gastrointestinal stromal tumors, neuroendocrine tumors of the thymus, well differentiated grade 3 neuroendocrine tumors, thyroid carcinoma (papillary, Hurthle cell, and follicular), endometrial carcinoma, histiocytic neoplasms (Rosai-Dorfman Disease, Erdheim-Chester Disease, Langerhans Cell Histiocytosis)  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For breast cancer: 1) The disease is recurrent, advanced, or metastatic hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, AND 2) The requested drug is prescribed in combination with exemestane, fulvestrant, or tamoxifen, AND 3) The requested drug is used for subsequent treatment. For renal cell carcinoma: The disease is relapsed, advanced, or stage IV. For subependymal giant cell astrocytoma (SEGA): The requested drug is given as adjuvant treatment. For gastrointestinal stromal tumor: The disease is recurrent, unresectable, or metastatic AND the patient failed an FDA-approved therapy (e.g., imatinib, sunitinib, regorafenib, ripretinib). For symptomatic or relapsed/refractory Erdheim-Chester Disease (ECD), symptomatic or relapsed/refractory Rosai-Dorfman Disease, and Langerhans Cell Histiocytosis (LCH): the patient must have a phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit alpha (PIK3CA) mutation. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | EXKIVITY  |
| <b>Drug Names</b>                   | EXKIVITY  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | FABRAZYME  |
| <b>Drug Names</b>                   | FABRAZYME  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient meets ANY of the following: 1) Diagnosis of Fabry disease was confirmed by an enzyme assay demonstrating a deficiency of alpha-galactosidase enzyme activity or by genetic testing, OR 2) The patient is a symptomatic obligate female carrier.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | FANAPT   |
| <b>Drug Names</b>                   | FANAPT, FANAPT TITRATION PACK  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For treatment of schizophrenia: 1) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following generic products: aripiprazole, asenapine, olanzapine, quetiapine, risperidone, ziprasidone AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following brand products: Latuda, Rexulti, Secuado, Vraylar. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | FASENRA  |
| <b>Drug Names</b>                   | FASENRA, FASENRA PEN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For severe asthma: For initial therapy: 1) Either a) Patient has baseline blood eosinophil count of at least 150 cells per microliter OR b) Patient is dependent on systemic corticosteroids, and 2) Patient has a history of severe asthma despite current treatment with both of the following medications: a) inhaled corticosteroid and b) additional controller (long-acting beta2-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline) unless patient has an intolerance or contraindication to such therapies. For continuation of therapy: Asthma control has improved on treatment with the requested drug, as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations or a reduction in the daily maintenance oral corticosteroid dose. |
| <b>Age Restrictions</b>             | 12 years of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | FENTANYL PATCH   |
| <b>Drug Names</b>                   | FENTANYL   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the patient meets all of the following: 1) The requested drug is being prescribed for pain severe enough to require daily, around-the-clock, long-term treatment in a patient who has been taking an opioid AND 2) The patient can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The patient has been evaluated and the patient will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a patient who has been receiving an extended-release opioid agent for at least 30 days OR the patient has taken an immediate-release opioid for at least one week. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | FETZIMA  |
| <b>Drug Names</b>                   | FETZIMA, FETZIMA TITRATION PACK  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to TWO of the following: serotonin and norepinephrine reuptake inhibitors (SNRIs), selective serotonin reuptake inhibitors (SSRIs), mirtazapine, bupropion. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | FINTEPLA   |
| <b>Drug Names</b>                   | FINTEPLA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | 2 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | FLUCYTOSINE  |
| <b>Drug Names</b>                   | FLUCYTOSINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 6 weeks  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | FORTEO   |
| <b>Drug Names</b>                   | FORTEO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For postmenopausal osteoporosis: patient has ONE of the following (1 or 2): 1) a history of fragility fracture, OR 2) A pre-treatment T-score of less than or equal to -2.5 or pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment Fracture Risk Assessment Tool (FRAX) fracture probability AND patient has ANY of the following: a) indicators for higher fracture risk (e.g., advanced age, frailty, glucocorticoid therapy, very low T-scores, or increased fall risk), OR b) patient has failed prior treatment with or is intolerant to a previous injectable osteoporosis therapy OR c) patient has had an oral bisphosphonate trial of at least 1-year duration or there is a clinical reason to avoid treatment with an oral bisphosphonate. For primary or hypogonadal osteoporosis in men: patient has one of the following: 1) a history of osteoporotic vertebral or hip fracture, OR 2) pre-treatment T-score of less than or equal to -2.5, OR 3) pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment FRAX fracture probability. For glucocorticoid-induced osteoporosis: Patient has had an oral bisphosphonate trial of at least 1-year duration unless patient has a contraindication or intolerance to an oral bisphosphonate, AND patient meets ANY of the following: 1) patient has a history of fragility fracture, OR 2) a pre-treatment T-score of less than or equal to -2.5, OR 3) pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment FRAX fracture probability. Continuation of therapy: If the patient has received greater than or equal to 24 months of therapy with any parathyroid hormone analog: 1) The patient remains at or has returned to having a high risk for fracture, AND 2) The benefit of therapy with this prescribed medication outweighs the potential risks for this patient. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Initial: 24 months, Continuation: Plan Year  |
| <b>Other Criteria</b>               | Patient has high FRAX fracture probability if the 10 year probability is either greater than or equal to 20 percent for any major osteoporotic fracture or greater than or equal to 3 percent for hip fracture. If glucocorticoid treatment is greater than 7.5 mg (prednisone equivalent) per day, the estimated risk score generated with FRAX should be multiplied by 1.15 for major osteoporotic fracture and 1.2 for hip fracture.  |

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| <b>Prior Authorization Group</b>    | FOTIVDA   |
| <b>Drug Names</b>                   | FOTIVDA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For advanced renal cell carcinoma: the following criteria must be met: 1) The disease is relapsed or refractory, 2) The requested drug must be used after at least two prior systemic therapies, and 3) The patient has experienced disease progression or an intolerable adverse event with a trial of cabozantinib (Cabometyx). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | FYCOMPA   |
| <b>Drug Names</b>                   | FYCOMPA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For treatment of partial-onset seizures: 1) The patient experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) The patient has experienced an inadequate treatment response, intolerance, or contraindication to any of the following: Aptiom, Vimpat, Xcopri, Spritam. For adjunctive treatment of primary generalized tonic-clonic seizures: 1) The patient experienced an inadequate treatment response, intolerance, or contraindication to a generic anticonvulsant AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following: Vimpat, Spritam. |
| <b>Age Restrictions</b>             | Partial-onset seizures: 4 years of age or older. Primary generalized tonic-clonic seizures: 12 years of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |



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| <b>Prior Authorization Group</b>    | GATTEX   |
| <b>Drug Names</b>                   | GATTEX   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For short bowel syndrome (SBS) initial therapy: Adult patients were dependent on parenteral support for at least 12 months. For SBS continuation: Requirement for parenteral support has decreased from baseline while on therapy with the requested drug. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with a gastroenterologist, gastrointestinal surgeon, or nutritional support specialist.   |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | GAVRETO  |
| <b>Drug Names</b>                   | GAVRETO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent rearranged during transfection (RET) rearrangement-positive non-small cell lung cancer   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For non-small cell lung cancer, patient must meet all of the following: 1) The disease is recurrent, advanced, or metastatic, and 2) The tumor is rearranged during transfection (RET) fusion-positive or RET rearrangement-positive.                      |
| <b>Age Restrictions</b>             | Non-small cell lung cancer: 18 years of age or older. Medullary thyroid cancer and thyroid cancer: 12 years of age or older.   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | GILENYA  |
| <b>Drug Names</b>                   | FINGOLIMOD, GILENYA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | GILOTRIF  |
| <b>Drug Names</b>                   | GILOTRIF  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For non-small cell lung cancer (NSCLC): Patient meets either of the following: 1) Patient has metastatic squamous NSCLC that progressed after platinum-based chemotherapy, OR 2) Patient has sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | GLATIRAMER  |
| <b>Drug Names</b>                   | GLATIRAMER ACETATE, GLATOPA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | GROWTH HORMONE   |
| <b>Drug Names</b>                   | GENOTROPIN, GENOTROPIN MINIQUICK   |
| <b>PA Indication Indicator</b>      | All Medically-accepted Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | Pediatric patients with closed epiphyses   |
| <b>Required Medical Information</b> | Pediatric growth hormone deficiency (GHD): Patient (pt) is a neonate or was diagnosed with GHD as a neonate OR meets any of the following: 1) younger than 2.5 years old (yo) with pre-treatment (pre-tx) height (ht) more than 2 standard deviations (SD) below mean and slow growth velocity OR 2) 2.5 yo or older AND one of the following: a) pre-tx 1-year ht velocity more than 2 SD below mean OR b) pre-tx ht more than 2 SD below mean and 1-year ht velocity more than 1 SD below mean, AND patient meets any of the following: 1) failed 2 pre-tx growth hormone (GH) stimulation tests (peak below 10 ng/mL), OR 2) pituitary/central nervous system (CNS) disorder (e.g., genetic defects, acquired structural abnormalities, congenital structural abnormalities) and pre-tx insulin-like growth factor-1 (IGF-1) more than 2 SD below mean. Turner syndrome: 1) Confirmed by karyotyping AND 2) pre-tx ht is less than the 5th percentile for age. Small for gestational age (SGA): 1) Birth weight (wt) less than 2500g at gestational age (GA) greater than 37 weeks, OR birth wt or length below 3rd percentile for GA or at least 2 SD below mean for GA, AND 2) did not manifest catch-up growth by age 2.   |
| <b>Age Restrictions</b>             | SGA: 2 years of age or older   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an endocrinologist, pediatric endocrinologist, nephrologist, infectious disease specialist, gastroenterologist/nutritional support specialist, or geneticist.  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Adult GHD: Pt meets any of the following: 1) failed 2 pre-tx GH stimulation tests, OR 2) pre-tx IGF-1 more than 2 SD below mean AND failed 1 pre-tx GH stimulation test. (Note: Stimulation tests include: a) insulin tolerance test [ITT] [peak GH less than or equal to 5 ng/ml], or b) Macrilen-stimulation test [peak GH level less than 2.8ng/ml], or c) glucagon-stimulation test [GST] [peak GH level less than or equal to 3 ng/ml] for pt with a body mass index [BMI] 25-30 kg/m <sup>2</sup> and high pretest probability of GHD [e.g., acquired structural abnormalities] or BMI less than 25 kg/m <sup>2</sup> , or d) GST [peak GH level less than or equal to 1 ng/ml] in pt with BMI 25-30 kg/m <sup>2</sup> and low pretest probability of GHD or BMI greater than 30 kg/m <sup>2</sup> ), OR 3) organic hypothalamic-pituitary disease (e.g., suprasellar mass with previous surgery and cranial irradiation) with 3 or more pituitary hormone deficiencies AND pre-tx IGF-1 more than 2 SD below mean, OR 4) genetic or structural hypothalamic-pituitary defects, OR 5) childhood-onset GHD with congenital (genetic or structural) abnormality of the hypothalamus/pituitary/CNS. Renewal for pediatric GHD, TS, SGA, and adult GHD: Patient is experiencing improvement. |

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| <b>Prior Authorization Group</b>    | HAEGARDA  |
| <b>Drug Names</b>                   | HAEGARDA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For hereditary angioedema: The requested drug is being used for the prevention of acute angioedema attacks. Patient has hereditary angioedema (HAE) with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has hereditary angioedema with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, EITHER 1) Patient tested positive for an F12, angiotensin-converting enzyme 1 (ACE1), plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosaminase 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation OR 2) Patient has a family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy for at least one month. |
| <b>Age Restrictions</b>             | 6 years of age or older   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an immunologist, allergist, or rheumatologist   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | HARVONI  |
| <b>Drug Names</b>                   | HARVONI  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases (AASLD) treatment guidelines. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Criteria applied consistent w/ current AASLD-IDSA guidance. Reminder for 8wk option if appropriate.  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | HERCEPTIN  |
| <b>Drug Names</b>                   | HERCEPTIN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |
| <b>Prior Authorization Group</b>    | HERCEPTIN HYLECTA  |
| <b>Drug Names</b>                   | HERCEPTIN HYLECTA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

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| <b>Prior Authorization Group</b>    | HERZUMA   |
| <b>Drug Names</b>                   | HERZUMA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The patient had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year   |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.  |
| <b>Prior Authorization Group</b>    | HETLIOZ   |
| <b>Drug Names</b>                   | HETLIOZ, TASIMELTEON  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For Non-24-Hour Sleep-Wake Disorder: 1) for initial therapy and continuation of therapy: a) diagnosis of total blindness in both eyes (e.g., nonfunctioning retinas) and b) unable to perceive light in either eye, AND 2) if currently on therapy with the requested drug, patient must meet at least one of the following: a) increased total nighttime sleep or b) decreased daytime nap duration. For nighttime sleep disturbances in Smith-Magenis Syndrome (SMS): 1) for initial therapy and continuation therapy, the patient has a confirmed diagnosis of SMS AND 2) if currently on therapy with the requested drug, the patient experiences improvement in the quality of sleep since starting therapy. |
| <b>Age Restrictions</b>             | Non-24: 18 years of age or older. SMS: 16 years of age or older   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with sleep disorder specialist or neurologist  |
| <b>Coverage Duration</b>            | Initiation: 6 Months, Renewal: Plan Year  |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | HRM-ANTICONVULSANTS  |
| <b>Drug Names</b>                   | PHENOBARBITAL, PHENOBARBITAL SODIUM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Epilepsy   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization requirement only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) |

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| <b>Prior Authorization Group</b>    | HRM-ANTIPARKINSON   |
| <b>Drug Names</b>                   | BENZTROPINE MESYLATE, TRIHEXYPHENIDYL HCL, TRIHEXYPHENIDYL HYDROCHLO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. EPS (extrapyramidal symptoms): 1) The patient has not tried the non-HRM alternative drug amantadine AND 2) The patient has a contraindication to the non-HRM alternative drug amantadine OR 3) The patient has tried the non-HRM alternative drug amantadine AND 4) The patient experienced an inadequate treatment response OR intolerance to the non-HRM alternative drug amantadine. Parkinson's: 1) The patient has tried two of the following non-HRM alternative drugs: amantadine, carbidopa/levodopa, pramipexole, or ropinirole. AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following non-HRM alternative drugs: amantadine, carbidopa/levodopa, pramipexole, or ropinirole. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)  |

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| <b>Prior Authorization Group</b>    | HRM-CYPROHEPTADINE  |
| <b>Drug Names</b>                   | CYPROHEPTADINE HCL, CYPROHEPTADINE HYDROCHLOR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Pruritus, spasticity due to spinal cord injury  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. For rhinitis: 1) The patient has tried two of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)  |

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| <b>Prior Authorization Group</b>    | HRM-DIPYRIDAMOLE   |
| <b>Drug Names</b>                   | DIPYRIDAMOLE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) |



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| <b>Prior Authorization Group</b>    | HRM-GUANFACINE ER  |
| <b>Drug Names</b>                   | GUANFACINE ER, GUANFACINE HYDROCHLORIDE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) |

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| <b>Prior Authorization Group</b>    | HRM-GUANFACINE IR  |
| <b>Drug Names</b>                   | GUANFACINE HCL, GUANFACINE HYDROCHLORIDE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) |

