

## PRIOR AUTHORIZATION CRITERIA

This list is current as of November 1, 2022, and pertains to the following formularies:

2022 Pharmacy Benefit Dimensions Prescription Drug Plan (PDP) Part D 3 Tier Formulary	Version 22
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Pharmacy Benefit Dimensions requires you (or your physician) to get prior authorization for certain drugs listed on the formularies above. This means that you will need to get approval from us before you fill your prescriptions. If you do not get approval, we may not cover the drug. These drugs are listed with a “PA” in the Requirements/Notes column on the formularies. This document contains the Prior Authorization requirements that are associated with the formularies listed above.

If you have any questions, please contact our Medicare Member Services Department at 1-800-667-5936 or, for TTY 711, October 1<sup>st</sup> – March 31<sup>st</sup>: Monday through Sunday from 8 a.m. to 8 p.m. ET, April 1<sup>st</sup> – September 30<sup>th</sup>: Monday through Friday from 8 a.m. to 8 p.m. ET.

Pharmacy Benefit Dimensions is a subsidiary of Independent Health. Independent Health is a PDP with a Medicare contract. Enrollment in Pharmacy Benefit Dimensions PDP depends on contract renewal between Independent Health and CMS.

The formulary may change at any time. You will receive notice when necessary.

## ABILIFY MYCITE (aripiprazole with sensor)

### Products Affected

- ABILIFY MYCITE MAINTENANCE KIT ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG
- ABILIFY MYCITE STARTER KIT ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG
- ABILIFY MYCITE ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Dementia-related psychosis
<b>Required Medical Information</b>	Diagnosis of covered use, documentation of previous aripiprazole use (see Other Criteria).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. For approval, the patient must have documentation of at least a one-month trial of generic aripiprazole solution, tablets, or orally-disintegrating tablets.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# ACTHAR (corticotropin)

## Products Affected

- ACTHAR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Request for IV administration, treatment of patients under 2 years of age in whom congenital infections are suspected, patients with scleroderma, osteoporosis, systemic fungal infections, ocular herpes simplex, recent surgery, a history of or presence of a peptic ulcer, congestive heart failure, uncontrolled hypertension, primary adrenocortical insufficiency, adrenocortical hyperfunction, or sensitivity to proteins of porcine origin
<b>Required Medical Information</b>	Diagnosis of covered use, submission of blood pressure reading and baseline serum sodium and potassium levels.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Updated blood pressure, sodium, and potassium levels since the previous authorization will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## ACTIMMUNE (interferon gamma-1b)

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### Products Affected

- ACTIMMUNE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ADEMPAS (riociguat)

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## Products Affected

- ADEMPAS

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy, severe (Child-Pugh class C) hepatic impairment, creatinine clearance below 15 mL/min or on dialysis, concurrent use with nitrates or nitric oxide donors in any form, concurrent use with phosphodiesterase inhibitors
Required Medical Information	Diagnosis of covered use, submission of negative pregnancy test result for female patients of childbearing age, creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance).
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology and pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# AKYNZEO (netupitant/palonosetron)

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## Products Affected

- AKYNZEO ORAL

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, severe renal impairment, end-stage renal disease
Required Medical Information	Diagnosis of covered use, confirmation patient will receive concurrent dexamethasone therapy as indicated based on level of chemotherapy regimen emetogenicity.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. If the medication is being administered related to cancer treatment and is a full replacement for intravenous administration of antiemetic therapy within 48 hours of cancer treatment, it is covered as a Part B benefit. To be eligible for Part B coverage, the prescribing physician must indicate this on the prescription. Otherwise it may be covered as a Part D benefit.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ALECENSA (alectinib)

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## Products Affected

- ALECENSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of ALK-positive tumor.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ALPHA-1-PROTEINASE INHIBITORS

## Products Affected

- ARALAST NP INTRAVENOUS SOLUTION RECONSTITUTED 1000 MG, 500 MG
- GLASSIA
- PROLASTIN-C INTRAVENOUS SOLUTION
- RECONSTITUTED ZEMAIRA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Individuals with immunoglobulin A (IgA) deficiency who have known antibodies against IgA
<b>Required Medical Information</b>	Diagnosis of covered use, confirmation that patient has clinically evident emphysema secondary to congenital alpha-1-PI deficiency by submission of pulmonary function testing (e.g., spirometry or body plethysmography), X-ray radiography, or diffusing capacity of the lung for carbon monoxide (DLCO).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to pulmonology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. Continuation of therapy requests require objective documentation from the prescriber that the patient's symptoms have improved. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# ALUNBRIG (brigatinib)

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## Products Affected

- ALUNBRIG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of ALK-positive tumor.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# AMANTADINE EXTENDED-RELEASE PRODUCTS

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## Products Affected

- GOCOVRI

PA Criteria	Criteria Details
Exclusion Criteria	End stage renal disease (creatinine clearance below 15 mL/min)
Required Medical Information	Diagnosis of covered use, documentation patient tried and failed immediate-release amantadine.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# AMVUTTRA (vutrisiran)

## Products Affected

- AMVUTTRA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Prior or scheduled liver transplant, New York Heart Association (NYHA) heart failure classification greater than 2
<b>Required Medical Information</b>	Diagnosis of covered use confirmed by (1) genetic testing including a mutation in the TTR gene and (2) signs and/or symptoms of polyneuropathy, including submission of baseline polyneuropathy disability (PND) score (required to be less than or equal to IIIb), submission of NYHA heart failure classification (required to be less than or equal to 2), previous medication(s) patient has tried and failed (at least one of either inotersen or patisiran).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology and specialists in genetic diseases
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. For approval, the patient must have tried and failed to have an adequate response to or had an intolerance to either inotersen or patisiran. Documentation of a positive response to therapy will be required for initial reauthorization after the first year. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## ANADROL-50 (oxymetholone)

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### Products Affected

- ANADROL-50

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Carcinoma of the prostate or breast in male patients, carcinoma of the breast in females with hypercalcemia, women who are or may become pregnant, nephrosis or the nephrotic phase of nephritis, severe hepatic dysfunction
<b>Required Medical Information</b>	Diagnosis of covered use, submission of CBC, liver function tests, and pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	6 months
<b>Other Criteria</b>	PA applies to all. Updated CBC since the previous authorization will be required for reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# ARANESP (darbepoetin alfa)

## Products Affected

- ARANESP (ALBUMIN FREE) INJECTION SOLUTION 100 MCG/ML, 200 MCG/ML, 25 MCG/ML, 300 MCG/ML, 40 MCG/ML, 60 MCG/ML
- ARANESP (ALBUMIN FREE) INJECTION SOLUTION

PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of hemoglobin or hematocrit level, serum iron, total iron-binding capacity (TIBC), and transferrin within 30 days of request date, documentation that the patient does not have uncontrolled hypertension.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	For non-ESRD-related conditions: 90 days. For ESRD-related conditions: 1 year.
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ARCALYST (rilonacept)

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## Products Affected

- ARCALYST

PA Criteria	Criteria Details
Exclusion Criteria	Active or chronic infection, coadministration with TNF-blocking agents
Required Medical Information	Diagnosis of covered use, TB skin test result obtained within the past 12 months.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ARIKAYCE (amikacin inhalation)

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### Products Affected

- ARIKAYCE

PA Criteria	Criteria Details
Exclusion Criteria	Non-refractory Mycobacterium avium complex (MAC) lung disease
Required Medical Information	Diagnosis of covered use, submission of other therapies that have been tried and failed or cannot be used because of a contraindication.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	This medication is covered as a Part B benefit except for enrollees residing in a long-term care facility. PA applies to all when covered as a Part D benefit.
Indications	All Medically-accepted Indications.
Off Label Uses	

## AURYXIA (ferric citrate)

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### Products Affected

- AURYXIA

PA Criteria	Criteria Details
Exclusion Criteria	Iron overload syndrome
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# AYVAKIT (avapritinib)

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## Products Affected

- AYVAKIT ORAL TABLET 100 MG, 200 MG, 25 MG, 300 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers or strong CYP3A inhibitors
Required Medical Information	Diagnosis of covered use. For gastrointestinal stromal tumor (GIST), submission of test result confirming presence of PDGFRA exon 18 mutation.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to allergy, immunology, and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BALVERSA (erdafitinib)

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## Products Affected

- BALVERSA

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP2C9 or CYP3A4 inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved companion test showing susceptible FGFR2 or FGFR3 genetic alterations, prior chemotherapy regimen(s) used.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BEMPEDOIC ACID

## Products Affected

- NEXLETOL
- NEXLIZET

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Concomitant pravastatin utilization with doses above 40 mg/day, concomitant simvastatin utilization with doses above 20 mg/day
<b>Required Medical Information</b>	Diagnosis of covered use, submission of current or previous lipid-lowering therapies.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. For approval, the patient must currently be using a statin plus ezetimibe or the patient must have tried and failed to have an adequate response to or had an intolerance to at least two statins or one statin and ezetimibe. At least one statin previously tried and failed must be a hydrophilic statin.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# BENLYSTA (belimumab)

## Products Affected

- BENLYSTA SUBCUTANEOUS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe active central nervous system lupus, patients using other biologic medications or intravenous cyclophosphamide
<b>Required Medical Information</b>	Diagnosis of covered use, confirmation that the patient is taking standard therapy defined as at least one of the following: corticosteroids, NSAIDs, antimalarials, or immunosuppressants.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# BESREMI (ropeginterferon alfa-2b-njft)

## Products Affected

- BESREMI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	History or presence of severe psychiatric disorders (including severe depression or suicidal ideation), history of presence of active serious or untreated autoimmune disease, moderate or severe hepatic impairment (Child-Pugh class B or C), immunosuppressed transplant recipients, severe or unstable cardiovascular disease (e.g., uncontrolled hypertension, NYHA class 2-4 congestive heart failure, serious cardiac arrhythmia, significant coronary artery stenosis, unstable angina), stroke or myocardial infarction within previous 6 months, severe renal impairment (eGFR less than 30 mL/min/1.73 m <sup>2</sup> )
<b>Required Medical Information</b>	Diagnosis of covered use, submission of eGFR, documentation patient has tried and failed, has a contraindication to, or could not tolerate hydroxyurea, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology and oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# BIOLOGIC RESPONSE MODIFIERS

## Products Affected

- ACTEMRA ACTPEN
- ACTEMRA SUBCUTANEOUS
- CIMZIA PREFILLED
- CIMZIA STARTER KIT
- CIMZIA SUBCUTANEOUS KIT 2 X 200 MG
- KEVZARA
- OTEZLA
- SIMPONI SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- SIMPONI SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- TREMFYA
- ZEPOSIA
- ZEPOSIA 7-DAY STARTER PACK
- ZEPOSIA STARTER KIT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	For Zeposia for the treatment of multiple sclerosis, only diagnosis of covered use is required. For all other drugs managed by this policy and for Zeposia for indications other than multiple sclerosis, diagnosis of covered use, submission of previous therapies. For all drugs managed by this policy except Otezla and Zeposia, submission of baseline latent tuberculosis screening test (Mantoux tuberculin skin test [a.k.a. PPD test] or interferon-gamma release assay [IGRA]).
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. With the exception of Zeposia for the treatment of multiple sclerosis only, for approval of a drug managed by this policy, the patient must have tried and failed to have an adequate response to or had an intolerance to at least two preferred agents (Cosentyx, Enbrel, Humira, Rinvoq, Skyrizi, Stelara, and Xeljanz/Xeljanz XR) for the indication submitted, where possible. For all drugs managed by this policy except Otezla and Zeposia, if TB screening test returns a positive result, coverage will be delayed until latent TB is treated. For re-authorization, yearly TB screening test or chest X-ray required for patients who live in, work in, or travel to areas where TB exposure is likely while on treatment or for those who have previously had a positive TB screening test.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BOSULIF (bosutinib)

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## Products Affected

- BOSULIF

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inhibitors or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of renal function testing. For accelerated or blast phase Ph+ CML, documentation of resistance or intolerance to at least one of the following prior therapies: imatinib, dasatinib, or nilotinib.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## BRAFTOVI/MEKTOVI (encorafenib/binimetinib)

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### Products Affected

- BRAFTOVI ORAL CAPSULE 75 MG
- MEKTOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of BRAF V600E or V600K mutation, serum potassium, and serum magnesium. For metastatic melanoma, confirmation that encorafenib and binimetinib will be co-administered. For metastatic colorectal cancer, confirmation that encorafenib and cetuximab will be co-administered.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# BRIVIACT (brivaracetam)

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## Products Affected

- BRIVIACT ORAL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	PA not required for neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BRONCHITOL (mannitol powder for inhalation)

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## Products Affected

- BRONCHITOL

PA Criteria	Criteria Details
Exclusion Criteria	Documented Bronchitol Tolerance Test failure
Required Medical Information	Diagnosis of covered use, documentation patient has passed the Bronchitol Tolerance Test, attestation patient will not be using in combination with hypertonic (7%) sodium chloride nebulized solution.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For approval, the patient must must have tried and failed to have an adequate response to or had an intolerance to hypertonic (7%) sodium chloride nebulized solution.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BRUKINSA (zanubrutinib)

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## Products Affected

- BRUKINSA

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of prior chemotherapy regimen(s) used.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# BUTALBITAL-CONTAINING PRODUCTS IN OLDER PATIENTS

## Products Affected

- *ascomp-codeine*
- BUPAP ORAL TABLET 50-300 MG
- BUTALBITAL-ACETAMINOPHEN ORAL TABLET 50-300 MG, 50-325 MG
- BUTALBITAL-APAP-CAFF-COD
- *butalbital-apap-caffeine oral capsule*
- *butalbital-apap-caffeine oral tablet 50-325-40 mg*
- *butalbital-asa-caff-codeine*
- *butalbital-aspirin-caffeine oral capsule*
- TENCON ORAL TABLET 50-325 MG
- VTOL LQ
- ZEBUTAL ORAL CAPSULE 50-325-40 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation patient has tried and failed a preferred alternative such as ibuprofen or rizatriptan, or has contraindications to all alternatives.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## BYLVAY (odevixibat)

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### Products Affected

- BYLVAY
- BYLVAY (PELLETS)

PA Criteria	Criteria Details
Exclusion Criteria	History of liver transplant, clinical evidence of decompensated cirrhosis
Required Medical Information	Diagnosis of covered use confirmed by molecular genetic testing, attestation drug-induced pruritus has been ruled out.
Age Restrictions	
Prescriber Restrictions	Restricted to gastroenterology and hepatology
Coverage Duration	1 year
Other Criteria	PA applies to all. Attestation of improvement in pruritus symptoms and submission of liver function testing, including serum bilirubin, since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

## CABLIVI (caplacizumab-yhdp)

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### Products Affected

- CABLIVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, confirmation drug will be given with plasma exchange and immunosuppressive therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology, hematology, and immunology
Coverage Duration	3 months
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CABOMETYX (cabozantinib)

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## Products Affected

- CABOMETYX

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment
Required Medical Information	Diagnosis of covered use.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CALQUENCE (acalabrutinib)

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## Products Affected

- CALQUENCE

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, coadministration with strong CYP3A inhibitors or proton pump inhibitors
Required Medical Information	Diagnosis of covered use, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# CAMZYOS (mavacamten)

## Products Affected

- *camzyos*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Left ventricular ejection fraction (LVEF) less than 55%, coadministration with a non-dihydropyridine (DHP) calcium channel blocker (CCB) plus disopyramide
<b>Required Medical Information</b>	Diagnosis of covered use including all three of the following: (1) attestation patient has exertional symptoms consistent with the definition of NYHA class II or III disease, (2) confirmation of left ventricular (LV) outflow tract obstruction gradient of at least 50 mmHg either at rest, during Valsalva maneuver testing, or after exercise, and (3) confirmation of LV wall thickness of at least 15 mm or at least 13 mm if condition is familial, submission of current LVEF, any previous or current therapies tried for the condition, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to cardiology
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. For approval, the patient must have tried and failed to have an adequate response to or had an intolerance/contraindication to both a beta-blocker and a non-DHP CCB. Documentation of a positive response to therapy will be required for initial reauthorization after the first 6 months. Maintenance of a clinical benefit and attestation that prescriber believes benefits of continuing therapy outweigh the potential risks to the patient will be required for subsequent annual reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# CAPLYTA (lumateperone)

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## Products Affected

- CAPLYTA ORAL CAPSULE 10.5 MG, 21 MG, 42 MG

PA Criteria	Criteria Details
Exclusion Criteria	Dementia-related psychosis, coadministration with moderate or strong CYP3A4 inhibitors or CYP3A4 inducers
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to psychiatry
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For approval for schizophrenia, the patient must have tried and failed to have an adequate response to or had an intolerance to aripiprazole and at least one other generic second-generation atypical antipsychotic (e.g., paliperidone, quetiapine, risperidone, etc.) or Latuda.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CAPRELSA (vandetanib)

## Products Affected

- CAPRELSA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	History of congenital long QT syndrome, Torsades de pointes, uncompensated heart failure, or bradyarrhythmias, moderate or severe hepatic impairment, QTcF interval greater than 450 msec
<b>Required Medical Information</b>	Diagnosis of covered use, submission of baseline serum potassium, calcium, magnesium, ALT, AST, bilirubin, TSH, creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance), ECG, and pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## CARBAGLU (carglumic acid)

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### Products Affected

- *carglumic acid*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of plasma ammonia level.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated plasma ammonia level since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CERDELGA (eliglustat)

## Products Affected

- CERDELGA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pre-existing cardiac disease, long QT syndrome, coadministration with Class Ia or Class III antiarrhythmics, patients who are extensive or intermediate CYP2D6 metabolizers taking a strong CYP2D6 inhibitor with a strong or moderate CYP3A inhibitor, intermediate and poor CYP2D6 metabolizers taking a strong CYP3A inhibitor
<b>Required Medical Information</b>	Diagnosis of covered use, submission of CYP2D6 metabolizer status as detected by an FDA-cleared test for determining CYP2D6 genotype, liver function testing or Child-Pugh score.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Updated liver function testing or Child-Pugh score since the previous authorization will be required for subsequent reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# CGRP INHIBITORS

## Products Affected

- AIMOVIG SUBCUTANEOUS SOLUTION AUTO-INJECTOR 140 MG/ML, 70 MG/ML
- AJOVY
- EMGALITY
- EMGALITY (300 MG DOSE)
- NURTEC
- QULIPTA
- UBRELVY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For treatment of migraine headache prevention, submission of baseline migraine days per month from medical chart, documentation patient has tried and failed at least two preferred FDA-approved (propranolol, timolol, topiramate, valproic acid) or compendial alternatives (e.g., amitriptyline, atenolol) for migraine prophylaxis, at least one alternative if they have contraindications to all other alternatives, or has contraindications to all alternatives. For Nurtec or Ubrelyvy for the treatment of acute migraine, documentation of prior use of at least one triptan.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	Initially 3 months, then 1 year
Other Criteria	PA applies to all. For all drugs in this policy except Ubrelyvy for migraine preventive treatment, submission of on-treatment headache days per month demonstrating improvement from baseline will be required for initial reauthorization after the first 3 months. Documentation of a clinically relevant response to therapy or maintenance of a clinical benefit will be required for subsequent reauthorizations. For approval of Nurtec for migraine headache prevention, the patient must have a diagnosis of episodic migraine, defined as fewer than 15 headache days per month. For approval of Emgality for migraine headache prevention, the patient must have tried and failed to have an adequate response to or had an intolerance to Aimovig and Ajovy. For Ajovy, a description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CHENODAL (chenodiol)

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## Products Affected

- CHENODAL

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy, known hepatocyte dysfunction, bile duct abnormalities such as intrahepatic cholestasis, primary biliary cirrhosis, or sclerosing cholangitis, radiopaque stones, nonvisualizing gallbladder confirmed as nonvisualizing after 2 consecutive single doses of dye, compelling reasons for gallbladder surgery
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	24 months
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## CHOLBAM (cholic acid)

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### Products Affected

- CHOLBAM

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of liver function testing.
Age Restrictions	
Prescriber Restrictions	Restricted to hepatology, gastroenterology, and pediatric gastroenterology
Coverage Duration	Initially 3 months, then 1 year
Other Criteria	PA applies to all. Documentation of liver function improvement without complete biliary obstruction or persistent clinical or laboratory indications of worsening liver function or cholestasis will be required for initial reauthorization after the first 3 months. Updated liver function testing since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	



# COMETRIQ (cabozantinib)

## Products Affected

- COMETRIQ (100 MG DAILY DOSE) ORAL KIT 80 & 20 MG
- COMETRIQ (140 MG DAILY DOSE) ORAL KIT 3 X 20 MG & 80 MG
- COMETRIQ (60 MG DAILY DOSE)

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment (Child-Pugh class C)
Required Medical Information	Diagnosis of covered use, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# COPIKTRA (duvelisib)

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## Products Affected

- COPIKTRA ORAL CAPSULE 15 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of at least two prior therapies tried and failed, submission of pregnancy status for female patients of childbearing potential, attestation patient will receive prophylaxis for <i>Pneumocystis jirovecii</i> pneumonia (PJP) and, if necessary, cytomegalovirus.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology or oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CORLANOR (ivabradine)

## Products Affected

- CORLANOR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Acute decompensated heart failure, clinically significant hypotension, clinically significant bradycardia, severe hepatic impairment, pacemaker dependence (heart rate maintained exclusively by the pacemaker), or sick sinus syndrome, sinoatrial block, or 3rd degree AV block unless a functioning demand pacemaker is present, coadministration with strong CYP3A4 inhibitors
<b>Required Medical Information</b>	Diagnosis of covered use described as is indicated (1) to reduce the risk of hospitalization for worsening heart failure in patients with stable, symptomatic chronic heart failure with left ventricular ejection fraction less than or equal to 35%, who are in sinus rhythm with resting heart rate greater than or equal to 70 beats per minute and either are on maximally tolerated doses of beta-blockers or have a contraindication to beta-blocker use or (2) for stable symptomatic heart failure due to dilated cardiomyopathy in patients who are in sinus rhythm with an elevated heart rate. Submission of current baseline blood pressure reading, confirmation that patient does not have any of the following: (1) acute decompensated heart failure, (2) sick sinus syndrome, sinoatrial block, or 3rd degree AV block, unless a functioning demand pacemaker is present, (3) resting heart rate less than 60 bpm prior to treatment, (4) severe hepatic impairment, (5) pacemaker dependence (heart rate maintained exclusively by the pacemaker). For patients under 18 years old, (1) left ventricular ejection fraction less than or equal to 45% and (2) resting heart rate greater than or equal to the following age-stratified requirements: (a) 105 beats per minute in ages 6 to 12 months old, (b) 95 beats per minute in ages 1 to 3 years old, (c) 75 beats per minute in ages 3 to 5 years old, and (d) 70 beats per minute in ages 5 to 18 years old.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	Restricted to cardiology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Continuation requires documentation of clinically relevant response to therapy.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# CORTROPHIN (corticotropin)

## Products Affected

- CORTROPHIN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Request for IV administration, treatment of patients under 2 years of age in whom congenital infections are suspected, patients with scleroderma, osteoporosis, systemic fungal infections, ocular herpes simplex, recent surgery, a history of or presence of a peptic ulcer, congestive heart failure, uncontrolled hypertension, primary adrenocortical insufficiency, adrenocortical hyperfunction, or sensitivity to proteins of porcine origin
<b>Required Medical Information</b>	Diagnosis of covered use, submission of blood pressure reading and baseline serum sodium and potassium levels.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Updated blood pressure, sodium, and potassium levels since the previous authorization will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# COTELLIC/ZELBORAF (cobimetinib/vemurafenib)

## Products Affected

- COTELLIC
- ZELBORAF

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inhibitors or inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of BRAF V600E or V600K mutation, submission of left ventricular ejection fraction, ECG, and serum potassium, magnesium, and calcium levels. For patients using cobimetinib, confirmation that it will be co-administered with vemurafenib.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Vemurafenib is not indicated in wild-type BRAF melanoma and will not be approved for this use.
Indications	All Medically-accepted Indications.
Off Label Uses	

# CYSTEAMINE EYE DROPS

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## Products Affected

- CYSTADROPS
- CYSTARAN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DALFAMPRIDINE

## Products Affected

- *dalfampridine er*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	History of seizure, moderate or severe renal impairment (CrCl less than or equal to 50 mL/min)
<b>Required Medical Information</b>	Diagnosis of covered use, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance) and objective measurement of walking speed, confirmation that patient is able to walk.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology
<b>Coverage Duration</b>	Initially 3 months, then 1 year
<b>Other Criteria</b>	PA applies to all. Documentation the patient has demonstrated an improvement in walking speed from baseline measure (or maintenance of improvement if patient has been on long-term therapy) since starting medication will be required for initial reauthorization after the first 3 months. Updated creatinine clearance since the previous authorization and confirmation patient is able to walk will be required for subsequent annual reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# DAURISMO (glasdegib)

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## Products Affected

- DAURISMO ORAL TABLET 100 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, confirmation patient will also be receiving cytarabine as part of chemotherapeutic regimen. If patient is under 75 years of age, documentation of comorbidities that preclude use of intensive induction chemotherapy, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# DAYVIGO (lemborexant)

## Products Affected

- DAYVIGO ORAL TABLET 10 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	Narcolepsy
Required Medical Information	Diagnosis of covered use. Patient must have tried and failed to tolerate or had an inadequate response to two covered alternative therapies recommended by the American Academy of Sleep Medicine (doxepin, eszopiclone, ramelteon, suvorexant, temazepam, zaleplon, zolpidem) including one non-suvorexant therapy for sleep maintenance (doxepin, eszopiclone, temazepam) if that is the diagnosis of covered use.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DEFERASIROX

## Products Affected

- *deferasirox granules*
- *deferasirox oral tablet*
- *deferasirox oral tablet soluble*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe hepatic impairment, estimated glomerular filtration rate less than 40 mL per min, platelet count below $50 \times 10^9/L$ , high-risk myelodysplastic syndromes, advanced malignancies
<b>Required Medical Information</b>	Diagnosis of covered use, submission of CBC, LFTs, serum creatinine, ferritin, and urine protein values, estimated glomerular filtration rate, documentation that member has had yearly ophthalmic and auditory testing.
<b>Age Restrictions</b>	2 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	3 months
<b>Other Criteria</b>	PA applies to all. Updated ferritin level within last 3 months and updated CBC, LFT, urine protein value, estimated glomerular filtration rate, and ophthalmic and auditory testing since the previous authorization will be required for subsequent reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# DEFERIPRONE

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## Products Affected

- *deferiprone*
- FERRIPROX ORAL SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of serum ferritin levels, CBC, ANC, platelet count, and serum ALT.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DIACOMIT (stiripentol)

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## Products Affected

- DIACOMIT

PA Criteria	Criteria Details
Exclusion Criteria	Moderate or severe renal impairment, moderate or severe hepatic impairment
Required Medical Information	Diagnosis of covered use, confirmation patient is also receiving clobazam.
Age Restrictions	2 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Monotherapy requests for Dravet syndrome will not be approved as there are no clinical data to support using stiripentol in this manner.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DICLOFENAC 1% GEL