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| <b>Prior Authorization Group</b>    | HRM-HYDROXYZINE  |
| <b>Drug Names</b>                   | HYDROXYZINE HCL, HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE PAMOATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For anxiety: 1) The patient has tried two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline, or venlafaxine extended-release AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline, or venlafaxine extended-release OR 3) The patient has not tried two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline or venlafaxine extended-release AND 4) The patient has acute anxiety. For all indications: 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. AND 2) If the patient is taking one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, cyclobenzaprine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the patient [Note: Use of multiple anticholinergic medications in older adults is associated with an increased risk of cognitive decline.]. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)   |

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| <b>Prior Authorization Group</b>    | HRM-HYDROXYZINE INJ  |
| <b>Drug Names</b>                   | HYDROXYZINE HCL, HYDROXYZINE HYDROCHLORIDE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | <p>Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. For alcohol withdrawal syndrome: 1) The patient has not tried one of the following alternative drugs: clorazepate or lorazepam AND 2) The patient has a contraindication to one of the following alternative drugs: clorazepate or lorazepam OR 3) The patient has tried one of the following alternative drugs: clorazepate or lorazepam AND 4) The patient experienced an inadequate treatment response OR intolerance to one of the following alternative drugs: clorazepate or lorazepam. For anxiety: 1) The patient has tried two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline or venlafaxine extended-release AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline or venlafaxine extended-release OR 3) The patient has not tried two of the following alternative drugs: buspirone, duloxetine, escitalopram, sertraline or venlafaxine extended-release AND 4) The patient has acute anxiety.</p> |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)   |

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| <b>Prior Authorization Group</b>    | HRM-HYPNOTICS   |
| <b>Drug Names</b>                   | ZOLPIDEM TARTRATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For insomnia: 1) The patient meets one of the following: a) the patient has a contraindication to the non-HRM (non-High Risk Medication) alternative drug doxepin (3 mg or 6 mg) OR b) The non-HRM (non-High Risk Medication) alternative drug doxepin (3 mg or 6 mg) has been tried AND the patient experienced an inadequate treatment response OR intolerance to the non-HRM (non-High Risk Medication) alternative drug doxepin (3 mg or 6 mg) AND 2) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient AND 3) If the patient is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, quetiapine, sertraline, clonazepam, escitalopram, alprazolam) with the requested drug, the prescriber has determined that taking multiple central nervous system (CNS) active medications is medically necessary for the patient [Note: Use of multiple central nervous system (CNS) active medications in older adults is associated with an increased risk of falls.]. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) APPLIES TO GREATER THAN CUMULATIVE 90 DAYS OF THERAPY PER YEAR.  |

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| <b>Prior Authorization Group</b>    | HRM-PROMETHAZINE  |
| <b>Drug Names</b>                   | PROMETHAZINE HCL, PROMETHAZINE HCL PLAIN, PROMETHAZINE HYDROCHLORID   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. For rhinitis: 1) The patient has tried two of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal AND 2) The patient experienced an inadequate treatment response OR intolerance to two of the following non-HRM alternative drugs: levocetirizine, azelastine nasal, fluticasone nasal, or flunisolide nasal. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)  |

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| <b>Prior Authorization Group</b>    | HRM-SCOPOLAMINE  |
| <b>Drug Names</b>                   | SCOPOLAMINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Excessive salivation   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) |

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| <b>Prior Authorization Group</b>    | HRM-SKELETAL MUSCLE RELAXANTS  |
| <b>Drug Names</b>                   | CYCLOBENZAPRINE HYDROCHLO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | 1) Prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for this patient. AND 2) If the patient is using one or more additional anticholinergic medications (e.g., oxybutynin, meclizine, paroxetine, amitriptyline, dicyclomine, hydroxyzine) with the requested drug, the prescriber has determined that taking multiple anticholinergic medications is medically necessary for the patient [Note: Use of multiple anticholinergic medications in older adults is associated with an increased risk of cognitive decline.]. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 3 months   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 70 years of age or older. (The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.)   |

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| <b>Prior Authorization Group</b>    | HUMIRA  |
| <b>Drug Names</b>                   | HUMIRA, HUMIRA PEDIATRIC CROHNS D, HUMIRA PEN, HUMIRA PEN-CD/UC/HS START, HUMIRA PEN-PEDIATRIC UC S, HUMIRA PEN-PS/UV STARTER   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Axial spondyloarthritis, Behcet's syndrome  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderately to severely active rheumatoid arthritis (new starts only): 1) Inadequate response, intolerance or contraindication to methotrexate (MTX) OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis and axial spondyloarthritis (new starts only): 1) Inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial OR 2) Intolerance or contraindication to NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) Patient meets any of the following: a) Patient has experienced an inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, OR b) Pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, OR c) Patient has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of the BSA or crucial body areas (e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas) are affected). For moderately to severely active Crohn's disease (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 2) Intolerance or contraindication to conventional therapy. For moderately to severely active ulcerative colitis (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 2) Intolerance or contraindication to conventional therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | IBRANCE  |
| <b>Drug Names</b>                   | IBRANCE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Unresectable well-differentiated/dedifferentiated liposarcoma of the retroperitoneum, recurrent hormone receptor-positive human epidermal growth factor receptor 2 (HER2)-negative breast cancer   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ICATIBANT  |
| <b>Drug Names</b>                   | ICATIBANT ACETATE, SAJAZIR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hereditary angioedema (HAE): The requested drug is being used for the treatment of acute angioedema attacks. Patient has HAE with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing OR patient has HAE with normal C1 inhibitor confirmed by laboratory testing. For patients with HAE with normal C1 inhibitor, EITHER 1) Patient tested positive for an F12, angiotensin-converting enzyme 1 (ACE1), plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosaminase 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation OR 2) Patient has a family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy for at least one month. |
| <b>Age Restrictions</b>             | 18 years of age or older   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an immunologist, allergist, or rheumatologist  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | ICLUSIG  |
| <b>Drug Names</b>                   | ICLUSIG  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Myeloid and/or lymphoid neoplasms with eosinophilia and FGFR1 or ABL1 rearrangement in the chronic phase or blast phase  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML) or acute lymphoblastic leukemia (ALL), including patients who have received a hematopoietic stem cell transplant: diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML, including patients who have received a hematopoietic stem cell transplant: 1) patient has accelerated or blast phase CML and no other kinase inhibitor is indicated, OR 2) patient has chronic phase CML and has experienced resistance or intolerance to at least 2 prior kinase inhibitors AND at least one of those was imatinib or dasatinib, OR 3) patient is positive for the T315I mutation. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | IDHIFA   |
| <b>Drug Names</b>                   | IDHIFA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Newly-diagnosed acute myeloid leukemia   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acute myeloid leukemia (AML) with an isocitrate dehydrogenase-2 (IDH2) mutation: 1) patient is 60 years of age or older with newly-diagnosed AML and meets one of the following: a) patient is not a candidate for intensive induction therapy, or b) patient declines intensive induction chemotherapy, OR 2) patient is 60 years of age or older and the requested drug will be used as post-induction therapy following response to induction therapy with the requested drug OR 3) patient has relapsed or refractory AML.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | IMATINIB  |
| <b>Drug Names</b>                   | IMATINIB MESYLATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Desmoid tumors, pigmented villonodular synovitis/tenosynovial giant cell tumor (PVNS/TGCT), recurrent chordoma, melanoma, Kaposi sarcoma, chronic myelomonocytic leukemia, chronic graft versus host disease (cGVHD), T-cell acute lymphoblastic leukemia with ABL-class translocation, aggressive systemic mastocytosis for well-differentiated systemic mastocytosis (WDSM) or when eosinophilia is present with FIP1L1-PDGFR A fusion gene, myeloid and/or lymphoid neoplasms with eosinophilia and ABL1, FIP1L1-PDGFR A, or PDGFR B rearrangement in the chronic phase or blast phase |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), including patients who have received a hematopoietic stem cell transplant: diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML: patient did not fail (excluding failure due to intolerance) prior therapy with a tyrosine kinase inhibitor. For melanoma: c-Kit mutation is positive.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | IMBRUVICA  |
| <b>Drug Names</b>                   | IMBRUVICA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Hairy cell leukemia, lymphoplasmacytic lymphoma, primary central nervous system (CNS) lymphoma, AIDS-related B-cell lymphoma, diffuse large B-cell lymphoma, post-transplant lymphoproliferative disorders, high-grade B-cell lymphoma   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For mantle cell lymphoma: 1) the requested drug will be used in a patient who has received at least one prior therapy, OR 2) the requested drug will be used in combination with rituximab as pretreatment to induction therapy with RHyperCVAD (rituximab, cyclophosphamide, vincristine, doxorubicin, and dexamethasone) regimen OR 3) the patient is 65 years of age or older AND the requested drug will be used in combination with rituximab. For marginal zone lymphoma (including gastric mucosa-associated lymphoid tissue [MALT] lymphoma, non-gastric MALT lymphoma, nodal marginal zone lymphoma, and splenic marginal zone lymphoma): the patient has received at least one prior therapy. For hairy cell leukemia: the requested drug will be used as a single agent for disease progression. For primary CNS lymphoma: 1) the disease is relapsed or refractory OR 2) the requested drug is used for induction therapy as a single agent. For diffuse large B-cell lymphoma and high-grade B-cell lymphoma: the requested drug will be used as second-line or subsequent therapy. For AIDS-related B-cell lymphoma: the requested drug will be used as a single agent and as second-line or subsequent therapy for relapsed disease. For post-transplant lymphoproliferative disorders: the requested drug will be used in patients who have received prior chemoimmunotherapy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | INCRELEX  |
| <b>Drug Names</b>                   | INCRELEX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | Pediatric patients with closed epiphyses  |
| <b>Required Medical Information</b> | For growth failure due to severe primary insulin-like growth factor-1 (IGF-1) deficiency or growth hormone (GH) gene deletion in patients who have developed neutralizing antibodies to GH, patient meets all of the following prior to beginning therapy with the requested drug (new starts only): 1) height 3 or more standard deviations (SD) below the mean for children of the same age and gender AND 2) basal IGF-1 level 3 or more SD below the mean for children of the same age and gender AND 3) provocative growth hormone test showing a normal or elevated growth hormone level. For growth failure due to severe primary IGF-1 deficiency or GH gene deletion in patients who have developed neutralizing antibodies to GH, continuation of therapy: patient is experiencing improvement. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | INGREZZA  |
| <b>Drug Names</b>                   | INGREZZA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | INLYTA  |
| <b>Drug Names</b>                   | INLYTA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Thyroid carcinoma (papillary, Hurthle cell, or follicular)  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For renal cell carcinoma: The disease is advanced, relapsed, or stage IV.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | INQOVI   |
| <b>Drug Names</b>                   | INQOVI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | INREBIC  |
| <b>Drug Names</b>                   | INREBIC  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and janus kinase 2 (JAK2) rearrangement, accelerated phase myelofibrosis, blast phase myelofibrosis/acute myeloid leukemia |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement: the disease is in chronic or blast phase.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | IR BEFORE ER   |
| <b>Drug Names</b>                   | HYDROCODONE BITARTRATE ER, HYSINGLA ER, METHADONE HCL, METHADONE HYDROCHLORIDE I, MORPHINE SULFATE ER  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug is being prescribed for pain associated with cancer, sickle cell disease, a terminal condition, or pain being managed through palliative care OR the patient meets all of the following: 1) The requested drug is being prescribed for pain severe enough to require daily, around-the-clock, long-term treatment in a patient who has been taking an opioid AND 2) The patient can safely take the requested dose based on their history of opioid use [Note: This drug should be prescribed only by healthcare professionals who are knowledgeable in the use of potent opioids for the management of chronic pain.] AND 3) The patient has been evaluated and the patient will be monitored for the development of opioid use disorder AND 4) This request is for continuation of therapy for a patient who has been receiving an extended-release opioid agent for at least 30 days OR the patient has taken an immediate-release opioid for at least one week. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | IRESSA   |
| <b>Drug Names</b>                   | IRESSA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Sensitizing epidermal growth factor receptor (EGFR) mutation-positive recurrent non-small cell lung cancer (NSCLC).  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For non-small cell lung cancer (NSCLC): 1) disease must be metastatic, advanced, or recurrent AND 2) patient must have a sensitizing EGFR mutation.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ISOTRETINOIN   |
| <b>Drug Names</b>                   | ACCUTANE, AMNESTEEM, CLARAVIS, ISOTRETINOIN, MYORISAN, ZENATANE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Refractory acne vulgaris, severe refractory rosacea, neuroblastoma, cutaneous T-cell lymphoma (CTCL) (e.g., mycosis fungoides, Sezary syndrome), high risk for developing skin cancer (squamous cell cancers), transient acantholytic dermatosis (Grover's Disease), keratosis follicularis (Darier Disease), lamellar ichthyosis, pityriasis rubra pilaris. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ITRACONAZOLE   |
| <b>Drug Names</b>                   | ITRACONAZOLE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Coccidioidomycosis, Coccidioidomycosis prophylaxis in HIV infection, Cryptococcosis, Microsporidiosis, Talaromycosis (formerly Penicilliosis), Histoplasmosis prophylaxis in HIV infection, Invasive fungal infection prophylaxis in liver transplant, chronic granulomatous disease (CGD), and hematologic malignancy, Sporotrichosis, Pityriasis versicolor, Tinea versicolor, Tinea corporis, Tinea cruris, Tinea capitis, Tinea manuum, Tinea pedis. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used orally. For the treatment of onychomycosis due to dermatophytes (Tinea unguium), the diagnosis has been confirmed by a fungal diagnostic test (e.g., potassium hydroxide [KOH] preparation, fungal culture, or nail biopsy).   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Disseminated/CNS histoplasmosis, Histoplasmosis/Coccidioidomycosis/CGD ppx: 12 mths. Others: 6 mths  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | IVERMECTIN TAB   |
| <b>Drug Names</b>                   | IVERMECTIN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Ascariasis, Cutaneous larva migrans, Mansonelliasis, Scabies, Gnathostomiasis, Pediculosis   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug is not being prescribed for the prevention or treatment of coronavirus disease 2019 (COVID-19).   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 1 month  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | IVIG   |
| <b>Drug Names</b>                   | BIVIGAM, FLEBOGAMMA DIF, GAMMAGARD LIQUID, GAMMAGARD S/D IGA LESS TH, GAMMAKED, GAMMAPLEX, GAMUNEX-C, OCTAGAM, PANZYGA, PRIVIGEN   |
| <b>PA Indication Indicator</b>      | All Medically-accepted Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For B-cell chronic lymphocytic leukemia (CLL): 1) serum IgG less than 500 mg/dL OR 2) a history of recurrent bacterial infections. For bone marrow transplant/hematopoietic stem cell transplant (BMT/HSCT): 1) IVIG is requested within the first 100 days post-transplant OR 2) serum IgG less than 400 mg/dL. For pediatric human immunodeficiency virus (HIV) infection: 1) serum IgG less than 400 mg/dL OR 2) history of recurrent bacterial infections. For dermatomyositis and polymyositis: 1) at least one standard first-line treatment (corticosteroid or immunosuppressant) has been tried but was unsuccessful or not tolerated OR 2) patient is unable to receive standard therapy because of a contraindication or other clinical reason. For pure red cell aplasia (PRCA): PRCA is secondary to parvovirus B19 infection. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |