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## Products Affected

- *diclofenac sodium external gel*

PA Criteria	Criteria Details
Exclusion Criteria	Use during the peri-operative period in the setting of coronary artery bypass graft (CABG) surgery
Required Medical Information	Diagnosis of covered use defined as the relief of pain of osteoarthritis of joints amenable to topical treatment, such as the knees and hands.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Not evaluated for use on joints of the spine, hip, or shoulder and therefore requests for use on these areas will not be approved.
Indications	All FDA-approved Indications.
Off Label Uses	

# DICLOFENAC 1.5% TOPICAL SOLUTION

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## Products Affected

- *diclofenac sodium external solution 1.5 %*

PA Criteria	Criteria Details
Exclusion Criteria	Use during the peri-operative period in the setting of coronary artery bypass graft (CABG) surgery
Required Medical Information	Diagnosis of covered use defined as the relief of pain of osteoarthritis of the knees.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. FDA-approved only for use on knee joints and therefore requests for other uses will not be approved.
Indications	All FDA-approved Indications.
Off Label Uses	

## DICLOFENAC 3% GEL

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### Products Affected

- *diclofenac sodium external gel*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to dermatology
Coverage Duration	90 days
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DICLOFENAC PATCH

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## Products Affected

- *diclofenac epolamine external*

PA Criteria	Criteria Details
Exclusion Criteria	Treatment of peri-operative pain in the setting of coronary artery bypass graft (CABG) surgery, use on non-intact or damaged skin resulting from any etiology including exudative dermatitis, eczema, infection lesions, burns, or wounds, pregnancy after 30 weeks gestation
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	PA applies to all. Product is approved for acute pain, defined as short-term pain not lasting longer than a 3-month period.
Indications	All FDA-approved Indications.
Off Label Uses	



## DIFICID (fidaxomicin)

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### Products Affected

- DIFICID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	10 days
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DIGOXIN IN OLDER PATIENTS

## Products Affected

- *digitek oral tablet 250 mcg*
- *digox oral tablet 250 mcg*
- *digoxin oral tablet 250 mcg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance) with result greater than or equal to 30 mL/min. Patient must have tried and failed to respond adequately to 0.125 mg of digoxin.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	PA not required for cardiology
Coverage Duration	1 year
Other Criteria	PA applies to all except cardiology. PA not required for doses less than or equal to 0.125 mg per day. Updated creatinine clearance since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DOPTELET (avatrombopag)

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## Products Affected

- DOPTELET ORAL TABLET 20 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For immune thrombocytopenia, submission of prior therapies tried and failed.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to gastroenterology, hematology, hepatology, and surgery
Coverage Duration	5 days for undergoing a procedure or 1 year for immune thrombocytopenia
Other Criteria	PA applies to all. This medication should not be administered to patients with chronic liver disease not scheduled to undergo a procedure in an attempt to normalize platelet counts and will not be approved for this indication.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DRONABINOL

## Products Affected

- *dronabinol*
- SYNDROS

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. If authorization is requested for treatment of nausea and vomiting associated with cancer therapy, documentation of previous conventional antiemetic therapies utilized is required.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. If the medication is being administered related to cancer treatment and is a full replacement for intravenous administration of antiemetic therapy within 48 hours of cancer treatment, it is covered as a Part B benefit. To be eligible for Part B coverage, the prescribing physician must indicate this on the prescription. If the medication is being requested for the use of anorexia associated with weight loss in patients with AIDS, approval may be covered under Part D.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DROXIDOPA

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## Products Affected

- *droxidopa*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## DUOBRII (halobetasol/tazarotene)

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### Products Affected

- DUOBRII

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential, documentation patient tried and failed augmented betamethasone dipropionate, clobetasol, fluocinonide 0.1%, halobetasol, or another Class I ultra-high potency topical steroid.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to dermatology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## DUOPA (carbidopa/levodopa enteral suspension)

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### Products Affected

- DUOPA ENTERAL

PA Criteria	Criteria Details
Exclusion Criteria	Administration of non-selective monoamine oxidase inhibitors within 14 days of initiation
Required Medical Information	Diagnosis of covered use, confirmation patient has a naso-jejunal tube for short-term administration or a PEG-J for long-term administration.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# DUPIXENT (dupilumab)

## Products Affected

- DUPIXENT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For atopic dermatitis, documentation of treatment with at least a moderate strength topical corticosteroid for at least four weeks, a contraindication to the use of topical corticosteroids, or therapy is not otherwise advisable. For moderate-to-severe asthma, either (1) submission of blood eosinophil count of at least 150 cells/mcL obtained within 6 weeks of therapy initiation or (2) documentation asthma requires daily oral corticosteroid for control. For chronic rhinosinusitis with nasal polyposis, documentation of treatment with an intranasal corticosteroid for at least three months, a contraindication to the use of intranasal corticosteroids, or therapy is not otherwise advisable.
Age Restrictions	
Prescriber Restrictions	Restricted to allergy, dermatology, immunology, otolaryngology/otorhinolaryngology, and pulmonology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Continuation requires documentation of a positive response to therapy. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	



## EGRIFTA SV (tesamorelin)

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### Products Affected

- EGRIFTA SV

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, active malignancy, disruption of HPA axis due to hypophysectomy, hypopituitarism, pituitary tumor/surgery, head irradiation, or head trauma, use for weight loss
<b>Required Medical Information</b>	Diagnosis of covered use. Submission of pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Continuation of therapy requests require confirmation that the patient has demonstrated a clinical improvement (or maintenance of improvement once achieved) from baseline.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## EMFLAZA (deflazacort)

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### Products Affected

- EMFLAZA

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, documentation of treatment failure with or intolerance to prednisone.
Age Restrictions	2 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# EMPAVELI (pegcetacoplan)

## Products Affected

- EMPAVELI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use confirmed by high-sensitivity flow cytometry, proof of vaccination against Streptococcus pneumoniae, Neisseria meningitidis, and Haemophilus influenzae type B or 2 weeks of antibacterial drug prophylaxis if the vaccines were administered within the last 2 weeks and therapy is required immediately, submission of lactate dehydrogenase level.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology, immunology, and nephrology
Coverage Duration	1 year
Other Criteria	Because this medication is delivered subcutaneously through an infusion pump, it covered as a Part B benefit except for enrollees residing in a long-term care facility. PA applies to all when covered as a Part D benefit.
Indications	All Medically-accepted Indications.
Off Label Uses	

## EMSAM (selegiline transdermal)

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### Products Affected

- EMSAM

PA Criteria	Criteria Details
Exclusion Criteria	Pheochromocytoma, coadministration with carbamazepine or serotonergic drugs
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ENDOTHELIN RECEPTOR ANTAGONISTS

## Products Affected

- *ambrisentan*
- *bosentan*
- OPSUMIT
- TRACLEER ORAL TABLET SOLUBLE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy. For ambrisentan, idiopathic pulmonary fibrosis and moderate or severe hepatic impairment.
<b>Required Medical Information</b>	Diagnosis of covered use, pregnancy status for female patients of childbearing potential, submission of baseline AST, ALT, and bilirubin. For ambrisentan and Opsumit, submission of baseline hemoglobin level.
<b>Age Restrictions</b>	For ambrisentan and Opsumit, 18 years of age or older. For bosentan, 3 years of age or older.
<b>Prescriber Restrictions</b>	Restricted to cardiology and pulmonology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# ENSPRYNG (satralizumab-mwge)

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## Products Affected

- ENSPRYNG

PA Criteria	Criteria Details
Exclusion Criteria	Active hepatitis B infection, active or untreated latent tuberculosis (TB)
Required Medical Information	Diagnosis of covered use, submission of confirmation patient has anti-aquaporin-4 (AQP4) antibody-positive NMOSD, submission of baseline latent TB screening test (Mantoux tuberculin skin test [a.k.a. PPD test] or interferon-gamma release assay [IGRA]), attestation patient does not have any active infection.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology and ophthalmology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# EPIDIOLEX (cannabidiol)

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## Products Affected

- EPIDIOLEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	1 year of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ERIVEDGE (vismodegib)

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### Products Affected

- ERIVEDGE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# ERLOTINIB

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## Products Affected

- *erlotinib hcl*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For non-small cell lung cancer, submission of FDA-approved test confirming presence of EGFR exon 19 deletion or exon 21 L858R substitution mutation and prior treatments used.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ESBRIET (pirfenidone)

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### Products Affected

- ESBRIET ORAL CAPSULE
- *pirfenidone oral tablet 267 mg, 801 mg*

PA Criteria	Criteria Details
Exclusion Criteria	End-stage renal disease on dialysis, severe (Child-Pugh class C) hepatic impairment
Required Medical Information	Diagnosis of covered use, submission of liver function tests or Child-Pugh status.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated liver function testing or Child-Pugh score since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# EVEROLIMUS

## Products Affected

- *everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg*
- *everolimus oral tablet soluble*

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with dual strong CYP3A4/P-glycoprotein inhibitors
Required Medical Information	Diagnosis of covered use and submission of pregnancy status for female patients of childbearing potential. For renal cell carcinoma, documented prior use of sunitinib or sorafenib. For postmenopausal women with advanced hormone receptor-positive, HER-2 negative breast cancer, documentation of treatment failure with letrozole or anastrozole and confirmation drug is being used in combination with exemestane.
Age Restrictions	1 year of age or older
Prescriber Restrictions	Restricted to neurology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## EVRYSDI (risdiplam)

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### Products Affected

- EVRYSDI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use confirmed by genetic testing including either (a) homozygous deletion of SMN1 exon 7 or (b) compound heterozygosity for SMN1 exon 7 deletion and small mutation, documentation of two or more copies of the SMN2 gene by genetic testing, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	
Prescriber Restrictions	Restricted to neurology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Maintenance of or improvement in any motor score or function compared to baseline will be required for reauthorization.
Indications	All Medically-accepted Indications.
Off Label Uses	

## EXKIVITY (mobocertinib)

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### Products Affected

- EXKIVITY

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A4 inducers, coadministration with strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of EGFR exon 20 insertion mutation and previous therapies used, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Part B before Part D Step Therapy. For approval, the patient must have documentation of failure or contraindication to both platinum-based chemotherapy and amivantamab.
Indications	All Medically-accepted Indications.
Off Label Uses	

# FENTANYL TRANSMUCOSAL

## Products Affected

- *fentanyl citrate buccal*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Patients not tolerant to the effects of a chronic opioid, treatment of acute or postoperative pain including headache, migraines, or dental pain
<b>Required Medical Information</b>	Diagnosis of covered use, verified claim or documentation of patient's morphine milligram equivalent opioid dose.
<b>Age Restrictions</b>	For the buccal tablet, 18 years of age or older. For the lozenge, 16 years of age or older.
<b>Prescriber Restrictions</b>	PA not required for oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all except oncology.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# FINTEPLA (fenfluramine)

## Products Affected

- FINTEPLA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Hepatic impairment, moderate or severe renal impairment, administration of monoamine oxidase inhibitors within 14 days of initiation
<b>Required Medical Information</b>	Diagnosis of covered use, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance) and liver function testing or Child-Pugh score.
<b>Age Restrictions</b>	2 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## FIRDAPSE (amifampridine)

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### Products Affected

- FIRDAPSE

PA Criteria	Criteria Details
Exclusion Criteria	History of seizure
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# FIRST-GENERATION ANTIHISTAMINES IN OLDER PATIENTS

## Products Affected

- *carbinoxamine maleate oral solution*
- *carbinoxamine maleate oral tablet 4 mg*
- *clemastine fumarate oral tablet 2.68 mg*
- *cyproheptadine hcl oral*
- *diphenhydramine hcl oral elixir*
- *hydroxyzine hcl oral tablet*
- *hydroxyzine pamoate oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For carbinoxamine or cyproheptadine for dermatographism, documentation patient tried and had an inadequate response to a second-generation antihistamine. For hydroxyzine for pruritus, documentation patient tried and had an inadequate response to a second-generation antihistamine. For hydroxyzine for anxiety, documentation patient has tried and had an inadequate response to at least 2 other FDA-approved products for the management of anxiety OR documentation medication is being used as a sedative before and after general anesthesia.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. First-generation antihistamines are anticholinergic medications considered high-risk in older patients due to risks of confusion, dry mouth, constipation, and decreased clearance with advanced age.
Indications	All Medically-accepted Indications.
Off Label Uses	

# FOTIVDA (tivozanib)

## Products Affected

- FOTIVDA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Uncontrolled hypertension, severe hepatic impairment, coadministration with strong CYP3A inducers
<b>Required Medical Information</b>	Diagnosis of covered use, submission of previous systemic therapies used to treat renal cell carcinoma including the failure of at least one prior VEGFR inhibitor, pregnancy status for female patients of childbearing potential, confirmation patient has not had episodes of symptomatic heart failure or unstable angina, a myocardial infarction, an arterial thrombotic event, or a significant bleeding event in the 6 months preceding the prior authorization request.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# FUMARATES FOR MULTIPLE SCLEROSIS

## Products Affected

- BAFIERTAM
- VUMERITY

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Hypersensitivity to dimethyl fumarate, coadministration with another fumarate. For Vumerity, moderate or severe renal impairment.
<b>Required Medical Information</b>	Diagnosis of covered use. For Vumerity, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. For approval of Bafiertam or Vumerity, the patient must have tried and failed to have an adequate response to or had an intolerance to dimethyl fumarate. Updated creatinine clearance since the previous authorization will be required for subsequent annual reauthorizations of Vumerity.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# GALAFOLD (migalastat)

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## Products Affected

- GALAFOLD

PA Criteria	Criteria Details
Exclusion Criteria	Severe renal impairment (eGFR less than 30 mL/min/1.73 m <sup>2</sup> ) or end-stage renal disease requiring dialysis
Required Medical Information	Diagnosis of covered use, documentation that the patient has an amenable galactosidase alpha gene variant (see section 12.1, table 2 of package insert for full list) based on in vitro assay data as interpreted by a clinical genetics professional.
Age Restrictions	16 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# GATTEX (teduglutide)