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| <b>Prior Authorization Group</b>    | JAKAFI   |
| <b>Drug Names</b>                   | JAKAFI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Lower-risk myelofibrosis, accelerated phase myelofibrosis, blast phase myelofibrosis/acute myeloid leukemia, acute lymphoblastic leukemia (ALL), chronic myelomonocytic leukemia (CMML)-2, BCR-ABL negative atypical chronic myeloid leukemia (aCML), essential thrombocythemia, and myeloid, lymphoid or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For polycythemia vera: patient had an inadequate response or intolerance to interferon therapy or hydroxyurea. For acute lymphoblastic leukemia: patient has a cytokine receptor-like factor 2 (CRLF2) mutation or a mutation associated with activation of the Janus kinase/signal transducers and activators of transcription (JAK/STAT) pathway. For CMML-2: the requested drug is used in combination with a hypomethylating agent. For BCR-ABL negative aCML: the requested drug is used as a single agent or in combination with a hypomethylating agent. For essential thrombocythemia: patient had an inadequate response or loss of response to hydroxyurea, interferon therapy, or anagrelide. For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and JAK2 rearrangement: the disease is in chronic or blast phase. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | KALYDECO   |
| <b>Drug Names</b>                   | KALYDECO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis (CF): The requested medication will not be used in combination with other medications containing ivacaftor.  |
| <b>Age Restrictions</b>             | 4 months of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | KANJINTI   |
| <b>Drug Names</b>                   | KANJINTI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |
| <b>Prior Authorization Group</b>    | KESIMPTA   |
| <b>Drug Names</b>                   | KESIMPTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

|                                     |  |
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| <b>Prior Authorization Group</b>    | KETOCONAZOLE   |
| <b>Drug Names</b>                   | KETOCONAZOLE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Cushing's syndrome   |
| <b>Exclusion Criteria</b>           | Acute or chronic liver disease. Concurrent use with drugs that are contraindicated with ketoconazole tablets: dofetilide, quinidine, pimozone, cisapride, methadone, disopyramide, dronedarone, ranolazine, ergot alkaloids, irinotecan, lurasidone, oral midazolam, alprazolam, triazolam, felodipine, nisoldipine, tolvaptan, eplerenone, lovastatin, simvastatin, or colchicine.                          |
| <b>Required Medical Information</b> | The potential benefits outweigh the risks of treatment with oral ketoconazole. For systemic fungal infections, the patient has any of the following diagnoses: blastomycosis, coccidioidomycosis, histoplasmosis, chromomycosis, or paracoccidioidomycosis. For Cushing's syndrome: the requested drug is being prescribed for a patient who cannot tolerate surgery or where surgery has not been curative. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 6 months   |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | KEVZARA  |
| <b>Drug Names</b>                   | KEVZARA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For moderately to severely active rheumatoid arthritis (new starts only): 1) Inadequate response, intolerance or contraindication to methotrexate (MTX) OR 2) Inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

**Prior Authorization Group** KEYTRUDA  
**Drug Names** KEYTRUDA  
**PA Indication Indicator** All Medically-accepted Indications  
**Off-label Uses** -  
**Exclusion Criteria** -  
**Required Medical Information** -  
**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

**Prior Authorization Group** KISQALI  
**Drug Names** KISQALI, KISQALI FEMARA 200 DOSE, KISQALI FEMARA 400 DOSE, KISQALI FEMARA 600 DOSE  
**PA Indication Indicator** All FDA-approved Indications, Some Medically-accepted Indications  
**Off-label Uses** Recurrent hormone receptor-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer, in combination with an aromatase inhibitor, or fulvestrant.  
**Exclusion Criteria** -  
**Required Medical Information** For treatment of breast cancer using Kisqali (ribociclib) in combination with an aromatase inhibitor or Kisqali Femara Co-Pack (ribociclib and letrozole) as initial endocrine-based therapy, one of the following criteria must be met: 1) the patient is pre- or peri-menopausal OR 2) the patient is a postmenopausal OR male AND the patient has experienced an intolerable adverse event to Ibrance (palbociclib) AND Verzenio (abemaciclib) or has a contraindication to Ibrance (palbociclib) AND Verzenio (abemaciclib). For treatment of breast cancer with Kisqali (ribociclib) in combination with fulvestrant, one of the following criteria must met: 1) the requested drug is being used with fulvestrant as initial endocrine-based therapy in a postmenopausal patient OR a male, OR 2) the requested drug is being used following disease progression on endocrine therapy in a postmenopausal patient OR a male, and the patient has experienced an intolerable adverse event to Ibrance (palbociclib) AND Verzenio (abemaciclib) OR has a contraindication to Ibrance (palbociclib) AND Verzenio (abemaciclib).  
**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

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| <b>Prior Authorization Group</b>    | KORLYM  |
| <b>Drug Names</b>                   | KORLYM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an endocrinologist  |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | KYNMOBI   |
| <b>Drug Names</b>                   | KYNMOBI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For continuation treatment of off episodes in Parkinson's disease: The patient is experiencing improvement on the requested drug.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | LAPATINIB   |
| <b>Drug Names</b>                   | LAPATINIB DITOSYLATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Brain metastases from human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent HER2-positive breast cancer, recurrent epidermal growth factor receptor (EGFR)-positive chordoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with trastuzumab.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For breast cancer, the patient meets all the following: a) the disease is recurrent, advanced, or metastatic (including brain metastases), b) the disease is human epidermal growth factor receptor 2 (HER2)-positive, c) the requested drug will be used in combination with any of the following: 1) aromatase inhibitor, 2) capecitabine, OR 3) trastuzumab. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | LENVIMA   |
| <b>Drug Names</b>                   | LENVIMA 10 MG DAILY DOSE, LENVIMA 12MG 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   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Medullary thyroid carcinoma, recurrent endometrial carcinoma, thymic carcinoma  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For differentiated thyroid cancer (follicular, papillary, or Hurthle cell): disease is not amenable to radioactive iodine therapy and unresectable, locally recurrent, persistent, or metastatic. For hepatocellular carcinoma: disease is unresectable or inoperable, local, metastatic or with extensive liver tumor burden. For renal cell carcinoma, the disease is advanced, relapsed, or stage IV. For endometrial carcinoma, the patient meets ALL of the following: 1) The disease is advanced, recurrent, or metastatic, 2) The patient experienced disease progression following prior systemic therapy, AND 3) The patient is not a candidate for curative surgery or radiation. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | LEUPROLIDE  |
| <b>Drug Names</b>                   | LEUPROLIDE ACETATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Use in combination with growth hormone for children with growth failure and advancing puberty, recurrent androgen receptor positive salivary gland tumors   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | LIDOCAINE PATCHES  |
| <b>Drug Names</b>                   | LIDOCAINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Pain associated with diabetic neuropathy, pain associated with cancer-related neuropathy (including treatment-related neuropathy [e.g., neuropathy associated with radiation treatment or chemotherapy]).  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | LONSURF  |
| <b>Drug Names</b>                   | LONSURF  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For colorectal cancer: The disease is advanced or metastatic. For gastric or gastroesophageal junction adenocarcinoma, all of the following criteria must be met: 1) The disease is unresectable locally advanced, recurrent, or metastatic, and 2) The patient has been previously treated with at least two prior lines of chemotherapy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | LORBRENA   |
| <b>Drug Names</b>                   | LORBRENA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Anaplastic lymphoma kinase (ALK)-positive recurrent non-small cell lung cancer (NSCLC), repressor of silencing (ROS)-1 rearrangement-positive recurrent, advanced, or metastatic NSCLC.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For recurrent, advanced, or metastatic non-small cell lung cancer: 1) Disease is anaplastic lymphoma kinase (ALK)-positive OR 2) Disease is positive for repressor of silencing (ROS)-1 rearrangement and the requested drug is being used following disease progression on crizotinib, entrectinib, or ceritinib.                         |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

|                                     |  |
|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | LUMAKRAS   |
| <b>Drug Names</b>                   | LUMAKRAS   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | LUMIZYME   |
| <b>Drug Names</b>                   | LUMIZYME   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Diagnosis of Pompe disease was confirmed by an enzyme assay demonstrating a deficiency of acid alpha-glucosidase (GAA) enzyme activity or by genetic testing.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | LUPRON PED   |
| <b>Drug Names</b>                   | LUPRON DEPOT-PED (1-MONTH, LUPRON DEPOT-PED (3-MONTH   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For central precocious puberty (CPP), patients not currently receiving therapy must meet all of the following criteria: 1) Diagnosis of CPP was confirmed by a pubertal response to a gonadotropin releasing hormone (GnRH) agonist test OR a pubertal level of a third generation luteinizing hormone (LH) assay, 2) Assessment of bone age versus chronological age supports the diagnosis of CPP, and 3) The onset of secondary sexual characteristics occurred prior to 8 years of age for female patients OR prior to 9 years of age for male patients. |
| <b>Age Restrictions</b>             | CPP: Patient must be less than 12 years old if female and less than 13 years old if male.  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | LUPRON-ENDOMETRIOSIS   |
| <b>Drug Names</b>                   | LUPRON DEPOT (1-MONTH), LUPRON DEPOT (3-MONTH)   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Breast cancer, epithelial ovarian cancer/fallopian tube cancer/primary peritoneal cancer   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For uterine fibroids, patient must meet one of the following: 1) Diagnosis of anemia (e.g., hematocrit less than or equal to 30 percent and/or hemoglobin less than or equal to 10g/dL), OR 2) the requested medication will be used prior to surgery for uterine fibroids. For breast cancer, the requested drug is used for hormone receptor (HR)-positive disease.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Fibroids: 3 months (mo), max 6 mo total. Endometriosis: 6 mo, max 12 mo total. Others: Plan Year   |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | LYNPARZA   |
| <b>Drug Names</b>                   | LYNPARZA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent HER2-negative, BRCA 1/2-germline mutated breast cancer, recurrent or metastatic HER2-positive, BRCA 1/2-germline mutated breast cancer, uterine leiomyosarcoma.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For recurrent or metastatic breast cancer: the disease is BRCA 1/2-germline mutated. For prostate cancer: The patient has progressed on prior treatment with an androgen receptor-directed therapy. For epithelial ovarian, fallopian tube, or primary peritoneal cancer: The requested drug is used for maintenance therapy for stage II-IV or recurrent disease who are in complete or partial response to chemotherapy. For uterine leiomyosarcoma: 1) the requested drug is used as second-line therapy AND 2) the patient has BRCA-altered disease. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | MAVYRET  |
| <b>Drug Names</b>                   | MAVYRET  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh [CTP] class B or C).  |
| <b>Required Medical Information</b> | For hepatitis C virus (HCV): Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [CTP class B or C]), presence or absence of human immunodeficiency virus (HIV) coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases (AASLD) treatment guidelines. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Criteria will be applied consistent with current AASLD-IDSa guidance   |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | MEGESTROL  |
| <b>Drug Names</b>                   | MEGESTROL ACETATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Cancer-related cachexia in adults  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Patient has experienced an inadequate treatment response or intolerance to megestrol 40 mg/mL oral suspension.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | MEKINIST   |
| <b>Drug Names</b>                   | MEKINIST   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Brain metastases from melanoma, uveal melanoma, central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma), low grade serous ovarian cancer, ovarian borderline epithelial tumors (low malignant potential) with invasive implants, Langerhans cell histiocytosis, Erdheim-Chester disease, Rosai-Dorfman disease, gallbladder cancer, intrahepatic cholangiocarcinoma, extrahepatic cholangiocarcinoma   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For adjuvant treatment of melanoma,: 1) The tumor is positive for a BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used in combination with dabrafenib. For unresectable or metastatic melanoma: 1) The tumor is positive for a BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used as a single agent or in combination with dabrafenib. For brain metastases from melanoma, central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma), non-small cell lung cancer, solid tumors, and anaplastic thyroid cancer: 1) The tumor is positive for a BRAF V600E mutation, and 2) The requested drug will be used in combination with dabrafenib. For uveal melanoma, the requested drug will be used as a single agent. For low grade serous ovarian cancer and ovarian borderline epithelial tumors (low malignant potential) with invasive implants: The requested drug will be used to treat persistent or recurrent disease. For gallbladder cancer, intrahepatic cholangiocarcinoma, and extrahepatic cholangiocarcinoma: 1) The tumor is positive for a BRAF V600E mutation, 2) the disease is unresectable or metastatic, and 3) The requested drug will be used in combination with dabrafenib. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | MEKTOVI   |
| <b>Drug Names</b>                   | MEKTOVI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Adjuvant systemic therapy for cutaneous melanoma  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For cutaneous melanoma: The patient must meet all of the following criteria: 1) Tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K), 2) The requested drug will be used in combination with encorafenib, and 3) The requested drug will be used for either of the following: a) unresectable or metastatic disease, or b) adjuvant systemic therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | MEMANTINE   |
| <b>Drug Names</b>                   | MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE E  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This edit only applies to patients less than 30 years of age.   |
| <b>Prior Authorization Group</b>    | METHYLPHENIDATE   |
| <b>Drug Names</b>                   | METADATE ER, METHYLPHENIDATE HYDROCHLO  |
| <b>PA Indication Indicator</b>      | All Medically-accepted Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The patient has a diagnosis of Attention-Deficit Hyperactivity Disorder (ADHD) or Attention Deficit Disorder (ADD) OR 2) The patient has a diagnosis of narcolepsy confirmed by a sleep study OR 3) The requested drug is being prescribed for the treatment of cancer-related fatigue after other causes of fatigue have been ruled out.                                |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | MIGLUSTAT   |
| <b>Drug Names</b>                   | MIGLUSTAT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For type 1 Gaucher disease (GD1): The diagnosis was confirmed by an enzyme assay demonstrating a deficiency of beta-glucocerebrosidase enzyme activity or by genetic testing. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | MONJUVI   |
| <b>Drug Names</b>                   | MONJUVI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b> | MVASI   |
| <b>Drug Names</b>                | MVASI   |
| <b>PA Indication Indicator</b>   | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>            | Breast cancer, central nervous system (CNS) tumor types: adult low-grade (WHO Grade I or II) glioma, adult intracranial and spinal ependymoma, anaplastic gliomas, adult medulloblastoma, primary central nervous system lymphoma, meningiomas, limited and extensive brain metastases, and metastatic spine tumors, malignant pleural mesothelioma, ovarian cancer/fallopian tube cancer/primary peritoneal cancer types: carcinosarcoma (malignant mixed Mullerian tumors), clear cell carcinoma, mucinous carcinoma, grade 1 endometrioid carcinoma, low-grade serous carcinoma, ovarian borderline epithelial tumors (low malignant potential) with invasive implants, and malignant sex cord-stromal tumors, soft tissue sarcoma types: angiosarcoma and solitary fibrous tumor/hemangiopericytoma, uterine neoplasms, endometrial carcinoma, vulvar squamous cell carcinoma, and ophthalmic-related disorders: diabetic macular edema, neovascular (wet) age-related macular degeneration including polypoidal choroidopathy and retinal angiomatous proliferation subtypes, macular edema following retinal vein occlusion, proliferative diabetic retinopathy, choroidal neovascularization, neovascular glaucoma and retinopathy of prematurity, hepatocellular carcinoma, small bowel adenocarcinoma. |

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| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual. |