## Products Affected

- GATTEX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline laboratory values including bilirubin, alkaline phosphatase, lipase, and amylase obtained within 6 months prior to starting therapy. For adults 18 years of age or older only, submission of documentation that a colonoscopy (or alternate imaging) of the entire colon with polyp removal was performed within 6 months prior to starting treatment.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. For adults 18 years of age or older, continuation of therapy requires submission of findings from a follow-up colonoscopy or alternate imaging result at the end of 1 year of teduglutide treatment. Subsequent imaging should be performed every 5 years, or sooner if polyps are found at the 1-year mark.
Indications	All Medically-accepted Indications.
Off Label Uses	

# GAVRETO (pralsetinib)

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## Products Affected

- GAVRETO

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of RET gene fusion or mutation, attestation patient does not have uncontrolled hypertension, pregnancy status for female patients of childbearing potential.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# GILOTRIF (afatinib)

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## Products Affected

- GILOTRIF

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For non-small cell lung cancer, submission of positive FDA-approved test for non-resistant epidermal growth factor receptor mutations. For metastatic, squamous non-small cell lung cancer, documentation of progression after platinum-based chemotherapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# GnRH ANTAGONISTS

## Products Affected

- CAMCEVI
- ELIGARD
- FIRMAGON (240 MG DOSE)
- FIRMAGON SUBCUTANEOUS SOLUTION RECONSTITUTED 80 MG
- *leuprolide acetate injection*
- LUPANETA PACK
- LUPRON DEPOT (1-MONTH)
- LUPRON DEPOT (3-MONTH)
- LUPRON DEPOT (4-MONTH)
- LUPRON DEPOT (6-MONTH)
- LUPRON DEPOT-PED (1-MONTH) INTRAMUSCULAR KIT 11.25 MG, 15 MG
- LUPRON DEPOT-PED (3-MONTH) INTRAMUSCULAR KIT 30 MG (PED)
- TRELSTAR MIXJECT

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential. For prostate cancer, documentation of baseline prostate-specific antigen and serum testosterone level.
Age Restrictions	
Prescriber Restrictions	Restricted to hematology, oncology, endocrinology, gynecology, and urology
Coverage Duration	For endometriosis and uterine fibroids, 6 months. For all other indications, 1 year.
Other Criteria	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	



# GROWTH HORMONE

## Products Affected

- GENOTROPIN
- GENOTROPIN MINIQUICK
- HUMATROPE
- NORDITROPIN FLEXPROM SUBCUTANEOUS SOLUTION PEN-INJECTOR
- NUTROPIN AQ NUSPIN 10 SUBCUTANEOUS SOLUTION PEN-INJECTOR
- NUTROPIN AQ NUSPIN 20 SUBCUTANEOUS SOLUTION PEN-INJECTOR
- NUTROPIN AQ NUSPIN 5 SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation of growth failure, submission of IGF-1 levels, height, weight, creatinine clearance (or serum creatinine), fasting blood glucose, and bone age if applicable based on patient age and diagnosis.
Age Restrictions	
Prescriber Restrictions	Restricted to endocrinology and nephrology
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated IGF-1 level, bone age (if applicable based on patient age and diagnosis) height, weight, creatinine clearance (or serum creatinine), and fasting glucose since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# HEREDITARY ANGIOEDEMA THERAPIES, ACUTE

## Products Affected

- *icatibant acetate*
- RUCONEST
- *sajazir*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Requests for prophylactic hereditary angioedema therapy. For Ruconest, acute laryngeal attacks.
<b>Required Medical Information</b>	Diagnosis of covered use. For Ruconest, documentation of the patient's typical attack presentation/symptoms.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	Restricted to allergy, dermatology, hematology, or immunology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# HEREDITARY ANGIOEDEMA THERAPIES, MAINTENANCE

## Products Affected

- HAEGARDA
- ORLADEYO
- *takhzyro subcutaneous solution*
- TAKHZYRO SUBCUTANEOUS SOLUTION PREFILLED

SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Requests for acute hereditary angioedema therapy (attacks). For Orladeyo, end-stage renal disease.
<b>Required Medical Information</b>	Diagnosis of covered use, submission of objective or subjective documentation that prophylactic therapy is medically necessary, including, but not limited to activity of disease and disease burden, the frequency of HAE attacks, and quality of life.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	Restricted to allergy, dermatology, hematology, or immunology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. For approval of either Haegarda or Orladeyo for patients 12 years of age and older, the patient must have tried and failed to have an adequate response to or had an intolerance or contraindication to Takhzyro. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# HETLIOZ (tasimelteon)

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## Products Affected

- HETLIOZ

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, coadministration with strong CYP1A2 inhibitors or CYP3A4 inducers
Required Medical Information	Diagnosis of covered use. For Smith-Magenis Syndrome patients only, documentation of genetic testing results confirming diagnosis is required.
Age Restrictions	
Prescriber Restrictions	Restricted to neurology and sleep medicine
Coverage Duration	1 year
Other Criteria	PA applies to all. For non-24-hour sleep-wake disorder, patients are required to be totally blind to match the population in which tasimelteon was studied.
Indications	All Medically-accepted Indications.
Off Label Uses	

# IBRANCE (palbociclib)

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## Products Affected

- IBRANCE

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HR-positive, HER2-negative, confirmation that the treatment regimen will include concomitant use of an aromatase inhibitor or fulvestrant, submission of baseline CBC.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ICLUSIG (ponatinib)

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## Products Affected

- ICLUSIG

PA Criteria	Criteria Details
Exclusion Criteria	Newly diagnosed chronic phase CML
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## IDHIFA (enasidenib)

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### Products Affected

- IDHIFA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of IDH2 mutation, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ILARIS

## Products Affected

- ILARIS SUBCUTANEOUS SOLUTION

PA Criteria	Criteria Details
Exclusion Criteria	Positive TB test, coadministration with TNF inhibitors
Required Medical Information	Diagnosis of covered use, submission of TB skin test result obtained within past 12 months, documentation that patient has received all recommended vaccinations as appropriate including pneumococcal vaccine and inactivated influenza vaccine prior to initiation of therapy. For CAPS, confirmed diagnosis including genetic testing for variant FCAS or MWS and documentation patient is not receiving concomitant TNF inhibitor therapy. For SJIA, submission of CBC including platelet count and confirmed diagnosis defined by prominence of systemic and inflammatory features including spiking fevers, rash, swelling and inflammation of lymph nodes, liver, and spleen, and high white blood cell and platelet counts.
Age Restrictions	2 years of age or older
Prescriber Restrictions	For SJIA, restricted to rheumatology
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated TB skin test result obtained within the past 12 months and objective documentation of positive patient response or maintenance of response will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	



# IMBRUVICA (ibrutinib)

## Products Affected

- IMBRUVICA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe hepatic impairment (Child-Pugh class C), coadministration with strong CYP3A inducers
<b>Required Medical Information</b>	Diagnosis of covered use, pregnancy status for female patients of childbearing potential. For chronic graft versus host disease, documentation of treatment failure with any other systemic immunosuppressive agent.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology, oncology, and transplant specialty
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# IMCIVREE (setmelanotide)

## Products Affected

- IMCIVREE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Moderate or severe renal impairment, end-stage renal disease
<b>Required Medical Information</b>	Diagnosis of covered use, including submission of genetic testing showing homozygous or compound heterozygous gene variants in POMC, PCSK1, or LEPR genes interpreted as pathogenic, likely pathogenic, or of uncertain clinical significance and body mass index (BMI) greater than 30 kg/m <sup>2</sup> in adults or greater than the 97th percentile in children.
<b>Age Restrictions</b>	6 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	Initially 16 weeks, then 1 year
<b>Other Criteria</b>	PA applies to all. For re-authorization at the 16-week point, submission of clinical documentation attesting to at least 5% weight loss from baseline (or at least 5% BMI from baseline in patients with continued growth potential) is required. Not FDA-approved for other types or causes of obesity, and therefore requests for these uses will not be approved.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

# IMMUNE GLOBULIN

## Products Affected

- BIVIGAM INTRAVENOUS SOLUTION 5 GM/50ML
- CARIMUNE NF INTRAVENOUS SOLUTION RECONSTITUTED 12 GM, 6 GM
- FLEBOGAMMA DIF
- GAMMAGARD
- GAMMAGARD S/D LESS IGA
- GAMMAKED INJECTION SOLUTION 1 GM/10ML, 10 GM/100ML, 20 GM/200ML, 5 GM/50ML
- GAMMAPLEX INTRAVENOUS SOLUTION 10 GM/100ML, 10 GM/200ML, 20 GM/200ML, 20 GM/400ML, 5 GM/100ML, 5 GM/50ML
- GAMUNEX-C
- OCTAGAM INTRAVENOUS SOLUTION 1 GM/20ML, 10 GM/100ML, 10 GM/200ML, 2 GM/20ML, 2.5 GM/50ML, 20 GM/200ML, 25 GM/500ML, 5 GM/100ML, 5 GM/50ML
- PRIVIGEN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	IgA-deficient patients with antibodies against IgA and a history of hypersensitivity. For IM forms, severe thrombocytopenia or coagulation disorder that would contraindicate an IM injection.
<b>Required Medical Information</b>	Diagnosis of covered use. For ITP, submission of platelet count. For CLL, IgG level less than 600 mg/dL and recent history of serious bacterial infection requiring either oral or IV antibiotic therapy. For CIDP, unequivocal diagnosis and documentation patient is refractory or intolerant to prednisone given in therapeutic doses over at least 3 months. For passive immunization against varicella, confirmation that the patient is immunosuppressed and cannot receive varicella-zoster immune globulin.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	For acute conditions/new starts, 3 months. For renewal of chronic conditions, 1 year.
<b>Other Criteria</b>	PA applies to all. For continuation of any diagnosis, documentation of the clinical response to therapy must be submitted. For IV formulations, covered as a Part B benefit if administered in the home for the treatment of primary immune deficiency. For any other combination of treatment site and indication, additional information may need to be submitted to determine if the immune globulin will be covered as a Part B or Part D benefit.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# INJECTABLE TESTOSTERONE

## Products Affected

- *testosterone cypionate injection solution 200 mg/ml*
- *testosterone cypionate intramuscular solution 100 mg/ml, 200 mg/ml*
- *testosterone enanthate intramuscular solution*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of serum testosterone level, documentation that patient has been evaluated for the presence of breast and prostate cancer prior to initiation of therapy.
Age Restrictions	
Prescriber Restrictions	PA not required for urology or endocrinology
Coverage Duration	1 year
Other Criteria	PA applies to all except when prescribed by urology or endocrinology. Documentation of clinically relevant response to therapy (including, but not limited to submission of updated serum testosterone level) will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# INLYTA (axitinib)

## Products Affected

- INLYTA

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment (Child-Pugh class C), uncontrolled hypertension, evidence of untreated brain metastasis, recent active gastrointestinal bleeding, coadministration with strong CYP3A4/5 inducers
Required Medical Information	Diagnosis of covered use, submission of laboratory values including baseline ALT, AST, bilirubin, submission of baseline blood pressure reading. If axitinib is being used as a single agent, submission of prior therapy or therapies tried and failed.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# INQOVI (decitabine/cedazuridine)

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## Products Affected

- INQOVI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of complete blood count, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to new starts only. Continuation of initial therapy beyond 6 months requires (a) confirmation of no disease progression and (b) attestation the patient is having no serious adverse events from treatment.
Indications	All Medically-accepted Indications.
Off Label Uses	

# INREBIC (fedratinib)

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## Products Affected

- INREBIC

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, coadministration with moderate or strong CYP3A4 inducers or dual CYP3A4/CYP2C19 inhibitors
Required Medical Information	Diagnosis of covered use, submission of thiamine level and platelet count.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For approval, the patient must have tried and failed to have an adequate response to or had an intolerance to ruxolitinib.
Indications	All Medically-accepted Indications.
Off Label Uses	

# INTERLEUKIN-5 ANTAGONISTS

## Products Affected

- FASENRA
- FASENRA PEN
- NUCALA SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- NUCALA SUBCUTANEOUS SOLUTION PREFILLED
- SYRINGE 100 MG/ML
- NUCALA SUBCUTANEOUS SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For eosinophilic asthma, documentation that patient's symptoms are poorly controlled with inhaled corticosteroids, submission of pulmonary function test results including FEV1, frequency of inhaled short-acting beta2-agonist therapy, frequency of daily and nighttime symptoms and exacerbations, and effect of exacerbations on activity. For Nucala (eosinophilic asthma diagnosis only), submission of blood eosinophil count of at least 150 cells/mcL obtained within 6 weeks of therapy initiation or at least 300 cells/mcL within 12 months of therapy initiation. For Nucala (for chronic rhinosinusitis with nasal polyps diagnosis only), documentation of treatment with an intranasal corticosteroid for at least 8 weeks, a contraindication to the use of intranasal corticosteroids, or therapy is not otherwise advisable. For Fasenra, submission of blood eosinophil count of at least 300 cells/mcL obtained within 6 weeks of therapy initiation or at least 150 cells/mcL within 6 weeks of therapy initiation if patient is dependent on a daily oral corticosteroid.
Age Restrictions	
Prescriber Restrictions	Restricted to allergy, hematology, immunology, otorhinolaryngology, pulmonology, and rheumatology
Coverage Duration	1 year
Other Criteria	PA applies to all. Continuation of therapy requests require objective documentation from the prescriber that the patient's symptoms have improved. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	



# INTRANASAL SEIZURE MEDICATIONS

## Products Affected

- NAYZILAM
- VALTOCO 10 MG DOSE
- VALTOCO 15 MG DOSE
- VALTOCO 20 MG DOSE
- VALTOCO 5 MG DOSE

PA Criteria	Criteria Details
Exclusion Criteria	Acute narrow-angle glaucoma
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# INTRON A (interferon alfa-2b)