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| <b>Prior Authorization Group</b>    | NAGLAZYME   |
| <b>Drug Names</b>                   | NAGLAZYME   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Diagnosis of Mucopolysaccharidosis VI (Maroteaux-Lamy syndrome) was confirmed by an enzyme assay demonstrating a deficiency of N-acetylgalactosamine 4-sulfatase (arylsulfatase B) enzyme activity or by genetic testing. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | NATPARA  |
| <b>Drug Names</b>                   | NATPARA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | Acute postsurgical hypoparathyroidism (within 6 months of surgery) and expected recovery from hypoparathyroidism.                    |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NERLYNX  |
| <b>Drug Names</b>                   | NERLYNX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent human epidermal growth factor receptor 2 (HER2)-positive breast cancer, brain metastases from HER2-positive breast cancer. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | NEXAVAR  |
| <b>Drug Names</b>                   | NEXAVAR, SORAFENIB TOSYLATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Acute myeloid leukemia, soft tissue sarcoma (angiosarcoma, desmoid tumors/aggressive fibromatosis, and solitary fibrous tumor subtypes), gastrointestinal stromal tumor, medullary thyroid carcinoma, osteosarcoma, recurrent chordoma, epithelial ovarian cancer, fallopian tube cancer, primary peritoneal cancer, lymphoid, myeloid, or mixed lineage neoplasms with eosinophilia   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acute myeloid leukemia: the disease is FMS-like tyrosine kinase 3-internal tandem duplication (FLT3-ITD) mutation-positive AND either of the following is met (1 OR 2): 1) the requested drug will be used as maintenance therapy after hematopoietic stem cell transplant, OR 2) the requested drug is used in combination with azacitidine or decitabine for low-intensity treatment induction or post-induction therapy AND either a) the patient has a physiologic age of 60 years of age or older or b) the disease is relapsed/refractory. For thyroid carcinoma: histology is follicular, papillary, Hurthle cell or medullary. For gastrointestinal stromal tumor (GIST): the disease is unresectable, recurrent, or metastatic AND the patient has failed on an FDA-approved therapy (e.g., imatinib, sunitinib, regorafenib, ripretinib). For renal cell carcinoma: the patient meets ALL of the following: 1) The disease is advanced, AND 2) The patient has experienced disease progression or an intolerable adverse event with a trial of cabozantinib or axitinib. For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia: 1) The disease has a FLT3 rearrangement AND 2) The disease is in chronic or blast phase. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NINLARO  |
| <b>Drug Names</b>                   | NINLARO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Relapsed/refractory systemic light chain amyloidosis, Waldenstrom macroglobulinemia, lymphoplasmacytic lymphoma  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | NITISINONE   |
| <b>Drug Names</b>                   | NITISINONE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hereditary tyrosinemia type 1 (HT-1): Diagnosis of HT-1 is confirmed by one of the following: 1) biochemical testing (e.g., detection of succinylacetone in urine) or 2) DNA testing (mutation analysis).  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NORTHERA   |
| <b>Drug Names</b>                   | DROXIDOPA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For neurogenic orthostatic hypotension (nOH): Prior to initial therapy, patient has a persistent, consistent decrease in systolic blood pressure of at least 20 mmHg OR decrease in diastolic blood pressure of at least 10 mmHg within 3 minutes of standing or head-up tilt test. For continuation of therapy for nOH, patient must experience a sustained reduction in symptoms of nOH (i.e., decrease in dizziness, lightheadedness, or feeling faint). For both initial and continuation of therapy for nOH, the requested drug will be used for patients with neurogenic orthostatic hypotension associated with one of the following diagnoses: 1) primary autonomic failure due to Parkinson's disease, multiple system atrophy, or pure autonomic failure, OR 2) dopamine beta-hydroxylase deficiency, OR 3) non-diabetic autonomic neuropathy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 3 months   |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | NOXAFIL SUSP   |
| <b>Drug Names</b>                   | NOXAFIL  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used orally. For treatment of oropharyngeal candidiasis: patient has experienced an inadequate treatment response, intolerance, or has a contraindication to fluconazole. |
| <b>Age Restrictions</b>             | 13 years of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Oropharyngeal candidiasis: 1 month. All other indications: 6 months  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NUBEQA   |
| <b>Drug Names</b>                   | NUBEQA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NUEDEXTA   |
| <b>Drug Names</b>                   | NUEDEXTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | NUPLAZID   |
| <b>Drug Names</b>                   | NUPLAZID   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hallucinations and delusions associated with Parkinson's disease psychosis, the diagnosis of Parkinson's disease must be made prior to the onset of psychotic symptoms.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | NURTEC   |
| <b>Drug Names</b>                   | NURTEC   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Acute migraine treatment: The patient has experienced an inadequate treatment response, intolerance, or the patient has a contraindication to one triptan 5-HT1 receptor agonist . Preventive treatment of migraine, initial: The patient meets either of the following: 1) The patient experienced an inadequate treatment response with a 4-week trial of any one of the following: antiepileptic drugs (AEDs), beta-adrenergic blocking agents, antidepressants OR 2) The patient experienced an intolerance or has a contraindication that would prohibit a 4-week trial of any one of the following: antiepileptic drugs (AEDs), beta-adrenergic blocking agents, antidepressants. Preventive treatment of migraine, continuation: The patient received at least 3 months of treatment with the requested drug, and the patient had a reduction in migraine days per month from baseline. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Preventive treatment of migraine - initial: 3 months, All other indications: Plan Year   |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | OCTREOTIDE   |
| <b>Drug Names</b>                   | OCTREOTIDE ACETATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Tumor control of thymomas and thymic carcinomas.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acromegaly (initial): 1) patient has a high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range, and 2) patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy. For acromegaly continuation of therapy: patient's IGF-1 level has decreased or normalized since initiation of therapy. For tumor control of thymomas and thymic carcinomas, the requested drug will be used for any of the following: 1) locally advanced or metastatic disease, 2) postoperatively following tumor resection. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ODOMZO   |
| <b>Drug Names</b>                   | ODOMZO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | OFEV   |
| <b>Drug Names</b>                   | OFEV   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | OGIVRI  |
| <b>Drug Names</b>                   | OGIVRI  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The patient had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year   |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.  |
| <b>Prior Authorization Group</b>    | OMNIPOD   |
| <b>Drug Names</b>                   | OMNIPOD 5 G6 INTRO KIT (G, OMNIPOD 5 G6 PODS (GEN 5), OMNIPOD CLASSIC PDM START, OMNIPOD CLASSIC PODS (GEN, OMNIPOD DASH INTRO KIT (G, OMNIPOD DASH PODS (GEN 4)  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Initial: 1) The patient has diabetes requiring insulin management with multiple daily injections AND 2) The patient is self-testing glucose levels 4 or more times per day OR the patient is using a continuous glucose monitor AND 3) The patient has experienced any of the following with the current diabetes regimen: inadequate glycemic control, recurrent hypoglycemia, wide fluctuations in blood glucose, dawn phenomenon with persistent severe early morning hyperglycemia, severe glycemic excursions.<br>Continuation: the patient has stable or improved glycemic control. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ONTRUZANT  |
| <b>Drug Names</b>                   | ONTRUZANT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient had an intolerable adverse event to Trazimera and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months, Other: Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |
| <b>Prior Authorization Group</b>    | ONUREG   |
| <b>Drug Names</b>                   | ONUREG   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | OPSUMIT  |
| <b>Drug Names</b>                   | OPSUMIT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1): Diagnosis was confirmed by right heart catheterization. For PAH new starts only: 1) Pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) Pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) Pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ORAL-INTRANASAL FENTANYL  |
| <b>Drug Names</b>                   | FENTANYL CITRATE ORAL TRA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The requested drug is indicated for the treatment of breakthrough CANCER-related pain only. The requested drug is being prescribed for the management of breakthrough pain in a CANCER patient with underlying CANCER pain AND 2) The International Classification of Diseases (ICD) diagnosis code provided supports the CANCER-RELATED diagnosis. [Note: For drug coverage approval, ICD diagnosis code provided MUST support the CANCER-RELATED diagnosis.] AND 3) The patient is currently receiving, and will continue to receive, around-the-clock opioid therapy for underlying CANCER pain AND 4) The requested drug is intended only for use in opioid tolerant patients. The patient can safely take the requested dose based on their current opioid use history. [Note: Patients considered opioid tolerant are those who are taking around-the-clock medicine consisting of at least 60 mg of oral morphine per day, at least 25 mcg per hour of transdermal fentanyl, at least 30 mg of oral oxycodone per day, at least 60 mg of oral hydrocodone per day, at least 8 mg of oral hydromorphone per day, at least 25 mg of oral oxymorphone per day, or an equianalgesic dose of another opioid medication daily for one week or longer.]. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ORGOVYX  |
| <b>Drug Names</b>                   | ORGOVYX  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | ORKAMBI  |
| <b>Drug Names</b>                   | ORKAMBI  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis (CF): The requested medication will not be used in combination with other medications containing ivacaftor.  |
| <b>Age Restrictions</b>             | 1 year of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | OTEZLA   |
| <b>Drug Names</b>                   | OTEZLA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For plaque psoriasis (new starts only): Patient meets either of the following: 1) Inadequate treatment response or intolerance to ANY of the following: a) a topical therapy (e.g., topical corticosteroids, calcineurin inhibitors, vitamin D analogs), b) phototherapy (e.g., UVB, PUVA), or c) pharmacologic treatment with methotrexate, cyclosporine, or acitretin, OR 2) pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | OXANDROLONE  |
| <b>Drug Names</b>                   | OXANDROLONE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications                                |
| <b>Off-label Uses</b>               | Cachexia associated with AIDS (HIV wasting), To enhance growth in patients with Turners Syndrome |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Turners Syndrome: Plan Year, All other diagnoses: 6 months                                       |
| <b>Other Criteria</b>               | Coverage will be denied if request is for an indication excluded from Medicare Part D.           |

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| <b>Prior Authorization Group</b>    | PANRETIN  |
| <b>Drug Names</b>                   | PANRETIN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications                       |
| <b>Off-label Uses</b>               | Topical treatment of cutaneous lesions in patients with non-AIDS-related Kaposi sarcoma |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | PAROXETINE SUSP              |
| <b>Drug Names</b>                   | PAROXETINE HYDROCHLORIDE     |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

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| <b>Prior Authorization Group</b>    | PEGASYS   |
| <b>Drug Names</b>                   | PEGASYS   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Myeloproliferative neoplasm (essential thrombocythemia, polycythemia vera, symptomatic lower-risk myelofibrosis), systemic mastocytosis, adult T-cell leukemia/lymphoma, mycosis fungoides/sezary syndrome, primary cutaneous CD30+ T-cell lymphoproliferative disorders, hairy cell leukemia, Erdheim-Chester disease. |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For chronic hepatitis C: Hepatitis C virus (HCV) confirmed by presence of hepatitis C virus HCV RNA in serum prior to starting treatment and the planned treatment regimen.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | HCV: 12-48wks. Criteria applied consistent w/current AASLD/IDSA guidance. HBV: 48wks. Other: Plan Yr  |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | PEMAZYRE  |
| <b>Drug Names</b>                   | PEMAZYRE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FGFR1 rearrangement: the disease is in chronic or blast phase.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | PHENYLBUTYRATE  |
| <b>Drug Names</b>                   | SODIUM PHENYLBUTYRATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For urea cycle disorders (UCD): Diagnosis of UCD was confirmed by enzymatic, biochemical or genetic testing.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | PHESGO   |
| <b>Drug Names</b>                   | PHESGO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent human epidermal growth factor receptor 2 (HER2)-positive breast cancer   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months. Other: Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | PIQRAY   |
| <b>Drug Names</b>                   | PIQRAY 200MG DAILY DOSE, PIQRAY 250MG DAILY DOSE, PIQRAY 300MG DAILY DOSE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative, PIK3CA-mutated breast cancer in combination with fulvestrant.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | POMALYST   |
| <b>Drug Names</b>                   | POMALYST   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Relapsed/refractory systemic light chain amyloidosis, primary central nervous system (CNS) lymphoma, POEMS syndrome.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For multiple myeloma: The patient has previously received at least two prior therapies for multiple myeloma, including an immunomodulatory agent AND a proteasome inhibitor. For Kaposi sarcoma, patient meets one of the following: 1) patient has acquired immunodeficiency syndrome (AIDS), or 2) patient is negative for human immunodeficiency virus (HIV). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | POSACONAZOLE   |
| <b>Drug Names</b>                   | POSACONAZOLE DR  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used orally. For prophylaxis of invasive Aspergillus and Candida infections: patient weighs greater than 40 kilograms.  |
| <b>Age Restrictions</b>             | Treatment of Invasive Aspergillosis: 13 years of age or older, Prophylaxis of Invasive Aspergillus and Candida Infections: 2 years of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 6 months   |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | PRALUENT   |
| <b>Drug Names</b>                   | PRALUENT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | PREGABALIN   |
| <b>Drug Names</b>                   | PREGABALIN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Cancer-related neuropathic pain, cancer treatment-related neuropathic pain   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For the management of postherpetic neuralgia, the management of neuropathic pain associated with diabetic peripheral neuropathy, cancer-related neuropathic pain, and cancer treatment-related neuropathic pain: The patient has experienced an inadequate treatment response, intolerance, or contraindication to gabapentin. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | PREVYMIS  |
| <b>Drug Names</b>                   | PREVYMIS  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For prophylaxis of cytomegalovirus (CMV) infection or disease: 1) the patient is CMV-seropositive, 2) the patient is a recipient of an allogeneic hematopoietic stem cell transplant (HSCT), AND 3) the requested drug will not be used beyond day 100 post-transplantation.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 6 months  |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | PROCRIT   |
| <b>Drug Names</b>                   | PROCRIT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Anemia due to myelodysplastic syndromes (MDS), anemia in congestive heart failure (CHF), anemia in rheumatoid arthritis (RA), anemia due to hepatitis C treatment (ribavirin in combination with either interferon alfa or peginterferon alfa)  |
| <b>Exclusion Criteria</b>           | Patients receiving chemotherapy with curative intent. Patients with myeloid cancer.   |
| <b>Required Medical Information</b> | Requirements regarding hemoglobin (Hgb) values exclude values due to a recent transfusion. For initial approval: 1) for all uses except anemia due to chemotherapy or myelodysplastic syndrome (MDS): patient has adequate iron stores (for example, a transferrin saturation [TSAT] greater than or equal to 20%), AND 2) for all uses except surgery: pretreatment (no erythropoietin treatment in previous month) Hgb is less than 10 g/dL (less than 9 g/dL for anemia in congestive heart failure), AND 3) for MDS: pretreatment serum erythropoietin level is 500 international units/L or less. For reauthorizations (patient received erythropoietin treatment in previous month) in all uses except surgery: 1) patient has received at least 12 weeks of erythropoietin therapy, AND 2) patient responded to erythropoietin therapy, AND 3) current Hgb is less than 12 g/dL, AND 4) for all uses except anemia due to chemotherapy or MDS: patient has adequate iron stores (for example, a transferrin saturation [TSAT] greater than or equal to 20%). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 16 weeks  |
| <b>Other Criteria</b>               | Coverage includes use in anemia in patients whose religious beliefs forbid blood transfusions. Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual (e.g., used for treatment of anemia for a patient with chronic renal failure who is undergoing dialysis, or furnished from physician's supply incident to a physician service).  |