## Products Affected

- INTRON A

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Autoimmune hepatitis, decompensated liver disease
<b>Required Medical Information</b>	Diagnosis of covered use, submission of triglyceride levels, hemoglobin, complete and differential white blood cell counts, platelet count, serum electrolytes, ALT, serum bilirubin level, serum albumin level, and TSH. For malignant melanoma, submission of the date of surgical treatment. For AIDS-related Kaposi's sarcoma, submission of total CD4 count. For chronic hepatitis C, submission of HCV RNA, prothrombin time, baseline serum creatinine level, and laboratory confirmation of hepatitis C virus, and documentation of previous response to therapy if applicable. For chronic hepatitis B, submission of prothrombin time and documentation patient has been serum HBsAG positive for at least 6 months with evidence of HBV replication.
<b>Age Restrictions</b>	For hairy cell leukemia, malignant melanoma, follicular lymphoma, condylomata acuminata, or AIDS-related Kaposi's sarcoma, 18 years of age or older. For chronic hepatitis C, 3 years of age or older. For chronic hepatitis B, 1 year of age or older.
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	Depends on covered use. See "Other Criteria" section.
<b>Other Criteria</b>	PA applies to new starts only. For hairy cell leukemia, the coverage duration is 6 months. For condylomata acuminata, 3 weeks per course, and at least 12 weeks must pass in between multiple courses in order to be reauthorized. For Kaposi's sarcoma, 16 weeks. For hepatitis B infection, 24 weeks. For all other indications/uses, 1 year. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# INVEGA HAFYERA (paliperidone 6-month injectable suspension)

## Products Affected

- INVEGA HAFYERA

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Dementia-related psychosis
<b>Required Medical Information</b>	Diagnosis of covered use, documentation of at least 4 months' treatment with 1-month paliperidone palmitate extended-release injectable suspension or at least one 3-month injection of 3-month paliperidone palmitate extended-release injectable suspension.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# INVEGA TRINZA (paliperidone 3-month injectable suspension)

## Products Affected

- INVEGA TRINZA INTRAMUSCULAR SUSPENSION  
PREFILLED SYRINGE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Dementia-related psychosis
<b>Required Medical Information</b>	Diagnosis of covered use, documentation of at least 4 months' treatment with 1-month paliperidone palmitate extended-release injectable suspension.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# IRESSA (gefitinib)

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## Products Affected

- IRESSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, confirmation of EGFR exon 19 deletions or exon 21 (L858R) substitution mutations as detected by an FDA-approved test, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ISTURISA (osilodrostat)

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## Products Affected

- ISTURISA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, attestation pituitary gland surgery is not an option for the patient or has not been curative, submission of baseline serum potassium and magnesium levels.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Continuation requires documentation of clinically relevant response to therapy, including, but not limited to urine free cortisol level.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ITRACONAZOLE

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## Products Affected

- *itraconazole oral*
- TOLSURA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, fungal culture result identifying causative organism or positive KOH result.
Age Restrictions	
Prescriber Restrictions	PA not required for infectious diseases
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# JAKAFI (ruxolitinib)

## Products Affected

- JAKAFI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	For myelofibrosis, a platelet count less than $50 \times 10^9/L$ with either concomitant estimated creatinine clearance between 15 and 59 mL/min, end-stage renal disease not on dialysis, or any degree of hepatic impairment
<b>Required Medical Information</b>	Diagnosis of covered use, submission of baseline platelet count, ALT, AST, and bilirubin, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance). For polycythemia vera, documented intolerance or inadequate response to hydroxyurea. For acute graft-versus-host disease, documented inadequate response to systemic corticosteroids.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	Restricted to hematology, oncology, and transplant specialty
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# JUXTAPID (lomitapide)

## Products Affected

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

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, moderate or severe hepatic impairment (Child-Pugh class B or C), active liver disease, coadministration with moderate or strong CYP3A4 inhibitors
<b>Required Medical Information</b>	Diagnosis of covered use, submission of baseline lab values including ALT, AST, alkaline phosphatase, total bilirubin, baseline LDL-C, total cholesterol (TC), apoB, and non-HDL-C, pregnancy status for female patients of childbearing potential, submission of renal indices, documentation of contraindication to or treatment failure with evolocumab.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to cardiology, lipidology, and endocrinology with experience in and a focus on lipid management
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. Submission of LDL level drawn after the initial LDL level submission documenting clinically significant response to therapy will be required for reauthorization. For approval, the patient must have tried and failed to have an adequate response to, had an intolerance to, or have a contraindication to therapy with evolocumab.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# JYNARQUE (tolvaptan)

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## Products Affected

- JYNARQUE

PA Criteria	Criteria Details
Exclusion Criteria	History of signs or symptoms of significant liver impairment or injury (not including uncomplicated polycystic liver disease), uncorrected abnormal blood sodium concentrations, inability to sense or respond to thirst, hypovolemia, uncorrected urinary outflow obstruction, anuria, coadministration with strong CYP3A inhibitors or inducers or desmopressin
Required Medical Information	Diagnosis of covered use, submission of serum sodium concentration.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to nephrology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# KALYDECO (ivacaftor)

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## Products Affected

- KALYDECO

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of cystic fibrosis mutation test result and baseline ALT and AST.
Age Restrictions	
Prescriber Restrictions	Restricted to pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# KERENDIA (finerenone)

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## Products Affected

- KERENDIA

PA Criteria	Criteria Details
Exclusion Criteria	Adrenal insufficiency, estimated glomerular filtration rate (eGFR) less than 25 mL/min/1.73 m <sup>2</sup> , severe (Child-Pugh class C) hepatic impairment, coadministration with strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of estimated glomerular filtration rate (eGFR) and baseline serum potassium level.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. For approval, the patient must have documentation of a trial of Farxiga.
Indications	All Medically-accepted Indications.
Off Label Uses	

# KETOCONAZOLE ORAL

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## Products Affected

- *ketoconazole oral*

PA Criteria	Criteria Details
Exclusion Criteria	Acute or chronic liver disease, treatment of fungal meningitis or fungal infections of the skin or nails
Required Medical Information	Ketoconazole is being requested for the treatment of culture-proven systemic blastomycosis, coccidioidomycosis, histoplasmosis, chromomycosis, or paracoccidioidomycosis, submission of baseline ALT, AST, total bilirubin, alkaline phosphatase, prothrombin time and INR, confirmation from the prescriber that the potential benefits of therapy outweigh the risks.
Age Restrictions	2 years of age or older
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## KEVEYIS (dichlorphenamide)

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### Products Affected

- KEVEYIS

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of high dose aspirin, severe pulmonary disease limiting compensation to metabolic acidosis, hepatic insufficiency
Required Medical Information	Diagnosis of covered use, submission of serum potassium.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	Initially 2 months, then 1 year
Other Criteria	PA applies to all. Documentation of a positive response to therapy will be required for initial reauthorization after the first 2 months. Maintenance of a clinical benefit will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# KISQALI (ribociclib)

## Products Affected

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

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Congenital long QT syndrome, QTcF interval greater than 450 msec at treatment initiation, uncorrected hypokalemia or hypomagnesemia, coadministration with strong CYP3A4 inducers or drugs that can prolong the QT interval
<b>Required Medical Information</b>	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HR-positive, HER2-negative, submission of ECG, serum electrolytes, CBC, and pregnancy status for female patients of childbearing potential. For patients receiving Kisqali alone, confirmation that the treatment regimen will include concomitant use of an aromatase inhibitor or fulvestrant.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# KORLYM (mifepristone)

## Products Affected

- KORLYM

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, severe hepatic impairment, uncorrected hypokalemia, female patients with a history of unexplained vaginal bleeding, endometrial hyperplasia with atypia, or endometrial carcinoma, patients using systemic corticosteroids for life-saving purposes, coadministration with strong CYP3A4 inducers, simvastatin, lovastatin, or CYP3A substrates with narrow therapeutic ranges
<b>Required Medical Information</b>	Diagnosis of covered use, attestation surgery is not an option for the patient or has not been curative, submission of baseline serum potassium, AST, ALT, and alkaline phosphatase, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to endocrinology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# KOSELUGO (selumetinib)

## Products Affected

- KOSELUGO

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with moderate or strong CYP3A4 inducers
<b>Required Medical Information</b>	Diagnosis of covered use, submission of left ventricular ejection fraction, Child-Pugh score or liver function testing results, and pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	Initiation: 2-17 years of age. Continuation: 2 years of age or older.
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to new starts only. Selumetinib is indicated in pediatric patients and will not be approved for adults unless the patient started on the medication prior to 18 years of age. Continuation of initial therapy beyond 6 months requires (a) documentation of any positive clinical response and (b) attestation the patient is having no serious adverse events to treatment.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# LAPATINIB

## Products Affected

- *lapatinib ditosylate*

PA Criteria	Criteria Details
Exclusion Criteria	Uncorrected hypokalemia, uncorrected hypomagnesemia
Required Medical Information	Diagnosis of covered use, submission of baseline left ventricular ejection fraction, potassium and magnesium levels, pregnancy status for female patients of childbearing potential, and depending on indication, confirmation that the treatment regimen will include concomitant use of either capecitabine or letrozole. For patients who will be using lapatinib with capecitabine, submission of prior therapies tried and failed.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LEDIPASVIR/SOFOSBUVIR

## Products Affected

- HARVONI ORAL PACKET
- HARVONI ORAL TABLET 45-200 MG, 90-400 MG
- *ledipasvir-sofosbuvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, laboratory confirmation of hepatitis C virus (HCV) genotype 1a, 1b, 4, 5, or 6 infection, submission of baseline HCV RNA level, documentation of whether cirrhosis is present or not and whether or not it is compensated or decompensated, confirmation of whether patient is treatment-naïve or treatment-experienced, confirmation a test for HBV infection (HBsAg and anti-HBc) was completed.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Treatment-experienced pts w/genotype 1 and compensated cirrhosis, 24 weeks. All others, 12 weeks.
Other Criteria	PA applies to all. For treatment-naïve patients without cirrhosis who have pre-treatment HCV RNA less than 6 million IU/mL, 8 weeks of therapy may be considered by the provider. For approval of brand Harvoni 90 mg/400 mg, the patient must have tried and failed to have an adequate response to or had an intolerance to ledipasvir/sofosbuvir 90 mg/400 mg.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LENVIMA (lenvatinib)

## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Uncorrected electrolyte abnormalities, uncontrolled hypertension
Required Medical Information	Diagnosis of covered use, submission of baseline blood pressure, comprehensive metabolic panel, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LEUKINE (sargramostim, GM-CSF)

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## Products Affected

- LEUKINE INJECTION SOLUTION RECONSTITUTED

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of WBC count and ANC.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	3 months
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LIDOCAINE TRANSDERMAL PATCHES

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## Products Affected

- *lidocaine external patch 5 %*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. FDA-approved only for postherpetic neuralgia. Requests for other indications will not be approved.
Indications	All FDA-approved Indications.
Off Label Uses	

## LIVMARLI (maralixibat)

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### Products Affected

- LIVMARLI

PA Criteria	Criteria Details
Exclusion Criteria	History of liver transplant, clinical evidence of decompensated cirrhosis
Required Medical Information	Diagnosis of covered use confirmed by molecular genetic testing, attestation drug-induced pruritus has been ruled out, attestation patient has tried and failed at least two of the following medications for pruritus: ursodiol, cholestyramine, naltrexone, rifampin.
Age Restrictions	
Prescriber Restrictions	Restricted to gastroenterology and hepatology
Coverage Duration	1 year
Other Criteria	PA applies to all. Attestation of improvement in pruritus symptoms and confirmation the patient has not progressed to portal hypertension or has had a hepatic decompensation event since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LIVTENCITY (maribavir)

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## Products Affected

- LIVTENCITY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of previous anti-CMV medication(s) patient has tried and failed (at least one of cidofovir, foscarnet, ganciclovir, valganciclovir), documented history of hematopoietic stem cell or solid organ transplant.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to hematology, infectious disease specialty, oncology, and transplant specialty
Coverage Duration	8 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# LONSURF (trifluridine/tipiracil)

## Products Affected

- LONSURF

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Moderate or severe hepatic impairment, febrile neutropenia
<b>Required Medical Information</b>	Diagnosis of covered use, submission of prior therapies used, submission of baseline CBC, absolute neutrophil count, ALT, AST, and bilirubin, pregnancy status for female patients of childbearing potential, documentation of KRAS status.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# LORBRENA (lorlatinib)

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## Products Affected

- LORBRENA ORAL TABLET 100 MG, 25 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers, uncontrolled hypertension
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of ALK-positive tumor, baseline blood pressure, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LUCEMYRA (lofexidine)

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## Products Affected

- LUCEMYRA

PA Criteria	Criteria Details
Exclusion Criteria	Congenital long QT syndrome, severe coronary insufficiency, recent myocardial infarction, cerebrovascular disease, chronic renal failure
Required Medical Information	Diagnosis of covered use, attestation patient will be using medication in conjunction with a comprehensive management program for the treatment of opioid use disorder.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	14 days
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LUMAKRAS (sotorasib)

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## Products Affected

- LUMAKRAS

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers, coadministration with proton pump inhibitors or H2 receptor antagonists
Required Medical Information	Diagnosis of covered use, submission of test result confirming presence of KRAS G12C mutations, submission of previous systemic treatment(s) tried.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LUPKYNIS (voclosporin)

## Products Affected

- LUPKYNIS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe hepatic impairment, coadministration with strong CYP3A4 inhibitors, moderate or strong CYP3A4 inducers, or cyclophosphamide, hypertensive emergency or a baseline blood pressure above 165/105 mmHg
<b>Required Medical Information</b>	Diagnosis of covered use, attestation patient will be taking concurrently with mycophenolate mofetil and corticosteroids, submission of estimated glomerular filtration rate (eGFR), baseline blood pressure, pregnancy status for female patients of childbearing potential. If the patient's eGFR is less than or equal to 45 mL/min/1.73 m <sup>2</sup> , attestation that prescriber believes benefits of therapy outweigh the potential risks to the patient.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. For approval, the patient must have tried and failed to have an adequate response to or had an intolerance/contraindication to Benlysta (belimumab). Documentation of a positive response to therapy will be required for initial reauthorization after the first 6 months. Maintenance of a clinical benefit, attestation that prescriber believes benefits of continuing therapy outweigh the potential risks to the patient, and updated estimated glomerular filtration rate (eGFR) and blood pressure since the previous authorization will be required for subsequent annual reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# LYBALVI (olanzapine/samidorphan)

## Products Affected

- LYBALVI

PA Criteria	Criteria Details
Exclusion Criteria	Dementia-related psychosis, coadministration with opioids or strong CYP3A inducers, acute opioid withdrawal, end-stage renal disease
Required Medical Information	Diagnosis of covered use, confirmation patient has previously tried and failed, had an intolerance to, or had a contraindication to at least one generic second-generation antipsychotic with low incidence of metabolic side effects (e.g., aripiprazole, ziprasidone), attestation patient has had a trial of generic olanzapine with documentation showing a positive therapeutic benefit but unacceptable weight gain (greater than or equal to a 7% gain from baseline body weight) while using olanzapine.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Reduction in or stabilization of body weight since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# LYNPARZA (olaparib)

## Products Affected

- LYNPARZA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of applicable mutations and previous therapies tried and failed depending on cancer type as necessary, submission of baseline CBC.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology and urology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MAVENCLAD (cladribine)

## Products Affected

- MAVENCLAD (10 TABS)
- MAVENCLAD (4 TABS)
- MAVENCLAD (5 TABS)
- MAVENCLAD (6 TABS)
- MAVENCLAD (7 TABS)
- MAVENCLAD (8 TABS)
- MAVENCLAD (9 TABS)

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Current malignancy, pregnancy, HIV or other active chronic infection (e.g., hepatitis or tuberculosis), lymphocyte count below normal limit before first course or less than 800 cells/microliter before second course, creatinine clearance below 60 mL/min, Child-Pugh score greater than 6
<b>Required Medical Information</b>	Diagnosis of covered use, pregnancy status for female patients of childbearing potential, submission of lymphocyte count, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Documentation of a positive response to therapy, confirmation the patient has no active infection, and updated lymphocyte count and creatinine clearance since the previous authorization will be required for reauthorization. After the completion of 2 treatment courses (2 years' treatment), additional treatment courses are not recommended over the following 2 years because of malignancy risk. Re-initiating treatment after those 2 years have passed has not been studied. Requests for therapy for a combined total of greater than 2 years will not be approved.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# MAVYRET (glecaprevir/pibrentasvir)

## Products Affected

- MAVYRET

PA Criteria	Criteria Details
Exclusion Criteria	Moderate or severe hepatic impairment (Child-Pugh class B or C), coadministration with rifampin or atazanavir
Required Medical Information	Diagnosis of covered use, laboratory confirmation of hepatitis C virus (HCV), documentation of whether cirrhosis is present or not and whether or not it is compensated or decompensated, confirmation a test for HBV infection (HBsAg and anti-HBc) was completed.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Criteria for coverage duration will be applied consistent with current AASLD-IDSA guidance.
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MECASERMIN

## Products Affected

- INCRELEX

PA Criteria	Criteria Details
Exclusion Criteria	Patients with closed epiphyses
Required Medical Information	Diagnosis of covered use, documentation of primary insulin-like growth factor (IGF-1) deficiency or growth hormone gene deletion in patients who have developed neutralizing antibodies to growth hormone, submission of IGF-1 level and growth hormone level.
Age Restrictions	
Prescriber Restrictions	Restricted to endocrinology and nephrology
Coverage Duration	6 months
Other Criteria	PA applies to all. Updated IGF-1 and growth hormone levels since the previous authorization will be required for subsequent reauthorizations. Mecasermin is not indicated as a growth hormone replacement and will not be approved for this indication.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MEGESTROL IN OLDER PATIENTS

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## Products Affected

- *megestrol acetate oral suspension 40 mg/ml, 400 mg/10ml, 625 mg/5ml*

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	PA not required for hematology or oncology
Coverage Duration	1 year
Other Criteria	PA applies to all except hematology and oncology.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MEKINIST (trametinib)

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## Products Affected

- MEKINIST

PA Criteria	Criteria Details
Exclusion Criteria	Progression of disease on prior BRAF-inhibitor therapy
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of BRAF V600E or V600K mutation. For non-small cell lung cancer and thyroid cancer, attestation that therapy will be used in combination with dabrafenib.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# METHAMPHETAMINE

## Products Affected

- *methamphetamine hcl*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Use for exogenous obesity, patients with glaucoma, advanced arteriosclerosis, symptomatic cardiovascular disease, moderate to severe hypertension, hyperthyroidism, or a history of drug abuse, administration of non-selective monoamine oxidase inhibitors within 14 days of initiation
<b>Required Medical Information</b>	Diagnosis of covered use. For patients 65 years of age and older, attestation provider is aware medication is considered a high-risk medication for elderly patients according to the Centers for Medicare and Medicaid Services (CMS) and that the benefits of methamphetamine therapy outweigh the potential risks to the patient.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. PA will not be authorized if using for exogenous obesity (excluded category per CMS).
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# METHOTREXATE INJECTABLE (SUBCUTANEOUS)

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## Products Affected

- OTREXUP SUBCUTANEOUS SOLUTION AUTO-INJECTOR 10 MG/0.4ML, 12.5 MG/0.4ML, 15 MG/0.4ML, 17.5 MG/0.4ML, 20 MG/0.4ML, 22.5 MG/0.4ML, 25 MG/0.4ML
- RASUVO SUBCUTANEOUS SOLUTION AUTO-INJECTOR 10 MG/0.2ML, 12.5 MG/0.25ML, 15 MG/0.3ML, 17.5 MG/0.35ML, 20 MG/0.4ML, 22.5 MG/0.45ML, 25 MG/0.5ML, 30 MG/0.6ML, 7.5 MG/0.15ML

- REDITREX

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use, documentation of intolerance or inadequate response to oral or non-subcutaneous injectable forms of methotrexate.
Age Restrictions	
Prescriber Restrictions	Restricted to rheumatology and dermatology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# METHYLTESTOSTERONE

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## Products Affected

- METHITEST
- *methyltestosterone oral*

PA Criteria	Criteria Details
Exclusion Criteria	Male patients with breast or prostate cancer, women who are or may become pregnant
Required Medical Information	Diagnosis of covered use. For patients 65 years of age and older, attestation provider is aware medication is considered a high-risk medication for elderly patients according to the Centers for Medicare and Medicaid Services (CMS) and that the benefits of methyltestosterone therapy outweigh the potential risks to the patient.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# MIGLUSTAT

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## Products Affected

- *miglustat*

PA Criteria	Criteria Details
Exclusion Criteria	Severe renal impairment (CrCl less than 30 mL/min)
Required Medical Information	Diagnosis of covered use, documentation that enzyme replacement is not a therapeutic option.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MYALEPT (metreleptin)

## Products Affected

- MYALEPT

PA Criteria	Criteria Details
Exclusion Criteria	General obesity not associated with congenital leptin deficiency
Required Medical Information	Diagnosis of covered use, submission of leptin level laboratory test result confirming leptin deficiency, baseline HbA1c, fasting glucose, fasting triglyceride levels, and weight.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated patient weight, HbA1c, fasting glucose, and fasting triglyceride levels since the previous authorization will be required for subsequent annual reauthorizations. Metreleptin is not established as a treatment for nonalcoholic steatohepatitis, complications of partial lipodystrophy, HIV-related lipodystrophy, or metabolic disease without generalized lipodystrophy, and submissions for these uses will not be approved.
Indications	All Medically-accepted Indications.
Off Label Uses	

## MYCAPSSA (octreotide)

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### Products Affected

- MYCAPSSA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of prior use of either injectable octreotide or lanreotide and attestation to its successful treatment of acromegaly using clinical biomarkers or chart notes.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# MYFEMBREE (relugolix/estradiol/norethindrone)

## Products Affected

- MYFEMBREE

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, known liver impairment or disease, known osteoporosis, undiagnosed abnormal uterine bleeding, women at increased risk of or current/a history of thrombotic or thromboembolic disorders (including women over 35 years of age who smoke and women with uncontrolled hypertension), current/history of breast cancer or other hormone-sensitive cancer
<b>Required Medical Information</b>	Diagnosis of covered use, attestation patient is premenopausal, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	Restricted to endocrinology and gynecology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Use of this drug for more than 2 years increases risk of bone loss and requests for therapy for more than 2 years will not be approved.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# MYTESI (crofelemer)

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## Products Affected

- MYTESI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, attestation infectious causes of diarrhea have been ruled out.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## NAMZARIC (memantine and donepezil)

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### Products Affected

- NAMZARIC

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of documentation that the patient has been stabilized on donepezil 10 mg daily.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# NATPARA (parathyroid hormone)

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## Products Affected

- NATPARA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation that (albumin-corrected) serum calcium is greater than 7.5 mg/dL and confirmation that 25-hydroxyvitamin D stores are sufficient.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## NERLYNX (neratinib)

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### Products Affected

- NERLYNX

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with proton pump inhibitors, strong CYP3A4 inhibitors, moderate CYP3A4 and P-glycoprotein dual inhibitors, or moderate or strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HER2-positive, confirmation member has completed adjuvant trastuzumab-based therapy or will be using in combination with capecitabine, submission of baseline liver function tests, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



## NEXAVAR (sorafenib)

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### Products Affected

- *sorafenib tosylate*

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# NINLARO (ixazomib)

## Products Affected

- NINLARO

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with strong CYP3A4 inducers
<b>Required Medical Information</b>	Diagnosis of covered use, documentation that medication will be administered concomitantly with lenalidomide and dexamethasone, documentation of prior therapy regimen for multiple myeloma, submission of baseline platelet count, absolute neutrophil count, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology and oncology
<b>Coverage Duration</b>	Initially 3 months, then 1 year
<b>Other Criteria</b>	PA applies to new starts only. For continuation, documentation of platelet count greater than 30,000/mm <sup>3</sup> , ANC greater than 500/mm <sup>3</sup> , and Grade 1 or lower non-hematological toxicities (including rash, peripheral neuropathies) required.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# NITISINONE

## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Updated liver function tests, urine succinylacetone levels, alpha-fetoprotein level, serum tyrosine level, and serum phenylalanine level since the previous authorization will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# NUEDEXTA (dextromethorphan and quinidine)

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## Products Affected

- NUEDEXTA

PA Criteria	Criteria Details
Exclusion Criteria	Prolonged QT interval, congenital long QT syndrome, heart failure, history suggestive of torsades de pointes, AV block without implanted pacemaker, administration of non-selective monoamine oxidase inhibitors within 14 days of initiation
Required Medical Information	Diagnosis of covered use, submission of ECG (specifically QT interval).
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology and psychiatry
Coverage Duration	1 year
Other Criteria	PA applies to all. The medication will not be approved for agitation or Alzheimer's disease without pseudobulbar affect as this is considered an off-label use.
Indications	All Medically-accepted Indications.
Off Label Uses	

# NUPLAZID (pimavanserin)

## Products Affected

- NUPLAZID ORAL CAPSULE
- NUPLAZID ORAL TABLET 10 MG

PA Criteria	Criteria Details
Exclusion Criteria	Dementia-related psychosis unrelated to Parkinson's disease psychosis, cardiac arrhythmias, symptomatic bradycardia, congenital QT prolongation, coadministration with moderate or strong CYP3A4 inducers, drugs that prolong the QT interval, hypokalemia, hypomagnesemia
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# OCALIVA (obeticholic acid)

## Products Affected

- OCALIVA ORAL TABLET 10 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	Complete biliary obstruction, decompensated cirrhosis (Child-Pugh B or C) or prior decompensation event, compensated cirrhosis with evidence of portal hypertension (e.g., ascites, gastroesophageal varices, persistent thrombocytopenia)
Required Medical Information	Diagnosis of covered use, documentation either (1) drug will be used in combination with ursodeoxycholic acid (UDCA) and UDCA has been used for 1 year or (2) patient had intolerance to UDCA, submission of baseline LFTs including ALP and total bilirubin, attestation patient does not have evidence of portal hypertension and has not had a prior decompensation event.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	Initially 3 months, then 1 year
Other Criteria	PA applies to all. Updated ALP obtained within the previous 3 months will be required for subsequent authorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ODOMZO (sonidegib)

### Products Affected

- ODOMZO

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, coadministration with strong CYP3A4 inhibitors or moderate or strong CYP3A4 inducers
<b>Required Medical Information</b>	Diagnosis of covered use, attestation patient is not a candidate for surgery or radiation therapy or carcinoma has recurred following surgery or radiation therapy, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## OFEV (nintedanib)

### Products Affected

- OFEV

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Moderate or severe (Child-Pugh class B or C) hepatic impairment, coadministration of a dual P-glycoprotein/CYP3A4 inducer
<b>Required Medical Information</b>	Diagnosis of covered use, submission of liver function tests or Child-Pugh status, pregnancy status in female patients of childbearing potential. For chronic fibrosing interstitial lung diseases with a progressive phenotype and systemic sclerosis-associated interstitial lung disease diagnoses, submission of HRCT scan showing fibrosis affecting at least 10% of the lungs within the previous 12 months.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to pulmonology or rheumatology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Updated liver function testing or Child-Pugh score since the previous authorization will be required for subsequent reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# ONUREG (azacitidine)