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| <b>Prior Authorization Group</b>    | PROMACTA   |
| <b>Drug Names</b>                   | PROMACTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic or persistent immune thrombocytopenia (ITP): 1) For new starts: a) Patient (pt) has had an inadequate response or is intolerant to a prior therapy such as corticosteroids or immunoglobulins, b) Untransfused platelet (plt) count at any point prior to the initiation of the requested medication is less than 30,000/mcL OR 30,000-50,000/mcL with symptomatic bleeding or risk factor(s) for bleeding (e.g., undergoing a medical or dental procedure where blood loss is anticipated, comorbidities such as peptic ulcer disease and hypertension, anticoagulation therapy, profession or lifestyle that predisposes pt to trauma) AND c) For chronic ITP only: pt has had an inadequate response or intolerance to avatrombopag. 2) For continuation of therapy, plt count response to the requested drug: a) Current plt count is less than or equal to 200,000/mcL OR b) Current plt count is greater than 200,000/mcL to less than or equal to 400,000/mcL and dosing will be adjusted to a plt count sufficient to avoid clinically important bleeding. For thrombocytopenia associated with chronic hepatitis C: 1) For new starts: the requested drug is used for initiation and maintenance of interferon-based therapy. 2) For continuation of therapy: pt is receiving interferon-based therapy. For severe aplastic anemia (AA): 1) For new starts: a) Pt will use the requested drug with standard immunosuppressive therapy for first line treatment OR b) the pt had an insufficient response to immunosuppressive therapy. 2) For continuation of therapy : 1) Current plt count is 50,000-200,000/mcL OR 2) Current plt count is less than 50,000/mcL and pt has not received appropriately titrated therapy for at least 16 weeks, OR 3) Current plt count is less than 50,000/mcL and pt is transfusion-independent, OR 4) Current plt count is greater than 200,000/mcL to less than or equal to 400,000/mcL and dosing will be adjusted to achieve and maintain an appropriate target plt count. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | HCV: 6mo, ITP/AA initial: 6mo, ITP reauth: Plan Year, AA reauth: APR-Plan Year, IPR-16 wks   |
| <b>Other Criteria</b>               | APR: adequate platelet response (greater than 50,000/mcL), IPR: inadequate platelet response (less than 50,000/mcL).   |

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| <b>Prior Authorization Group</b>    | PULMOZYME  |
| <b>Drug Names</b>                   | PULMOZYME  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis: Diagnosis of cystic fibrosis was confirmed by appropriate diagnostic or genetic testing.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual. |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | QINLOCK  |
| <b>Drug Names</b>                   | QINLOCK  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | QUETIAPINE XR  |
| <b>Drug Names</b>                   | QUETIAPINE FUMARATE ER   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Maintenance monotherapy treatment in bipolar I disorder, monotherapy treatment of generalized anxiety disorder, monotherapy treatment of major depressive disorder   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For schizophrenia, acute treatment of manic or mixed episodes associated with bipolar I disorder, both as monotherapy and as an adjunct to lithium or divalproex, the acute treatment of depressive episodes associated with bipolar disorder, maintenance treatment of bipolar I disorder, as an adjunct to lithium or divalproex, adjunctive treatment of major depressive disorder, or maintenance monotherapy treatment in bipolar I disorder: The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following generic products: A) aripiprazole, B) asenapine, C) olanzapine, D) quetiapine immediate-release, E) risperidone, F) ziprasidone. For all indications: If the patient is 65 years of age or older AND is using two or more additional central nervous system (CNS) active medications (e.g., lorazepam, sertraline, clonazepam, escitalopram, alprazolam, zolpidem) with the requested drug, the prescriber determined that taking multiple central nervous system (CNS) active medications is medically necessary. [Note: Use of multiple central nervous system (CNS) active medications in older adults is associated with an increased risk of falls.]. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | QUININE SULFATE  |
| <b>Drug Names</b>                   | QUININE SULFATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Babesiosis, uncomplicated Plasmodium vivax malaria.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 1 month  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | REGRANEX                     |
| <b>Drug Names</b>                   | REGRANEX                     |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | 20 weeks                     |
| <b>Other Criteria</b>               | -                            |

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| <b>Prior Authorization Group</b>    | RELISTOR INJ  |
| <b>Drug Names</b>                   | RELISTOR  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For the treatment of opioid-induced constipation in a patient with chronic non-cancer pain, including chronic pain related to prior cancer or its treatment who does not require frequent (e.g., weekly) opioid dosage escalation: 1) the patient is unable to tolerate oral medications OR 2) the patient meets one of the following criteria A) experienced an inadequate treatment response or intolerance to an oral drug indicated for opioid-induced constipation in a patient with chronic non-cancer pain (e.g., Movantik) OR B) the patient has a contraindication that would prohibit a trial of an oral drug indicated for opioid-induced constipation in a patient with chronic non-cancer pain (e.g., Movantik). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 4 months  |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | REMICADE   |
| <b>Drug Names</b>                   | INFLIXIMAB, REMICADE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Behcet's syndrome, granulomatosis with polyangiitis (Wegener's granulomatosis), hidradenitis suppurativa, juvenile idiopathic arthritis, pyoderma gangrenosum, sarcoidosis, Takayasu's arteritis, uveitis.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For moderately to severely active Crohn's disease (new starts only): 1) Pt has fistulizing disease, OR 2) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 3) Intolerance or contraindication (CI) to conventional therapy. For moderately to severely active ulcerative colitis (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids) OR 2) Intolerance or CI to conventional therapy. For moderately to severely active rheumatoid arthritis (new starts only): 1) Pt meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) or leflunomide OR b) intolerance or CI to MTX AND leflunomide AND 2) Pt meets ANY of the following: a) inadequate response, intolerance or CI to MTX OR b) inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis (new starts only): Inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial OR intolerance or CI to NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at time of diagnosis, AND 2) Pt meets ANY of the following: a) pt has experienced inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with MTX, cyclosporine, or acitretin, OR b) pharmacologic treatment with MTX, cyclosporine, or acitretin is contraindicated, OR c) pt has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of BSA or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | For hidradenitis suppurativa (new starts only): Pt has severe, refractory disease. For uveitis (new starts only): Inadequate response or intolerance or has a CI to a trial of immunosuppressive therapy for uveitis. For FDA-approved indications and off-label uses that overlap: The patient had an intolerable adverse event to Renflexis and that adverse event was NOT attributed to the active ingredient as described in the prescribing information.  |

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| <b>Prior Authorization Group</b>    | RENFLEXIS  |
| <b>Drug Names</b>                   | RENFLEXIS  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Behcet's syndrome, granulomatosis with polyangiitis (Wegener's granulomatosis), hidradenitis suppurativa, juvenile idiopathic arthritis, pyoderma gangrenosum, sarcoidosis, Takayasu's arteritis, uveitis  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For moderately to severely active Crohn's disease (new starts only): 1) Pt has fistulizing disease, OR 2) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 3) Intolerance or contraindication (CI) to conventional therapy. For moderately to severely active ulcerative colitis (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids) OR 2) Intolerance or CI to conventional therapy. For moderately to severely active rheumatoid arthritis (new starts only): 1) Pt meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) or leflunomide OR b) intolerance or CI to MTX AND leflunomide AND 2) pt meets ANY of the following: a) inadequate response, intolerance or CI to MTX OR b) inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. For active ankylosing spondylitis (new starts only): Inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial OR intolerance or CI to NSAIDs. For moderate to severe plaque psoriasis (new starts only): 1) At least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at time of diagnosis, AND 2) Pt meets ANY of the following: a) pt has experienced inadequate response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with MTX, cyclosporine, or acitretin, OR b) pharmacologic treatment with MTX, cyclosporine, or acitretin is contraindicated, OR c) pt has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of BSA or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | For hidradenitis suppurativa (new starts only): pt has severe, refractory disease. For uveitis (new starts only): Inadequate response or intolerance or has a CI to a trial of immunosuppressive therapy for uveitis.  |

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| <b>Prior Authorization Group</b>    | RETEVMO   |
| <b>Drug Names</b>                   | RETEVMO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent rearranged during transfection (RET)-rearrangement positive non-small cell lung cancer, Langerhans Cell Histiocytosis with a RET gene fusion, symptomatic or relapsed/refractory Erdheim-Chester Disease with a RET gene fusion, symptomatic or relapsed/refractory Rosai-Dorfman Disease with a RET gene fusion.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For non-small cell lung cancer, patient must meet all of the following: 1) The disease is recurrent, advanced or metastatic, and 2) Tumor is RET fusion-positive or RET rearrangement-positive.   |
| <b>Age Restrictions</b>             | Non-small cell lung cancer: 18 years of age or older. Medullary thyroid cancer and thyroid cancer: 12 years of age or older.  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | REVLIMID  |
| <b>Drug Names</b>                   | LENALIDOMIDE, REVLIMID  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Systemic light chain amyloidosis, classical Hodgkin lymphoma, myelodysplastic syndrome without the 5q deletion cytogenetic abnormality, myelofibrosis-associated anemia, POEMS syndrome, myeloproliferative neoplasms, Kaposi Sarcoma, Langerhans cell histiocytosis, peripheral T-Cell lymphomas not otherwise specified, angioimmunoblastic T-cell lymphoma (AITL), enteropathy-associated T-cell lymphoma, monomorphic epitheliotropic intestinal T-cell lymphoma, nodal peripheral T-cell lymphoma, adult T-cell leukemia/lymphoma, hepatosplenic T-cell lymphoma, primary central nervous system (CNS) lymphoma, chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL), acquired immunodeficiency syndrome (AIDS)-related non-germinal center diffuse large B-cell lymphoma, monomorphic post-transplant lymphoproliferative disorder, diffuse large B-cell lymphoma, multicentric Castlemans disease, high-grade B-cell lymphomas, histologic transformation of nodal marginal zone lymphoma to diffuse large B-cell lymphoma, histologic transformation of follicular lymphoma to diffuse large B-cell lymphoma. |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For myelodysplastic syndrome (MDS): Lower risk MDS with symptomatic anemia per the Revised International Prognostic Scoring System (IPSS-R), International Prognostic Scoring System (IPSS), or World Health organization (WHO) classification-based Prognostic Scoring System (WPSS).  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | REZUROCK                     |
| <b>Drug Names</b>                   | REZUROCK                     |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | 12 years of age or older     |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

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| <b>Prior Authorization Group</b>    | RINVOQ  |
| <b>Drug Names</b>                   | RINVOQ  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderately to severely active rheumatoid arthritis (new starts only): patient has experienced an inadequate treatment response, intolerance or has a contraindication to methotrexate (MTX) AND at least one tumor necrosis factor (TNF) inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). For active psoriatic arthritis (new starts only): patient has experienced an inadequate treatment response, intolerance or has a contraindication to at least one TNF inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). For moderately to severely active ulcerative colitis (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one TNF inhibitor (e.g., Humira [adalimumab]). For atopic dermatitis, continuation of therapy: the patient achieved or maintained positive clinical response. For active ankylosing spondylitis (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one TNF inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). For non-radiographic axial spondyloarthritis (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one TNF inhibitor. |
| <b>Age Restrictions</b>             | Atopic dermatitis: 12 years of age or older   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Atopic dermatitis (initial): 4 months, All others: Plan Year  |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ROZLYTREK   |
| <b>Drug Names</b>                   | ROZLYTREK   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent ROS1-positive non-small cell lung cancer (NSCLC), Non-metastatic neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, first-line treatment of NTRK gene fusion-positive solid tumors.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For all neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, the disease is without a known acquired resistance mutation. For ROS1-positive non-small cell lung cancer, the patient has recurrent, advanced, or metastatic disease.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | RUBRACA   |
| <b>Drug Names</b>                   | RUBRACA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Uterine leiomyosarcoma, advanced (stage II-IV) epithelial ovarian, fallopian tube, or primary peritoneal cancer   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For metastatic castration-resistant prostate cancer with a deleterious breast cancer susceptibility gene (BRCA) mutation (germline and/or somatic): 1) patient has been treated with androgen receptor-directed therapy, 2) patient has been treated with a taxane-based chemotherapy or the patient is not fit for chemotherapy, 3) the requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy. For maintenance treatment of BRCA mutated epithelial ovarian, fallopian tube, primary peritoneal cancer: 1) the patient has advanced (stage II-IV) disease and is in complete or partial response to primary therapy or 2) the patient has recurrent disease and is in complete or partial response to platinum-based chemotherapy. For uterine leiomyosarcoma: 1) the requested drug is used as second-line therapy AND 2) the patient has BRCA-altered disease. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | RYDAPT   |
| <b>Drug Names</b>                   | RYDAPT   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Relapsed or refractory acute myeloid leukemia (AML), myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FGFR1 or FLT3 rearrangements, post-induction therapy for AML, re-induction in residual disease for AML  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acute myeloid leukemia (AML): AML is FLT3 mutation-positive. For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FGFR1 or FLT3 rearrangements: the disease is in chronic or blast phase.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | SAPROPTERIN  |
| <b>Drug Names</b>                   | JAVYGTOR, SAPROPTERIN DIHYDROCHLORI  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For phenylketonuria (PKU): For patients who have not yet received a therapeutic trial of the requested drug, the patient's pretreatment, including before dietary management, phenylalanine level is greater than 6 mg/dL (360 micromol/L). For patients who completed a therapeutic trial of the requested drug, the patient must have experienced improvement (e.g., reduction in blood phenylalanine levels, improvement in neuropsychiatric symptoms). |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Initial: 2 months. All others: Plan Year.  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | SCEMBLIX   |
| <b>Drug Names</b>                   | SCEMBLIX   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML) in the chronic phase: 1) the diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene AND the patient meets either of the following: A) the patient has previously been treated with 2 or more tyrosine kinase inhibitors (TKIs) AND at least one of those was imatinib or dasatinib, OR B) the patient is positive for the T315I mutation.                        |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | SIGNIFOR   |
| <b>Drug Names</b>                   | SIGNIFOR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an endocrinologist   |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | SILDENAFIL   |
| <b>Drug Names</b>                   | SILDENAFIL CITRATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (PAH) (World Health Organization [WHO] Group 1): Diagnosis was confirmed by right heart catheterization. For PAH new starts only: 1) Pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) Pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, and 3) Pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | SIRTURO   |
| <b>Drug Names</b>                   | SIRTURO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with an infectious disease specialist. |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | SKYRIZI   |
| <b>Drug Names</b>                   | SKYRIZI, SKYRIZI PEN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderate to severe plaque psoriasis (new starts only): 1) at least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) patient meets any of the following: a) patient has experienced an inadequate treatment response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, or b) pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, or c) patient has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of the body surface area or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected). For moderately to severely active Crohn's disease (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 2) Intolerance or contraindication to conventional therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | SKYRIZI-CD  |
| <b>Drug Names</b>                   | SKYRIZI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderate to severe plaque psoriasis (new starts only): 1) at least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis, AND 2) patient meets any of the following: a) patient has experienced an inadequate treatment response or intolerance to either phototherapy (e.g., UVB, PUVA) or pharmacologic treatment with methotrexate, cyclosporine, or acitretin, or b) pharmacologic treatment with methotrexate, cyclosporine, or acitretin is contraindicated, or c) patient has severe psoriasis that warrants a biologic as first-line therapy (i.e., at least 10% of the body surface area or crucial body areas [e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas] are affected). For moderately to severely active Crohn's disease (new starts only): 1) Inadequate response to at least one conventional therapy (e.g., corticosteroids), OR 2) Intolerance or contraindication to conventional therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | SOMATULINE DEPOT  |
| <b>Drug Names</b>                   | SOMATULINE DEPOT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Tumor control of neuroendocrine tumors (NETs) of the lung, thymus or unresected primary gastrinoma, well-differentiated grade 3 neuroendocrine tumors not of gastroenteropancreatic origin, pheochromocytoma/paraganglioma.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For acromegaly (initial): 1) patient has a high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range, and 2) patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy. For acromegaly continuation of therapy: patient's IGF-1 level has decreased or normalized since initiation of therapy. For tumor control of neuroendocrine tumors (NETs) of the thymus or lung: patient has locoregional unresectable disease and/or distant metastatic disease. For tumor control of well-differentiated grade 3 unresectable locally advanced or metastatic NETs (not of gastroenteropancreatic origin): patient has favorable biology (e.g., relatively low Ki-67 [less than 55%] and somatostatin receptor [SSR] positive imaging). For tumor control of pheochromocytomas or paragangliomas: 1) patient has symptomatic locally unresectable disease with SSR positive imaging or 2) patient has distant metastases that are secreting tumors. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | SOMAVERT  |
| <b>Drug Names</b>                   | SOMAVERT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For acromegaly (initial): 1) patient has a high pretreatment insulin-like growth factor-1 (IGF-1) level for age and/or gender based on the laboratory reference range, and 2) patient had an inadequate or partial response to surgery or radiotherapy OR there is a clinical reason for why the patient has not had surgery or radiotherapy. For acromegaly (continuation of therapy): patient's IGF-1 level has decreased or normalized since initiation of therapy.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | SPRYCEL  |
| <b>Drug Names</b>                   | SPRYCEL  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Gastrointestinal stromal tumor (GIST), metastatic chondrosarcoma, recurrent chordoma, T-cell acute lymphoblastic leukemia (ALL), and Philadelphia (Ph)-like B-ALL, myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in the chronic phase or blast phase  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML), including patients who have received a hematopoietic stem cell transplant: 1) Diagnosis was confirmed by detection of the Philadelphia (Ph) chromosome or BCR-ABL gene, and 2) If patient experienced resistance to an alternative tyrosine kinase inhibitor, patient is negative for all of the following mutations: T315I/A, F317L/V/I/C, and V299L. For acute lymphoblastic leukemia (ALL), the patient has a diagnosis of one of the following: 1) Philadelphia chromosome positive ALL, including patients who have received a hematopoietic stem cell transplant: diagnosis that has been confirmed by detection of the Ph chromosome or BCR-ABL gene, and if patient experienced resistance to an alternative tyrosine kinase inhibitor, patient is negative for all of the following mutations: T315I/A, F317L/V/I/C, and V299L, OR 2) Ph-like B-ALL with ABL-class kinase fusion, OR 3) relapsed or refractory T-cell ALL with ABL-class translocation. For GIST, 1) the disease has progressed on imatinib in patients with PDGFRA D842V mutation, OR 2) the patient has failed at least 2 FDA-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib) |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | STIVARGA   |
| <b>Drug Names</b>                   | STIVARGA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Progressive gastrointestinal stromal tumors (GIST), osteosarcoma, glioblastoma, angiosarcoma, retroperitoneal/intra-abdominal soft tissue sarcoma, rhabdomyosarcoma, and soft tissue sarcomas of the extremities, body wall, head and neck, advanced colorectal cancer.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For gastrointestinal stromal tumors: The disease is progressive, locally advanced, unresectable, or metastatic.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | SUTENT   |
| <b>Drug Names</b>                   | SUNITINIB MALATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Thyroid carcinoma (follicular, medullary, papillary, and Hurthle cell), soft tissue sarcoma (angiosarcoma, solitary fibrous tumor, and alveolar soft part sarcoma subtypes), recurrent chordoma, thymic carcinoma, lymphoid, myeloid, or mixed lineage neoplasms with eosinophilia   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For renal cell carcinoma (RCC): 1) The disease is relapsed, advanced, or stage IV OR 2) the requested drug is being used as adjuvant treatment for patients that are at high risk of recurrent RCC following nephrectomy. For gastrointestinal stromal tumor (GIST): the disease is unresectable, recurrent, or metastatic AND the patient has failed on an FDA-approved therapy (e.g., imatinib, sunitinib, regorafenib, ripretinib). For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia: 1) The disease has a FLT3 rearrangement AND 2) The disease is in chronic or blast phase. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | SYMDEKO  |
| <b>Drug Names</b>                   | SYMDEKO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis: The requested medication will not be used in combination with other medications containing ivacaftor.   |
| <b>Age Restrictions</b>             | 6 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | SYMPAZAN   |
| <b>Drug Names</b>                   | SYMPAZAN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | 2 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