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## Products Affected

- ONUREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of absolute neutrophil count, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to new starts only. Attestation of clinical benefit or stabilization and absence of unacceptable toxicity will be required for reauthorization. This dosage form is not intended to be a substitute for or substituted for injectable azacitidine.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ORENITRAM (treprostinil)

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## Products Affected

- ORENITRAM

PA Criteria	Criteria Details
Exclusion Criteria	Moderate or severe (Child-Pugh class B or C) hepatic impairment
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology and pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ORGOVYX (relugolix)

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## Products Affected

- ORGOVYX

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology and urology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ORIAHNN (elagolix/estradiol/norethindrone)

### Products Affected

- ORIAHNN

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, known liver impairment or disease, known osteoporosis, undiagnosed abnormal uterine bleeding, women at increased risk of or a history of thrombotic or thromboembolic disorders (including women over 35 years of age who smoke and women with uncontrolled hypertension), current/history of breast cancer or other hormone-sensitive cancer, coadministration with OATP1B1 inhibitors or strong CYP3A4 inhibitors
<b>Required Medical Information</b>	Diagnosis of covered use, attestation patient is premenopausal, submission of baseline blood pressure, pregnancy status for female patients of childbearing potential.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to endocrinology and gynecology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Use of this drug for more than 2 years increases risk of bone loss and requests for therapy for more than 2 years will not be approved.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# ORILISSA (elagolix)

## Products Affected

- ORILISSA ORAL TABLET 150 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy, severe hepatic impairment (Child-Pugh class C), known osteoporosis, coadministration with OATP1B1 inhibitors
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology and gynecology
Coverage Duration	Up to 24 months based on liver function and coexisting dyspareunia. See "Other Criteria" section.
Other Criteria	PA applies to all. For endometriosis with dyspareunia or in women with moderate hepatic impairment, 6 months. For endometriosis without dyspareunia, 150 mg daily for 24 months. Use of this drug for more than 2 years increases risk of bone loss and requests for therapy for more than 2 years will not be approved.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ORKAMBI (lumacaftor/ivacaftor)

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### Products Affected

- ORKAMBI

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, documentation that the patient is homozygous for the F508del mutation in the CFTR gene provided from an FDA-cleared CF mutation test, attestation baseline and follow-up ophthalmologic exams will be performed in pediatric patients starting on therapy.
Age Restrictions	2 years of age or older
Prescriber Restrictions	Restricted to pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# OSMOLEX (amantadine)

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## Products Affected

- OSMOLEX ER ORAL TABLET ER 24 HOUR THERAPY PACK
- OSMOLEX ER ORAL TABLET EXTENDED RELEASE 24 HOUR 129 MG, 193 MG, 258 MG

PA Criteria	Criteria Details
Exclusion Criteria	End-stage renal disease (creatinine clearance below 15 mL/min)
Required Medical Information	Diagnosis of covered use, documentation patient tried and failed immediate-release amantadine.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## OXBRYTA (voxelotor)

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### Products Affected

- OXBRYTA

PA Criteria	Criteria Details
Exclusion Criteria	Hemoglobin greater than 10.5 g/dL
Required Medical Information	Diagnosis of covered use, submission of hemoglobin level.
Age Restrictions	
Prescriber Restrictions	Restricted to hematology
Coverage Duration	6 months
Other Criteria	PA applies to all. Submission of improved hemoglobin level from baseline will be required for initial reauthorization after the first 6 months. Documentation of continued hemoglobin level improvement or maintenance of initial hemoglobin level improvement will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	



## OXERVATE (cenegermin-bkbj)

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### Products Affected

- OXERVATE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	Restricted to optometry and ophthalmology
Coverage Duration	8 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# OXYBATE SALT MEDICATIONS

## Products Affected

- XYREM
- XYWAV

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with sedative hypnotics
Required Medical Information	Diagnosis of covered use confirmed with documentation from a sleep study.
Age Restrictions	7 years of age or older
Prescriber Restrictions	Restricted to neurology, psychiatry, and sleep medicine
Coverage Duration	1 year
Other Criteria	PA applies to all. For adults with excessive daytime sleepiness associated with narcolepsy, Xyrem will be authorized only if the patient previously tried and had an inadequate clinical response or an intolerance to armodafinil or modafinil. Xywav will be authorized only if the patient has used Xyrem and prescriber submits a clinical reason detailing the need to switch to Xywav. Neither medication covered in this policy is indicated to treat insomnia and will not be approved for this use.
Indications	All Medically-accepted Indications.
Off Label Uses	

## PALYNZIQ (pegvaliase-pqpz)

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### Products Affected

- PALYNZIQ

PA Criteria	Criteria Details
Exclusion Criteria	Blood phenylalanine concentration below 600 micromol/L
Required Medical Information	Diagnosis of covered use, submission of blood phenylalanine concentration.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Reduction in blood phenylalanine concentration from pre-treatment baseline will be required for initial reauthorization after the first year. Documentation of continued phenylalanine level improvement or maintenance of initial phenylalanine level improvement will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PANRETIN (alitretinoin)

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## Products Affected

- PANRETIN

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy, requirement for systemic Kaposi's sarcoma therapy (more than 10 new Kaposi's Sarcoma lesions in the prior month, symptomatic lymphedema, symptomatic pulmonary Kaposi's sarcoma, or symptomatic visceral involvement)
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to dermatology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PARKINSON'S DISEASE "OFF" EPISODE (AS NEEDED) THERAPIES

## Products Affected

- *apomorphine hcl subcutaneous*
- INBRIJA
- KYNMOBI
- KYNMOBI TITRATION KIT

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	For Inbrija, administration of non-selective monoamine oxidase inhibitors within 14 days of initiation, asthma, COPD, or other chronic underlying lung disease. For Kynmobi, severe renal impairment or end-stage renal disease, severe hepatic impairment.
<b>Required Medical Information</b>	Diagnosis of covered use, attestation patient is experiencing "off" episodes despite carbidopa/levodopa therapy, prescription claims or documentation from physician showing patient is (a) currently taking at least one other medication to help reduce "off" episodes (from COMT inhibitor, dopamine agonist, or monoamine oxidase B inhibitor drug classes) or (b) has tried and failed, had an intolerance to, or has a contraindication to medications from at least two different drug classes (COMT inhibitors, dopamine agonists, monoamine oxidase B inhibitors) that can help reduce "off" episodes.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# PCSK9 INHIBITORS

## Products Affected

- PRALUENT SUBCUTANEOUS SOLUTION AUTO-INJECTOR
- REPATHA
- REPATHA PUSHTRONEX SYSTEM
- REPATHA SURECLICK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	For all indications, submission of LDL level obtained within the previous 6 months. For primary hyperlipidemia (including HeFH) and ASCVD indications, submission of current or previous lipid-lowering therapies. For HeFH, documentation of genetic test result documenting HeFH or diagnosis by clinical criteria using Simon Broom or WHO/Dutch Lipid Network criteria. For ASCVD, documented history of MI, ACS, stable or unstable angina, coronary or other arterial revascularization, stroke, TIA, or PAD.
Age Restrictions	For Repatha, 13 years of age or older. For Praluent, 18 years of age or older.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For approval for primary hyperlipidemia (including HeFH) and ASCVD indications, the patient must currently be using a statin plus ezetimibe or the patient must have tried and failed to have an adequate response to or had an intolerance to at least two statins or one statin and ezetimibe. At least one statin previously tried and failed must be a hydrophilic statin.
Indications	All Medically-accepted Indications.
Off Label Uses	

## PDE5 INHIBITORS (PAH)

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### Products Affected

- *alyq*
- *sildenafil citrate oral suspension reconstituted*
- *sildenafil citrate oral tablet 20 mg*
- *tadalafil (pah)*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	For tadalafil, diagnosis of severe (Child-Pugh class C) hepatic impairment, creatinine clearance below 30 mL/min or on hemodialysis
<b>Required Medical Information</b>	Diagnosis of covered use. For tadalafil, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to cardiology and pulmonology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# PEGFILGRASTIM

## Products Affected

- NEULASTA ONPRO
- NEULASTA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- UDENYCA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of FDA-approved indication.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All FDA-approved Indications.
Off Label Uses	



# PEMAZYRE (pemigatinib)

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## Products Affected

- PEMAZYRE ORAL TABLET 13.5 MG, 4.5 MG, 9 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use, submission of test result confirming presence of FGFR2 fusion or rearrangement, submission of previous systemic treatment(s) tried, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PIQRAY (alpelisib)

## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HR-positive, HER2-negative, PIK3CA-mutated, attestation that patient has advanced or metastatic disease and will be taking concurrently with fulvestrant, submission of prior therapies tried, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# POMALYST (pomalidomide)

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## Products Affected

- POMALYST

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use, submission of baseline serum bilirubin, AST, ALT, CBC including ANC and platelet count, prior therapies, when prior therapy was completed, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PRETOMANID

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## Products Affected

- PRETOMANID

PA Criteria	Criteria Details
Exclusion Criteria	Inability to use bedaquiline or linezolid
Required Medical Information	Diagnosis of covered use, attestation pretomanid will be used in combination with bedaquiline and linezolid.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to infectious diseases and pulmonology.
Coverage Duration	26 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PREVYMIS (letermovir)

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## Products Affected

- PREVYMIS ORAL

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, coadministration with ergot alkaloids, pimozone, pitavastatin, or simvastatin
Required Medical Information	Diagnosis of covered use, submission of day number post-HSCT, documentation of any previous doses of letermovir.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology, oncology, transplant specialist, and infectious diseases
Coverage Duration	100 days
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PRIOR AUTHORIZATION TO OVERRIDE SPECIALTY RESTRICTIONS

## Products Affected

- FABIOR
- PEG-INTRON REDIPEN SUBCUTANEOUS KIT 50 MCG/0.5ML
- PEG-INTRON SUBCUTANEOUS KIT 50 MCG/0.5ML
- PEGASYS PROCLICK SUBCUTANEOUS SOLUTION 135 MCG/0.5ML
- PEGASYS PROCLICK SUBCUTANEOUS SOLUTION AUTO-INJECTOR 135 MCG/0.5ML
- PEGASYS SUBCUTANEOUS SOLUTION
- PEGASYS SUBCUTANEOUS SOLUTION PREFILLED SYRINGE
- PEGINTRON SUBCUTANEOUS KIT 50 MCG/0.5ML
- *tazarotene external cream*
- *tazarotene external foam*
- TAZORAC EXTERNAL CREAM 0.05 %
- TAZORAC EXTERNAL GEL
- VABOMERE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	The following physician specialties are exempt from prior authorization (by drug): (a) for Fabior, tazarotene, and Tazorac: dermatology exempt, (b) for Pegasys: gastroenterology, hepatology, or infectious diseases exempt, (c) for Vabomere: infectious diseases or nephrology exempt
Coverage Duration	1 year
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PROCYSBI (cysteamine)

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## Products Affected

- PROCYSBI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation that patient has tried and failed or had an intolerance to immediate-release cysteamine.
Age Restrictions	1 year of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. For approval, the patient must have tried and failed to have an adequate response to, had an intolerance to, or have a contraindication to therapy with immediate-release cysteamine.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PROLIA (denosumab)

## Products Affected

- PROLIA SUBCUTANEOUS SOLUTION PREFILLED SYRINGE

PA Criteria	Criteria Details
Exclusion Criteria	Hypocalcemia, pregnancy
Required Medical Information	Diagnosis of covered use, submission of calcium level, pregnancy status for female patients of childbearing potential. "High risk for fracture" is defined as (1) a history of osteoporotic fracture or (2) multiple risk factors for fracture or (3) patients who have failed or are intolerant of other available osteoporosis therapies, confirmation of osteoporosis diagnosis either through densitometry (T-score less than or equal to -2.5 at the total hip, femoral neck, or lumbar spine) or clinically (documented presence of fragility fracture).
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated serum calcium level since the previous authorization will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	



# PROMACTA (eltrombopag)

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## Products Affected

- PROMACTA ORAL PACKET
- PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG, 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use (including cause of thrombocytopenia if being used for that indication), documentation of previous therapies tried (corticosteroids, immunoglobulins), submission of platelet count.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	6 months
Other Criteria	PA applies to all. Updated platelet count since the previous authorization will be required for subsequent reauthorizations. Not indicated for treatment of patients with myelodysplastic syndrome and will not be approved for this use.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PROMETHAZINE IN OLDER PATIENTS

## Products Affected

- *promethazine hcl oral*
- PROMETHAZINE HCL RECTAL SUPPOSITORY 12.5 MG
- *promethazine hcl rectal suppository 25 mg*
- *promethazine vc plain*
- *promethazine-phenylephrine*
- *promethegan rectal suppository 25 mg, 50 mg*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For allergic conditions, documentation must be submitted showing patient has tried and failed or had an inadequate response to a second-generation antihistamine.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	Promethazine is a potent anticholinergic considered high-risk in older patients due to risks of confusion, dry mouth, constipation, and decreased clearance with advanced age.
Indications	All Medically-accepted Indications.
Off Label Uses	

# PROSTATE CANCER ORAL MEDICATIONS

## Products Affected

- *abiraterone acetate oral tablet 250 mg*
- ERLEADA
- NUBEQA
- XTANDI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	For abiraterone, severe hepatic impairment (Child-Pugh class C)
<b>Required Medical Information</b>	Diagnosis of covered use. For Nubeqa, documentation of other treatments tried. For abiraterone, confirmation patient will receive concurrent prednisone, submission of baseline ALT, AST, bilirubin, and serum potassium level.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to oncology and urology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only. Nubeqa will be authorized only if the patient previously tried and had an inadequate clinical response or an intolerance to both Erleada and Xtandi.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# PYRUKYND (mitapivat)