**Prior Authorization Group** SYNRIBO  
**Drug Names** SYNRIBO  
**PA Indication Indicator** All FDA-approved Indications  
**Off-label Uses** -  
**Exclusion Criteria** -  
**Required Medical Information** -  
**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

**Prior Authorization Group** TABRECTA  
**Drug Names** TABRECTA  
**PA Indication Indicator** All FDA-approved Indications, Some Medically-accepted Indications  
**Off-label Uses** Recurrent non-small cell lung cancer (NSCLC).  
**Exclusion Criteria** -  
**Required Medical Information** For recurrent, advanced, or metastatic NSCLC: Tumor is positive for mesenchymal-epithelial transition (MET) exon 14 skipping mutation.  
**Age Restrictions** -  
**Prescriber Restrictions** -  
**Coverage Duration** Plan Year  
**Other Criteria** -

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| <b>Prior Authorization Group</b>    | TAFINLAR   |
| <b>Drug Names</b>                   | TAFINLAR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Brain metastases from melanoma, thyroid carcinoma (papillary carcinoma, follicular carcinoma, and Hurthle cell carcinoma), central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma), gallbladder cancer, extrahepatic cholangiocarcinoma, intrahepatic cholangiocarcinoma, Langerhans cell histiocytosis, Erdheim-Chester disease   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For brain metastases from melanoma, adjuvant treatment of melanoma, and central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma): 1) The tumor is positive for a BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used in combination with trametinib. For unresectable or metastatic melanoma: 1) The tumor is positive for a BRAF V600 activating mutation (e.g., V600E or V600K), and 2) The requested drug will be used as a single agent or in combination with trametinib. For non-small cell lung cancer: 1) The tumor is positive for a BRAF V600E mutation, and 2) The requested drug will be used as a single agent or in combination with trametinib. For thyroid carcinoma with papillary, follicular, or Hurthle histology: The tumor is positive for BRAF activating mutation (e.g., V600E or V600K). For Langerhans Cell Histiocytosis and Erdheim-Chester Disease: The disease is positive for a BRAF V600E mutation. For gallbladder cancer, extrahepatic cholangiocarcinoma, and intrahepatic cholangiocarcinoma: 1) The disease is positive for a BRAF V600E mutation and 2) The disease is unresectable or metastatic and 3) The requested drug will be used in combination with trametinib. For solid tumors: ) The tumor is positive for a BRAF V600E mutation, and 2) The requested drug will be used in combination with trametinib. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | TAGRISSO   |
| <b>Drug Names</b>                   | TAGRISSO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Sensitizing epidermal growth factor receptor (EGFR) mutation-positive recurrent non-small cell lung cancer (NSCLC), brain metastases from sensitizing EGFR mutation-positive NSCLC, leptomeningeal metastases from EGFR mutation-positive NSCLC.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For NSCLC, the requested drug is used in any of the following settings: 1) The patient meets both of the following: a) patient has metastatic, advanced, or recurrent NSCLC (including brain and/or leptomeningeal metastases from NSCLC) and b) patient has a sensitizing EGFR mutation OR 2) The patient meets both of the following: a) request is for adjuvant treatment of NSCLC following tumor resection and b) patient has EGFR mutation-positive disease. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | TALTZ   |
| <b>Drug Names</b>                   | TALTZ   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderate to severe plaque psoriasis (new starts only): 1) at least 3% of body surface area (BSA) is affected OR crucial body areas (e.g., feet, hands, face, neck, groin, intertriginous areas) are affected at the time of diagnosis AND 2) the patient has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following products: Enbrel (etanercept), Humira (adalimumab), Otezla (apremilast), Skyrizi (risankizumab-rzaa). For active ankylosing spondylitis (new starts only): the patient has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following products: Enbrel (etanercept), Humira (adalimumab), Rinvoq (upadacitinib), Xeljanz (tofacitinib)/Xeljanz XR (tofacitinib extended-release). For active psoriatic arthritis (PsA) (new starts only): the patient has experienced an inadequate treatment response, intolerance, or has a contraindication to one of the following products: Enbrel (etanercept), Humira (adalimumab), Otezla (apremilast), Rinvoq (upadacitinib), Skyrizi (risankizumab-rzaa), Xeljanz (tofacitinib)/Xeljanz XR (tofacitinib extended-release). For active non-radiographic axial spondyloarthritis (new starts only): Patient meets any of the following: 1) has had an inadequate response to a non-steroidal anti-inflammatory drug (NSAID) trial or 2) has an intolerance or contraindication to NSAIDs. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | TALZENNA  |
| <b>Drug Names</b>                   | TALZENNA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent germline breast cancer susceptibility gene (BRCA)-mutated breast cancer   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TARGRETIN TOPICAL   |
| <b>Drug Names</b>                   | BEXAROTENE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Stage 2 or higher mycosis fungoides/Sezary syndrome, chronic or smoldering adult T-cell leukemia/lymphoma, primary cutaneous marginal zone lymphoma, primary cutaneous follicle center lymphoma.  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | TASIGNA   |
| <b>Drug Names</b>                   | TASIGNA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), gastrointestinal stromal tumor (GIST), myeloid and/or lymphoid neoplasms with eosinophilia and ABL1 rearrangement in the chronic phase or blast phase.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For chronic myeloid leukemia (CML) or acute lymphoblastic leukemia (ALL), diagnosis was confirmed by detection of the Philadelphia chromosome or BCR-ABL gene. For CML, including patients newly diagnosed with CML and patients who have received a hematopoietic stem cell transplant: patient has experienced resistance or intolerance to imatinib or dasatinib. If patient experienced resistance to an alternative tyrosine kinase inhibitor for CML, patient is negative for T315I, Y253H, E255K/V, and F359V/C/I mutations. For GIST, patient must have progressed on imatinib, sunitinib, and regorafenib. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TAZAROTENE  |
| <b>Drug Names</b>                   | TAZAROTENE, TAZORAC   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For plaque psoriasis: 1)The requested drug is being prescribed to treat less than or equal to 20 percent of the patient's body surface area (BSA) AND 2) the patient experienced an inadequate treatment response or intolerance to at least one topical corticosteroid OR has a contraindication that would prohibit a trial of topical corticosteroids. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | TAZVERIK  |
| <b>Drug Names</b>                   | TAZVERIK  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | Epithelioid sarcoma: 16 years of age or older, Follicular lymphoma: 18 years of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TECENTRIQ  |
| <b>Drug Names</b>                   | TECENTRIQ  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent non-small cell lung cancer, single agent maintenance for extensive small cell lung cancer following combination treatment with etoposide and carboplatin, urothelial carcinoma.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For urothelial carcinoma, patient meets one of the following criteria: 1) Patient is ineligible for cisplatin therapy and tumors express PD-L1 (defined as PD-L1 stained tumor-infiltrating immune cells [IC] covering greater than or equal to 5 percent of the tumor area) OR 2) Patient is ineligible for any platinum containing chemotherapy. For non-small cell lung cancer (NSCLC): 1) the patient has recurrent, advanced or metastatic disease AND the requested drug will be used as any of the following: a) first-line treatment of tumors with high PD-L1 expression (defined as PD-L1 stained greater than or equal to 50 percent of tumor cells or PD-L1 stained tumor-infiltrating immune cells [IC] covering greater than or equal to 10 percent of the tumor area) and no EGFR or ALK genomic tumor aberrations, b) used in combination with carboplatin, paclitaxel, and bevacizumab, or in combination with carboplatin and albumin-bound paclitaxel for nonsquamous NSCLC, or c) the requested drug will be used as subsequent therapy or continuation maintenance therapy, OR 2) the patient has stage II to IIIA disease AND the requested drug will be used as adjuvant treatment following resection and platinum-based chemotherapy for tumors with PD-L1 expression on greater than or equal to 1 percent of tumor cells. For hepatocellular carcinoma, the requested drug will be used as initial treatment in combination with bevacizumab. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | TEMAZEPAM   |
| <b>Drug Names</b>                   | TEMAZEPAM   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For short-term treatment of insomnia: 1) The prescriber must acknowledge that the benefit of therapy with this prescribed medication outweighs the potential risks for the patient. (Note: The American Geriatrics Society identifies the use of this medication as potentially inappropriate in older adults, meaning it is best avoided, prescribed at reduced dosage, or used with caution or carefully monitored.) AND 2) The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to doxepin (3 mg or 6 mg). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | This Prior Authorization only applies to patients 65 years of age or older.   |
| <b>Prior Authorization Group</b>    | TEPMETKO  |
| <b>Drug Names</b>                   | TEPMETKO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent non-small cell lung cancer (NSCLC).   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For recurrent, advanced, or metastatic NSCLC: Tumor is positive for mesenchymal-epithelial transition (MET) exon 14 skipping mutation.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TERIPARATIDE   |
| <b>Drug Names</b>                   | TERIPARATIDE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For postmenopausal osteoporosis: patient has ONE of the following (1 or 2): 1) a history of fragility fracture, OR 2) A pre-treatment T-score of less than or equal to -2.5 or pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment Fracture Risk Assessment Tool (FRAX) fracture probability AND patient has ANY of the following: a) indicators for higher fracture risk (e.g., advanced age, frailty, glucocorticoid therapy, very low T-scores, or increased fall risk), OR b) patient has failed prior treatment with or is intolerant to a previous injectable osteoporosis therapy OR c) patient has had an oral bisphosphonate trial of at least 1-year duration or there is a clinical reason to avoid treatment with an oral bisphosphonate. For primary or hypogonadal osteoporosis in men: patient has one of the following: 1) a history of osteoporotic vertebral or hip fracture, OR 2) pre-treatment T-score of less than or equal to -2.5, OR 3) pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment FRAX fracture probability. For glucocorticoid-induced osteoporosis: Patient has had an oral bisphosphonate trial of at least 1-year duration unless patient has a contraindication or intolerance to an oral bisphosphonate, AND patient meets ANY of the following: 1) patient has a history of fragility fracture, OR 2) a pre-treatment T-score of less than or equal to -2.5, OR 3) pre-treatment T-score greater than -2.5 and less than -1 with a high pre-treatment FRAX fracture probability. Continuation of therapy: If the patient has received greater than or equal to 24 months of therapy with any parathyroid hormone analog: 1) The patient remains at or has returned to having a high risk for fracture, AND 2) The benefit of therapy with this prescribed medication outweighs the potential risks for this patient. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Initial: 24 months, Continuation: Plan Year  |
| <b>Other Criteria</b>               | Patient has high FRAX fracture probability if the 10 year probability is either greater than or equal to 20 percent for any major osteoporotic fracture or greater than or equal to 3 percent for hip fracture. If glucocorticoid treatment is greater than 7.5 mg (prednisone equivalent) per day, the estimated risk score generated with FRAX should be multiplied by 1.15 for major osteoporotic fracture and 1.2 for hip fracture.  |

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| <b>Prior Authorization Group</b>    | TESTOSTERONE CYPIONATE INJ  |
| <b>Drug Names</b>                   | TESTOSTERONE CYPIONATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Gender Dysphoria  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Primary or hypogonadotropic hypogonadism: 1) Request is for continuation of testosterone therapy and the patient had a confirmed low morning testosterone level according to current practice guidelines or your standard lab reference values before starting testosterone therapy [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.] OR 2) Request is not for continuation of testosterone therapy and the patient has at least two confirmed low morning testosterone levels according to current practice guidelines or your standard lab reference values [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.]. Gender dysphoria: The patient is able to make an informed decision to engage in hormone therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TESTOSTERONE ENANTHATE INJ  |
| <b>Drug Names</b>                   | TESTOSTERONE ENANTHATE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Gender Dysphoria  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Primary or hypogonadotropic hypogonadism: 1) Request is for continuation of testosterone therapy and the patient had a confirmed low morning testosterone level according to current practice guidelines or your standard lab reference values before starting testosterone therapy [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.] OR 2) Request is not for continuation of testosterone therapy and the patient has at least two confirmed low morning testosterone levels according to current practice guidelines or your standard lab reference values [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.]. Gender dysphoria: The patient is able to make an informed decision to engage in hormone therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | TETRABENAZINE  |
| <b>Drug Names</b>                   | TETRABENAZINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Tic disorders, tardive dyskinesia, hemiballismus, chorea not associated with Huntington's disease.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For treatment of chorea associated with Huntington's disease: The patient must have a prior inadequate response or intolerable adverse event with deutetrabenazine therapy.<br>For treatment of tardive dyskinesia: The patient must have a prior inadequate response or intolerable adverse event with deutetrabenazine or valbenazine therapy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | TETRACYCLINE   |
| <b>Drug Names</b>                   | TETRACYCLINE HYDROCHLORID  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The patient will use the requested drug orally.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | THALOMID   |
| <b>Drug Names</b>                   | THALOMID   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Myelofibrosis-related anemia, AIDS-related aphthous stomatitis, Kaposi sarcoma, chronic graft-versus-host disease, Crohn's disease, multicentric Castleman's disease., Rosai-Dorfman disease, Langerhans cell histiocytosis  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |



|                                     |  |
|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | TIBSOVO  |
| <b>Drug Names</b>                   | TIBSOVO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Conventional (grades 1-3) or dedifferentiated chondrosarcoma. Newly-diagnosed acute myeloid leukemia (AML) if 60-74 years of age and without comorbidities.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Patient has disease with a susceptible isocitrate dehydrogenase-1 (IDH1) mutation. For acute myeloid leukemia (AML): 1) patient has newly-diagnosed AML and meets one of the following: a) 75 years of age or older, b) patient has comorbidities that preclude use of intensive induction chemotherapy, or c) patient is 60 years of age or older and declines intensive induction chemotherapy, OR 2) patient is 60 years of age or older and the requested drug will be used as post-induction therapy following response to induction therapy with the requested drug, OR 3) patient has relapsed or refractory AML. For locally advanced, unresectable, or metastatic cholangiocarcinoma: the requested drug will be used as subsequent treatment for progression on or after systemic treatment. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | TOBRAMYCIN   |
| <b>Drug Names</b>                   | TOBRAMYCIN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Non-cystic fibrosis bronchiectasis   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis and non-cystic fibrosis bronchiectasis, the patient must meet one of the following: 1) Pseudomonas aeruginosa is present in the patient's airway cultures, OR 2) the patient has a history of Pseudomonas aeruginosa infection or colonization in the airways.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | TOPICAL LIDOCAINE   |
| <b>Drug Names</b>                   | GLYDO, LIDOCAINE, LIDOCAINE HCL, LIDOCAINE HCL JELLY, LIDOCAINE/PRILOCAINE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | 1) The requested drug is being used for topical anesthesia, AND 2) If the requested drug will be used as part of a compounded product, then all the active ingredients in the compounded product are Food and Drug Administration (FDA) approved for topical use. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 3 months  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.  |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | TOPICAL TESTOSTERONES   |
| <b>Drug Names</b>                   | TESTOSTERONE, TESTOSTERONE PUMP   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Gender Dysphoria  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Primary or hypogonadotropic hypogonadism: 1) Request is for continuation of testosterone therapy and the patient had a confirmed low morning testosterone level according to current practice guidelines or your standard lab reference values before starting testosterone therapy [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.] OR 2) Request is not for continuation of testosterone therapy and the patient has at least two confirmed low morning testosterone levels according to current practice guidelines or your standard lab reference values [Note: Safety and efficacy of testosterone products in patients with "age-related hypogonadism" (also referred to as "late-onset hypogonadism") have not been established.]. Gender dysphoria: The patient is able to make an informed decision to engage in hormone therapy. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

|                                     |  |
|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | TOPICAL TRETINOIN  |
| <b>Drug Names</b>                   | AVITA, TRETINOIN   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <br>                                |  |
| <b>Prior Authorization Group</b>    | TRAZIMERA  |
| <b>Drug Names</b>                   | TRAZIMERA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neoadjuvant treatment for human epidermal growth factor receptor 2 (HER2)-positive breast cancer, recurrent or advanced unresectable HER2-positive breast cancer, leptomeningeal metastases from HER2-positive breast cancer, brain metastases from HER2-positive breast cancer, HER2-positive esophageal and esophagogastric junction adenocarcinoma, HER2-positive advanced, recurrent, or metastatic uterine serous carcinoma, HER2-amplified and RAS and BRAF wild-type colorectal cancer in combination with pertuzumab or lapatinib, HER2-positive recurrent salivary gland tumor. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Neoadjuvant therapy for breast cancer: 6 months. Other: Plan Year.   |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

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| <b>Prior Authorization Group</b>    | TREPROSTINIL INJ   |
| <b>Drug Names</b>                   | TREPROSTINIL   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (World Health Organization [WHO] Group 1), the diagnosis was confirmed by right heart catheterization. For new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND 3) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

|                                     |                              |
|-------------------------------------|------------------------------|
| <b>Prior Authorization Group</b>    | TRIENTINE                    |
| <b>Drug Names</b>                   | TRIENTINE HYDROCHLORIDE      |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

|                                     |  |
|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | TRIKAFTA   |
| <b>Drug Names</b>                   | TRIKAFTA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For cystic fibrosis: The requested medication will not be used in combination with other medications containing ivacaftor. |
| <b>Age Restrictions</b>             | 6 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

|                                     |                              |
|-------------------------------------|------------------------------|
| <b>Prior Authorization Group</b>    | TRUSELTIQ                    |
| <b>Drug Names</b>                   | TRUSELTIQ                    |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications |
| <b>Off-label Uses</b>               | -                            |
| <b>Exclusion Criteria</b>           | -                            |
| <b>Required Medical Information</b> | -                            |
| <b>Age Restrictions</b>             | -                            |
| <b>Prescriber Restrictions</b>      | -                            |
| <b>Coverage Duration</b>            | Plan Year                    |
| <b>Other Criteria</b>               | -                            |

**Prior Authorization Group**

**Drug Names**

**PA Indication Indicator**

**Off-label Uses**

TRUXIMA

TRUXIMA

All FDA-approved Indications, Some Medically-accepted Indications

Non-Hodgkin's lymphoma subtypes [small lymphocytic lymphoma (SLL), mantle cell lymphoma, marginal zone lymphomas (nodal, splenic, gastric mucosa-associated lymphoid tissue [MALT], nongastric MALT), Burkitt lymphoma, primary cutaneous B-cell lymphoma, high-grade B-cell lymphoma with translocations of MYC and BCL2 and/or BCL6 (double/triple hit lymphoma), high-grade B-cell lymphoma not otherwise specified, histological transformation from follicular lymphoma to diffuse large B-cell lymphoma, histological transformation from nodal marginal zone lymphoma to diffuse large B-cell lymphoma, Castleman's disease, acquired immunodeficiency syndrome (AIDS)-related B-cell lymphoma, hairy cell leukemia, post-transplant lymphoproliferative disorder (PTLD), B-cell lymphoblastic lymphoma], refractory immune or idiopathic thrombocytopenic purpura (ITP), autoimmune hemolytic anemia, Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma, chronic graft-versus-host disease (GVHD), Sjogren syndrome, thrombotic thrombocytopenic purpura, refractory myasthenia gravis, Hodgkin's lymphoma (nodular lymphocyte-predominant), primary central nervous system (CNS) lymphoma, leptomeningeal metastases from lymphomas, acute lymphoblastic leukemia, prevention of Epstein-Barr virus (EBV)-related PTL, multiple sclerosis, immune checkpoint inhibitor-related toxicities, pemphigus vulgaris, pediatric aggressive mature B-cell lymphomas, and Rosai-Dorfman disease, and pediatric mature B-cell acute leukemia.

**Exclusion Criteria**

-

**Required Medical Information**

For moderately to severely active rheumatoid arthritis (new starts only): 1) patient meets ANY of the following: a) requested drug will be used in combination with methotrexate (MTX) OR b) patient has intolerance or contraindication to MTX, AND 2) patient meets ANY of the following: a) inadequate response, intolerance, or contraindication to MTX OR b) inadequate response or intolerance to a prior biologic disease-modifying antirheumatic drug (DMARD) or a targeted synthetic DMARD. Hematologic malignancies must be CD20-positive. For multiple sclerosis: 1) patient has a diagnosis of relapsing remitting multiple sclerosis and 2) patient has had an inadequate response to two or more disease-modifying drugs indicated for multiple sclerosis despite adequate duration of treatment.

**Age Restrictions**

-

**Prescriber Restrictions**

-

**Coverage Duration**

Immune checkpoint inhibitor-related toxicities: 3 months, All other: Plan Year

**Other Criteria**

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|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | TUKYSA  |
| <b>Drug Names</b>                   | TUKYSA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent human epidermal growth factor receptor 2 (HER2)-positive breast cancer  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | TURALIO   |
| <b>Drug Names</b>                   | TURALIO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Langerhans Cell Histiocytosis, Erdheim-Chester Disease, Rosai-Dorfman Disease   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For Langerhans Cell Histiocytosis: 1) disease has colony stimulating factor 1 receptor (CSF1R) mutation. For Erdheim-Chester Disease and Rosai-Dorfman Disease: 1) disease has CSF1R mutation AND patient has any of the following: a) symptomatic disease OR b) relapsed/refractory disease. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | UCERIS  |
| <b>Drug Names</b>                   | BUDESONIDE ER   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one 5-aminosalicylic acid (5-ASA) therapy.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 2 months  |
| <b>Other Criteria</b>               | -   |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | V-GO  |
| <b>Drug Names</b>                   | V-GO 20, V-GO 30, V-GO 40   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | Initial: 1) The patient has diabetes requiring insulin management with multiple daily injections AND 2) The patient is self-testing glucose levels 4 or more times per day OR the patient is using a continuous glucose monitor AND 3) The patient has experienced any of the following with the current diabetes regimen: inadequate glycemic control, recurrent hypoglycemia, wide fluctuations in blood glucose, dawn phenomenon with persistent severe early morning hyperglycemia, severe glycemic excursions.<br>Continuation: the patient has stable or improved glycemic control. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | VALCHLOR  |
| <b>Drug Names</b>                   | VALCHLOR  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Chronic or smoldering adult T-cell leukemia/lymphoma, Stage 2 or higher mycosis fungoides/Sezary syndrome, primary cutaneous marginal zone lymphoma, primary cutaneous follicle center lymphoma, lymphomatoid papulosis, unifocal Langerhans cell histiocytosis (LCH) with isolated skin disease.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | VARENICLINE TAB   |
| <b>Drug Names</b>                   | VARENICLINE STARTING MONT, VARENICLINE TARTRATE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 6 months  |
| <b>Other Criteria</b>               | -   |



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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | VELCADE  |
| <b>Drug Names</b>                   | BORTEZOMIB   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Systemic light chain amyloidosis, Waldenstrom's macroglobulinemia/lymphoplasmacytic lymphoma, multicentric Castleman's disease, adult T-cell leukemia/lymphoma, acute lymphoblastic leukemia, Kaposi's sarcoma, Hodgkin lymphoma, POEMS syndrome |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

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| <b>Prior Authorization Group</b>    | VENCLEXTA  |
| <b>Drug Names</b>                   | VENCLEXTA, VENCLEXTA STARTING PACK   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Mantle cell lymphoma, blastic plasmacytoid dendritic cell neoplasm (BPDCN), multiple myeloma, relapsed or refractory acute myeloid leukemia (AML), Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma, relapsed or refractory systemic light chain amyloidosis with translocation t(11:14)   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For acute myeloid leukemia (AML): 1) patient is 60 years of age or older OR 2) patient is less than 60 years of age with unfavorable risk genetics and TP53-mutation OR 3) patient has comorbidities that preclude use of intensive induction chemotherapy OR 4) patient has relapsed or refractory disease. For blastic plasmacytoid dendritic cell neoplasm (BPDCN): 1) patient has systemic disease being treated with palliative intent OR 2) patient has relapsed or refractory disease. For multiple myeloma: 1) the disease is relapsed or progressive AND 2) the requested drug will be used in combination with dexamethasone AND 3) patient has t(11:14) translocation. For Waldenstrom macroglobulinemia/lymphoplasmacytic lymphoma: 1) patient has previously treated disease that did not respond to primary therapy OR 2) patient has progressive or relapsed disease. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | VENTAVIS   |
| <b>Drug Names</b>                   | VENTAVIS   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For pulmonary arterial hypertension (World Health Organization [WHO] Group 1), the diagnosis was confirmed by right heart catheterization. For new starts only: 1) pretreatment mean pulmonary arterial pressure is greater than 20 mmHg, 2) pretreatment pulmonary capillary wedge pressure is less than or equal to 15 mmHg, AND 3) pretreatment pulmonary vascular resistance is greater than or equal to 3 Wood units. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | VERSACLOZ   |
| <b>Drug Names</b>                   | VERSACLOZ   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For the treatment of a severely ill patient with schizophrenia who failed to respond adequately to standard antipsychotic treatment (i.e., treatment-resistant schizophrenia): 1) the patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following generic products: A) aripiprazole, B) asenapine, C) olanzapine, D) quetiapine, E) risperidone, F) ziprasidone AND 2) The patient experienced an inadequate treatment response, intolerance, or contraindication to one of the following brand products: A) Latuda, B) Rexulti, C) Secuado, D) Vraylar. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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|-------------------------------------|--|
| <b>Prior Authorization Group</b>    | VERZENIO   |
| <b>Drug Names</b>                   | VERZENIO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative breast cancer in combination with fulvestrant or an aromatase inhibitor, or as a single agent if progression on prior endocrine therapy and prior chemotherapy in the metastatic setting. |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | VIGABATRIN   |
| <b>Drug Names</b>                   | VIGABATRIN, VIGADRONE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For complex partial seizures (CPS): patient has experienced an inadequate treatment response to at least two antiepileptic drugs for CPS.  |
| <b>Age Restrictions</b>             | Infantile Spasms: 1 month to 2 years of age. CPS: 2 years of age or older  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | VITRAKVI   |
| <b>Drug Names</b>                   | VITRAKVI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Non-metastatic neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, first-line treatment of NTRK gene fusion-positive solid tumors.   |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For all neurotrophic tyrosine receptor kinase (NTRK) gene fusion-positive solid tumors, the disease is without a known acquired resistance mutation.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | VIZIMPRO  |
| <b>Drug Names</b>                   | VIZIMPRO  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent non-small cell lung cancer (NSCLC).   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For non-small cell lung cancer (NSCLC): 1) the disease is recurrent, advanced or metastatic, and 2) the patient has sensitizing EGFR mutation-positive disease. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | VONJO   |
| <b>Drug Names</b>                   | VONJO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | VORICONAZOLE  |
| <b>Drug Names</b>                   | VORICONAZOLE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | The patient will use the requested drug orally or intravenously.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 6 months  |
| <b>Other Criteria</b>               | -   |

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|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | VOSEVI  |
| <b>Drug Names</b>                   | VOSEVI  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | Decompensated cirrhosis/moderate or severe hepatic impairment (Child Turcotte Pugh class B or C)  |
| <b>Required Medical Information</b> | For hepatitis C: Infection confirmed by presence of HCV RNA in the serum prior to starting treatment. Planned treatment regimen, genotype, prior treatment history, presence or absence of cirrhosis (compensated or decompensated [Child Turcotte Pugh class B or C]), presence or absence of HIV coinfection, presence or absence of resistance-associated substitutions where applicable, transplantation status if applicable. Coverage conditions and specific durations of approval will be based on current American Association for the Study of Liver Diseases (AASLD) treatment guidelines. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Criteria will be applied consistent with current AASLD-IDSA guidance.   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | VOTRIENT  |
| <b>Drug Names</b>                   | VOTRIENT  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Thyroid carcinoma (follicular, papillary, Hurthle cell, or medullary), uterine sarcoma, chondrosarcoma, gastrointestinal stromal tumor.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For renal cell carcinoma: The disease is advanced, relapsed, or stage IV. For gastrointestinal stromal tumor (GIST): the disease is unresectable, recurrent, or metastatic AND the patient has failed on an FDA-approved therapy (e.g., imatinib, sunitinib, regorafenib, ripretinib). For soft tissue sarcoma (STS): The patient does not have an adipocytic soft tissue sarcoma. For uterine sarcoma: The disease is recurrent or metastatic.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

|                                     |   |
|-------------------------------------|---|
| <b>Prior Authorization Group</b>    | WELIREG   |
| <b>Drug Names</b>                   | WELIREG   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | XALKORI   |
| <b>Drug Names</b>                   | XALKORI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Recurrent non-small cell lung cancer (NSCLC), NSCLC with high-level MET amplification or MET exon 14 skipping mutation, symptomatic or relapsed/refractory anaplastic lymphoma kinase (ALK)-fusion positive Erdheim-Chester Disease, symptomatic or relapsed/refractory (ALK)-fusion positive Rosai-Dorfman Disease, (ALK)-fusion positive Langerhans Cell Histiocytosis.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For NSCLC, the requested drug is used in any of the following settings: 1) the patient has recurrent, advanced or metastatic ALK-positive NSCLC, 2) the patient has recurrent, advanced or metastatic ROS-1 positive NSCLC, or 3) the patient has NSCLC with high-level MET amplification or MET exon 14 skipping mutation. For IMT, the disease is ALK-positive. For ALCL, the disease is relapsed or refractory and ALK-positive. |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | XELJANZ   |
| <b>Drug Names</b>                   | XELJANZ, XELJANZ XR   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For moderately to severely active rheumatoid arthritis (new starts only): 1) patient has experienced an inadequate treatment response, intolerance or has a contraindication to methotrexate (MTX) AND at least one tumor necrosis factor (TNF) inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). For active psoriatic arthritis (new starts only): 1) patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one TNF inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]) AND 2) the requested drug is used in combination with a nonbiologic DMARD. For active ankylosing spondylitis (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one tumor necrosis factor (TNF) inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). For moderately to severely active ulcerative colitis (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one tumor necrosis factor (TNF) inhibitor (e.g., Humira [adalimumab]). For active polyarticular course juvenile idiopathic arthritis (pcJIA) (new starts only): patient has experienced an inadequate treatment response, intolerance, or has a contraindication to at least one tumor necrosis factor (TNF) inhibitor (e.g., Enbrel [etanercept], Humira [adalimumab]). |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | XERMELO   |
| <b>Drug Names</b>                   | XERMELO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | XGEVA  |
| <b>Drug Names</b>                   | XGEVA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For hypercalcemia of malignancy: condition is refractory to intravenous (IV) bisphosphonate therapy or there is a clinical reason to avoid IV bisphosphonate therapy.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.   |
| <b>Prior Authorization Group</b>    | XHANCE   |
| <b>Drug Names</b>                   | XHANCE   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Patient has experienced an inadequate treatment response to generic fluticasone nasal spray.   |
| <b>Age Restrictions</b>             | 18 years of age or older   |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | XIFAXAN  |
| <b>Drug Names</b>                   | XIFAXAN  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For irritable bowel syndrome with diarrhea (IBS-D): 1) The patient has not previously received treatment with the requested drug OR 2) The patient has previously received treatment with the requested drug AND a) the patient is experiencing a recurrence of symptoms AND b) the patient has not already received an initial 14-day course of treatment and two additional 14-day courses of treatment with the requested drug. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Reduction in risk of overt HE recurrence: 6 Months, IBS-D: 14 Days   |
| <b>Other Criteria</b>               | -  |



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| <b>Prior Authorization Group</b>    | XOLAIR  |
| <b>Drug Names</b>                   | XOLAIR  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For allergic asthma initial therapy: 1) Patient has positive skin test (or blood test) to at least 1 perennial aeroallergen, 2) Patient has baseline IgE level greater than or equal to 30 IU/mL, and 3) Patient has inadequate asthma control despite current treatment with both of the following medications: a) Inhaled corticosteroid, and b) Additional controller (long acting beta2-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline) unless patient has an intolerance or contraindication to such therapies. For allergic asthma continuation therapy only: Patient's asthma control has improved on treatment with the requested drug since initiation of therapy. For chronic idiopathic urticaria (CIU) initial therapy: 1) Patient has been evaluated for other causes of urticaria, including bradykinin-related angioedema and IL-1-associated urticarial syndromes (auto-inflammatory disorders, urticarial vasculitis), and 2) Patient has experienced a spontaneous onset of wheals, angioedema, or both, for at least 6 weeks. For CIU continuation therapy: Patient has experienced a response (e.g., improved symptoms) since initiation of therapy. For nasal polyps: 1) the requested drug is used as add-on maintenance treatment, AND 2) the patient has experienced an inadequate treatment response to Xhance (fluticasone). |
| <b>Age Restrictions</b>             | For CIU: 12 years of age or older. For allergic asthma: 6 years of age or older. For nasal polyps: 18 years of age or older.  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | CIU initial: 6 months. All others: Plan Year.   |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | XOSPATA   |
| <b>Drug Names</b>                   | XOSPATA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FLT3 rearrangement  |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and FLT3 rearrangement: the disease is in chronic or blast phase.   |
| <b>Age Restrictions</b>             | 18 years of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | XPOVIO   |
| <b>Drug Names</b>                   | XPOVIO, XPOVIO 60 MG TWICE WEEKLY, XPOVIO 80 MG TWICE WEEKLY   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | XTANDI   |
| <b>Drug Names</b>                   | XTANDI   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The requested drug will be used in combination with a gonadotropin-releasing hormone (GnRH) analog or after bilateral orchiectomy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | XYREM  |
| <b>Drug Names</b>                   | XYREM  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | The diagnosis has been confirmed by sleep lab evaluation.<br>EDS: 1)The patient experienced an inadequate treatment response or intolerance to at least one central nervous system (CNS) stimulant drug (e.g., amphetamine, dextroamphetamine, or methylphenidate) OR has a contraindication that would prohibit a trial of central nervous system (CNS) stimulant drugs (e.g., amphetamine, dextroamphetamine, or methylphenidate) [Note: Coverage of amphetamines may require prior authorization.] AND 2) If the patient is 18 years of age or older, the patient experienced an inadequate treatment response or intolerance to at least one central nervous system (CNS) wakefulness promoting drug (e.g., armodafinil) OR has a contraindication that would prohibit a trial of central nervous system (CNS) wakefulness promoting drugs (e.g., armodafinil) [Note: coverage of armodafinil may require prior authorization.]. |
| <b>Age Restrictions</b>             | 7 years of age or older  |
| <b>Prescriber Restrictions</b>      | Prescribed by or in consultation with a sleep disorder specialist or neurologist.  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | If the request is for a continuation of therapy, then the patient experienced a decrease in daytime sleepiness with narcolepsy or a decrease in cataplexy episodes with narcolepsy.  |
| <b>Prior Authorization Group</b>    | ZARXIO   |
| <b>Drug Names</b>                   | ZARXIO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Neutropenia in myelodysplastic syndromes (MDS), agranulocytosis, neutropenia in aplastic anemia, human immunodeficiency virus (HIV)-related neutropenia, neutropenia related to renal transplant.  |
| <b>Exclusion Criteria</b>           | Use of the requested product within 24 hours prior to or following chemotherapy.   |
| <b>Required Medical Information</b> | For prophylaxis or treatment of myelosuppressive chemotherapy-induced febrile neutropenia (FN) patient must meet both of the following: 1) Patient has a solid tumor or non-myeloid cancer, and 2) Patient has received, is currently receiving, or will be receiving treatment with myelosuppressive anti-cancer therapy.   |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | 6 months   |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ZEJULA  |
| <b>Drug Names</b>                   | ZEJULA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | In combination with bevacizumab for persistent or recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer for platinum-sensitive disease, uterine leiomyosarcoma |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For uterine leiomyosarcoma: 1) the requested drug is used as second-line therapy AND 2) the patient has BRCA-altered disease.   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ZELBORAF   |
| <b>Drug Names</b>                   | ZELBORAF   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Non-small cell lung cancer, hairy cell leukemia, thyroid carcinoma (i.e., papillary carcinoma, follicular carcinoma, and Hurthle cell carcinoma), central nervous system cancer (i.e., glioma, meningioma, astrocytoma), adjuvant systemic therapy for cutaneous melanoma, Langerhans cell histiocytosis.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For adjuvant treatment of melanoma, and central nervous system (CNS) cancer (i.e., glioma, meningioma, astrocytoma): 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K) and 2) The requested drug will be used in combination with cobimetinib. For unresectable or metastatic melanoma: 1) The tumor is positive for BRAF V600 activating mutation (e.g., V600E or V600K) and 2) the requested drug will be used as a single agent, or in combination with cobimetinib (with or without atezolizumab). For Erdheim-Chester Disease and Langerhans Cell Histiocytosis: Tumor is positive for BRAF V600 mutation. For non-small cell lung cancer: 1) Tumor is positive for the BRAF V600E mutation, and 2) The patient has recurrent, advanced, or metastatic disease. For thyroid carcinoma: 1) Tumor is positive for BRAF mutation, and 2) Patient has radioiodine refractory follicular, Hurthle cell, or papillary thyroid carcinoma. For hairy cell leukemia: The requested drug will be used for subsequent therapy. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |

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| <b>Prior Authorization Group</b>    | ZIEXTENZO   |
| <b>Drug Names</b>                   | ZIEXTENZO   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Stem cell transplantation-related indications   |
| <b>Exclusion Criteria</b>           | Use of the requested product less than 24 hours before or after chemotherapy.   |
| <b>Required Medical Information</b> | For prophylaxis of myelosuppressive chemotherapy-induced febrile neutropenia: the patient must meet both of the following: 1) Patient has a solid tumor or non-myeloid cancer, and 2) Patient is currently receiving or will be receiving treatment with myelosuppressive anti-cancer therapy.  |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | 6 months  |
| <b>Other Criteria</b>               | -   |
| <b>Prior Authorization Group</b>    | ZIRABEV   |
| <b>Drug Names</b>                   | ZIRABEV   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Breast cancer, central nervous system (CNS) tumor types: adult low-grade (WHO Grade I or II) glioma, adult intracranial and spinal ependymoma, anaplastic gliomas, adult medulloblastoma, primary central nervous system lymphoma, meningiomas, limited and extensive brain metastases, and metastatic spine tumors, malignant pleural mesothelioma, ovarian cancer/fallopian tube cancer/primary peritoneal cancer types: carcinosarcoma (malignant mixed Mullerian tumors), clear cell carcinoma, mucinous carcinoma, grade 1 endometrioid carcinoma, low-grade serous carcinoma, ovarian borderline epithelial tumors (low malignant potential) with invasive implants, and malignant sex cord-stromal tumors, soft tissue sarcoma types: angiosarcoma and solitary fibrous tumor/hemangiopericytoma, uterine neoplasms, endometrial carcinoma, vulvar squamous cell carcinoma, and ophthalmic-related disorders: diabetic macular edema, neovascular (wet) age-related macular degeneration including polypoidal choroidopathy and retinal angiomatous proliferation subtypes, macular edema following retinal vein occlusion, proliferative diabetic retinopathy, choroidal neovascularization, neovascular glaucoma and retinopathy of prematurity, hepatocellular carcinoma, small bowel adenocarcinoma. |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | Coverage under Part D will be denied if coverage is available under Part A or Part B as the medication is prescribed and dispensed or administered for the individual.  |

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| <b>Prior Authorization Group</b>    | ZOLINZA   |
| <b>Drug Names</b>                   | ZOLINZA   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications   |
| <b>Off-label Uses</b>               | Mycosis fungoides, Sezary syndrome.   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | -   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | ZONISADE  |
| <b>Drug Names</b>                   | ZONISADE  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | For adjunctive treatment of partial-onset seizures (i.e., focal-onset seizures): 1) The patient has experienced an inadequate treatment response, intolerance, or has a contraindication to a generic anticonvulsant AND the patient has experienced an inadequate treatment response, intolerance, or has a contraindication to any of the following: Aptiom, Xcopri, Spritam OR 2) The patient has difficulty swallowing solid oral dosage forms (e.g., tablets, capsules). |
| <b>Age Restrictions</b>             | 16 years of age or older  |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |
| <br>                                |   |
| <b>Prior Authorization Group</b>    | ZTALMY  |
| <b>Drug Names</b>                   | ZTALMY  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications  |
| <b>Off-label Uses</b>               | -   |
| <b>Exclusion Criteria</b>           | -   |
| <b>Required Medical Information</b> | -   |
| <b>Age Restrictions</b>             | 2 years of age or older   |
| <b>Prescriber Restrictions</b>      | -   |
| <b>Coverage Duration</b>            | Plan Year   |
| <b>Other Criteria</b>               | -   |

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| <b>Prior Authorization Group</b>    | ZYDELIG  |
| <b>Drug Names</b>                   | ZYDELIG  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Refractory chronic lymphocytic leukemia (CLL), relapsed or refractory small lymphocytic lymphoma (SLL).  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | -  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ZYKADIA  |
| <b>Drug Names</b>                   | ZYKADIA  |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications, Some Medically-accepted Indications  |
| <b>Off-label Uses</b>               | Recurrent ALK-positive non-small cell lung cancer (NSCLC), recurrent, advanced, or metastatic ROS1-positive NSCLC, inflammatory myofibroblastic tumor (IMT), brain metastases from NSCLC.  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | For NSCLC: the patient has recurrent, advanced, or metastatic ALK-positive or ROS1-positive disease. For inflammatory myofibroblastic tumor: the disease is ALK-positive. For brain metastases from NSCLC: the patient has ALK-positive NSCLC. |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |
| <b>Prior Authorization Group</b>    | ZYPREXA RELPREVV   |
| <b>Drug Names</b>                   | ZYPREXA RELPREVV   |
| <b>PA Indication Indicator</b>      | All FDA-approved Indications   |
| <b>Off-label Uses</b>               | -  |
| <b>Exclusion Criteria</b>           | -  |
| <b>Required Medical Information</b> | Tolerability with oral olanzapine has been established.  |
| <b>Age Restrictions</b>             | -  |
| <b>Prescriber Restrictions</b>      | -  |
| <b>Coverage Duration</b>            | Plan Year  |
| <b>Other Criteria</b>               | -  |