## Products Affected

- PYRUKYND
- PYRUKYND TAPER PACK ORAL TABLET THERAPY PACK  
5 MG, 7 X 20 MG & 7 X 5 MG, 7 X 50 MG & 7 X 20 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with hematopoietic stimulating agents
<b>Required Medical Information</b>	Diagnosis of covered use, submission of test confirming presence of at least two mutant alleles in the PKLR gene, of which at least one is a missense mutation, and where the mutations are not a homozygous R479H mutation, hemoglobin level within the previous 3 months, past-year history of red blood cell (RBC) transfusions.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology or specialists in inborn errors of metabolism
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. Positive history of RBC transfusions is required for initial coverage. For initial reauthorization, improvement of hemoglobin level and/or reductions in annualized rate of RBC transfusions is required. Continued improvement in either hemoglobin level or reductions in RBC transfusional burden from baseline will be required for subsequent reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## QINLOCK (ripretinib)

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### Products Affected

- QINLOCK

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of previous kinase inhibitor therapies, baseline blood pressure reading, attestation patient does not have uncontrolled hypertension, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## QUVIVIQ (daridorexant)

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### Products Affected

- QUVIVIQ

PA Criteria	Criteria Details
Exclusion Criteria	Narcolepsy
Required Medical Information	Diagnosis of covered use. Patient must have tried and failed to tolerate or had an inadequate response to two covered alternative therapies recommended by the American Academy of Sleep Medicine (doxepin, eszopiclone, ramelteon, suvorexant, temazepam, zaleplon, zolpidem) including one non-suvorexant therapy for sleep maintenance (doxepin, eszopiclone, temazepam) if that is the diagnosis of covered use.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# RADICAVA ORS (edaravone)

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## Products Affected

- RADICAVA ORS
- RADICAVA ORS STARTER KIT

PA Criteria	Criteria Details
Exclusion Criteria	ALS duration of greater than 2 years
Required Medical Information	Diagnosis of covered use, submission of ALS Functional Rating Scale - Revised (ALSFRS-R) scoring (patient is required to have scores of 2 points or better on each of the 12 individual ALSFRS-R items).
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# RAVICTI (glycerol phenylbutyrate)

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## Products Affected

- RAVICTI

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline fasting plasma ammonia level.
Age Restrictions	2 months of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# RECORLEV (levoketoconazole)

## Products Affected

- RECORLEV

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Cirrhosis, acute, poorly-controlled chronic, or extensive metastatic liver disease, baseline AST or ALT greater than 3 times the upper limit of normal, recurrent symptomatic cholelithiasis, a prior history of drug-induced liver injury due to ketoconazole or any azole antifungal therapy that required discontinuation of treatment, prolonged QTcF interval greater than 470 msec at baseline, history of torsades de pointes, ventricular tachycardia, ventricular fibrillation, or prolonged QT syndrome, coadministration with drugs that cause QT prolongation associated with ventricular arrhythmias
<b>Required Medical Information</b>	Diagnosis of covered use, attestation pituitary gland surgery is not an option for the patient or has not been curative, submission of cortisol level (e.g., 24-hour urine free cortisol test), electrocardiogram (including QTcF), and liver function tests all performed within 3 months of prior authorization request, documentation patient tried and failed at least one other therapy for Cushing's syndrome (e.g., mifepristone, osilodrostat, pasireotide).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to endocrinology
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. Continuation requires documentation of clinically relevant response to therapy, including, but not limited to urine free cortisol level. Recorlev is not approved for the treatment of fungal infections and will not be approved for this use.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# RETACRIT (epoetin alfa-epbx)

## Products Affected

- RETACRIT INJECTION SOLUTION 10000 UNIT/ML, 2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML, 4000 UNIT/ML, 40000 UNIT/ML

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of hemoglobin or hematocrit level, serum iron, total iron-binding capacity (TIBC), and transferrin within 30 days of request date, documentation that the patient does not have uncontrolled hypertension.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	For non-ESRD-related conditions: 90 days. For ESRD-related conditions: 1 year.
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# RETEVMO (selpercatinib)

## Products Affected

- RETEVMO ORAL CAPSULE 40 MG, 80 MG

PA Criteria	Criteria Details
Exclusion Criteria	Uncontrolled hypertension, coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of RET gene fusion or mutation, baseline blood pressure reading, pregnancy status for female patients of childbearing potential. For patients with RET fusion-positive thyroid cancer, submission of date or year of previous previous radioactive iodine treatment or reason why radioactive iodine therapy is not appropriate.
Age Restrictions	12 years of age or older based on indication
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# REVCOVI (elapegademase-lvlr)

## Products Affected

- REVCOVI

PA Criteria	Criteria Details
Exclusion Criteria	Severe thrombocytopenia
Required Medical Information	Diagnosis of covered use, submission of plasma ADA activity and platelet count.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated plasma ADA level and platelet count since the previous authorization will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All FDA-approved Indications.
Off Label Uses	

# REVLIMID (lenalidomide)

## Products Affected

- *lenalidomide*
- REVLIMID

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Pregnancy, chronic lymphocytic leukemia (outside of a controlled clinical trial)
<b>Required Medical Information</b>	Diagnosis of covered use, submission of CBC including ANC and platelet count, pregnancy status for female patients of childbearing potential. For mantle cell lymphoma, documentation of at least two prior therapies tried, one of which included bortezomib (or a documented contraindication to bortezomib). For follicular lymphoma and marginal zone lymphoma, submission of prior treatments tried and attestation medication will be coadministered with a rituximab product.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology and oncology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# REZUROCK (belumosudil)

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## Products Affected

- REZUROCK

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of at least 2 previous therapies tried and failed for chronic graft-versus-host disease, pregnancy status for female patients of childbearing potential.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to hematology, oncology, and transplant specialty
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All FDA-approved Indications.
Off Label Uses	

# ROZLYTREK (entrectinib)

## Products Affected

- ROZLYTREK ORAL CAPSULE 100 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential. For non-small cell lung cancer, submission of results showing tumor is ROS1-positive as detected by an FDA-approved test. For solid tumors, submission of evidence of a neurotrophic receptor tyrosine kinase (NTRK) gene fusion without a known acquired resistance mutation.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# RUBRACA (rucaparib)

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## Products Affected

- RUBRACA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline CBC. For BRCA mutation-associated ovarian, fallopian tube, primary peritoneal or metastatic castration-resistant prostate cancer, confirmation of deleterious BRCA mutation as detected by FDA-approved companion diagnostic test, documentation that the patient has been treated with two or more chemotherapies.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology and urology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For initial approval, patient must have recovered from hematological toxicity caused by previous chemotherapy (Grade 1 or less).
Indications	All Medically-accepted Indications.
Off Label Uses	



# RYDAPT (midostaurin)

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## Products Affected

- RYDAPT

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use. For acute myeloid leukemia, submission of FDA-approved test confirming presence of FLT3 mutation, documentation of other chemotherapy that will be coadministered with midostaurin.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SAPROPTERIN

## Products Affected

- *sapropterin dihydrochloride oral packet*
- *sapropterin dihydrochloride oral tablet*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of blood phenylalanine level.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Submission of blood phenylalanine concentration documenting a reduction in blood phenylalanine concentration from pre-treatment baseline will be required for initial reauthorization. Documentation of continued phenylalanine level improvement or maintenance of initial phenylalanine level improvement will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SCEMBLIX (asciminib)

### Products Affected

- SCEMBLIX ORAL TABLET 20 MG, 40 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of pregnancy status for female patients of childbearing potential. For use in patients with a T315I mutation, documentation patient has first tried and failed or become intolerant to ponatinib. For use in patients without a T315I mutation, documentation of other tyrosine kinase inhibitors tried and failed.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. For approval in T315I-mutation-positive CML, the patient must have tried and failed to have an adequate response to or had an intolerance to ponatinib.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SEDATIVE HYPNOTICS IN OLDER PATIENTS

## Products Affected

- *eszopiclone*
- *zaleplon*
- *zolpidem tartrate oral*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation at least two of the following medications were tried and deemed ineffective or intolerable: Belsomra, doxepin tablets, ramelteon, and trazodone.
Age Restrictions	PA applies to patients 65 years of age or older. PA does not apply to patients 64 years of age or younger.
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Sedative hypnotic medications are high-risk medications in older patients due to increased risks of cognitive impairment, delirium, unsteady gait, syncope, falls, fractures, and motor vehicle accidents.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SEROSTIM (somatropin)

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## Products Affected

- SEROSTIM SUBCUTANEOUS SOLUTION  
RECONSTITUTED 4 MG, 5 MG, 6 MG

PA Criteria	Criteria Details
Exclusion Criteria	Active malignancy, acute critical illness, active proliferative or severe non-proliferative diabetic retinopathy
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. Serostim is indicated only for the treatment of HIV-associated cachexia/wasting and uses outside of this indication will not be approved.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SIGNIFOR (pasireotide)

## Products Affected

- SIGNIFOR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe hepatic impairment (Child-Pugh class C), uncorrected hypokalemia or hypomagnesemia
<b>Required Medical Information</b>	Diagnosis of covered use, submission of ALT, aspartate aminotransferase, alkaline phosphatase, total bilirubin, and serum potassium and magnesium levels, attestation that pituitary radiation/surgery was not curative or is not an option.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to endocrinology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Updated liver function testing and serum potassium and magnesium levels since the previous authorization will be required for subsequent reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# SIMVASTATIN 80 mg per day

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## Products Affected

- *ezetimibe-simvastatin oral tablet 10-80 mg*
- *simvastatin oral tablet 80 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Any new start to therapy. Not recommended as initial therapy nor for patients already taking lower doses of simvastatin whose response is inadequate.
Required Medical Information	Diagnosis of covered use, documentation that patient has been taking simvastatin 80 mg daily for 12 months or longer without ill effect, submission of lipid panel, liver function tests, and serum creatinine level all obtained within the past 12 months.
Age Restrictions	10 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SIRTURO (bedaquiline)

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## Products Affected

- SIRTURO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline ECG, serum potassium, calcium, magnesium, ALT, AST, alkaline phosphatase, and bilirubin, confirmation that Sirturo will be co-administered with pretomanid and linezolid or at least 3 other drugs proven to be or at least 4 other drugs suspected to be effective against the patient's M. tuberculosis isolate and submission of susceptibility testing, if available.
Age Restrictions	5 years of age or older
Prescriber Restrictions	Restricted to infectious diseases and pulmonology
Coverage Duration	26 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# SIVEXTRO (tedizolid)

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## Products Affected

- SIVEXTRO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation of a culture and sensitivity showing that the suspected causative agent is susceptible to this medication.
Age Restrictions	12 years of age or older
Prescriber Restrictions	Restricted to infectious diseases
Coverage Duration	6 days
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SOFOSBUVIR/VELPATASVIR

## Products Affected

- EPCLUSA ORAL PACKET
- EPCLUSA ORAL TABLET 200-50 MG, 400-100 MG
- *sofosbuvir-velpatasvir*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, laboratory confirmation of hepatitis C virus (HCV) infection, submission of baseline HCV RNA level, documentation of whether cirrhosis is present or not and whether it is compensated or decompensated, confirmation that patients with decompensated cirrhosis will receive concomitant ribavirin therapy unless ribavirin therapy is otherwise clinically not indicated, submission of eGFR (safety and efficacy of sofosbuvir/velpatasvir has not been established in patients with eGFR less than 30 mL/min/1.73 m <sup>2</sup> ), confirmation a test for HBV infection (HBsAg and anti-HBc) was completed.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 weeks
Other Criteria	PA applies to all. For approval of brand Epclusa 400 mg/100 mg, the patient must have tried and failed to have an adequate response to or had an intolerance to sofosbuvir/velpatasvir 400 mg/100 mg.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SOMATULINE DEPOT (lanreotide)

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## Products Affected

- SOMATULINE DEPOT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation of baseline serum GH, IGF-1, TSH, and blood glucose levels. For acromegaly, attestation that pituitary radiation/surgery was not curative or is not an option.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology and oncology
Coverage Duration	Initially 3 months, then up to 1 year
Other Criteria	PA applies to new starts only. Continuation of therapy requires documentation of a positive clinical response. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SOMAVERT (pegvisomant)

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## Products Affected

- SOMAVERT

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline IGF-1, ALT, AST, alkaline phosphatase, and serum total bilirubin, attestation that surgery or radiation was not curative or is not an option.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to endocrinology
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated IGF-1 level and liver function tests since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SOVALDI (sofosbuvir)

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## Products Affected

- SOVALDI ORAL PACKET
- SOVALDI ORAL TABLET 200 MG, 400 MG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, laboratory confirmation of hepatitis C virus (HCV) genotype 1a, 1b, 2, 3, or 4 infection, documentation of whether cirrhosis is present or not and whether or not it is compensated or decompensated, confirmation a test for HBV infection (HBsAg and anti-HBc) was completed.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	Criteria for coverage duration will be applied consistent with current AASLD-IDSA guidance.
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SPRYCEL (dasatinib)

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### Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Uncorrected hypokalemia, uncorrected hypomagnesemia
Required Medical Information	Diagnosis of covered use, submission of serum potassium and magnesium, pregnancy status for female patients of childbearing potential. For adults with resistance or intolerance to prior therapy, documentation of prior therapy.
Age Restrictions	
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## STIVARGA (regorafenib)

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### Products Affected

- STIVARGA

PA Criteria	Criteria Details
Exclusion Criteria	Severe or uncontrolled hypertension, coadministration with strong CYP3A4 inhibitors or inducers
Required Medical Information	Diagnosis of covered use, submission of previous therapies, submission of baseline ALT, AST, serum bilirubin, blood pressure reading, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SUCRAID (sacrosidase)

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## Products Affected

- SUCRAID

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of laboratory-confirmed congenital sucrase-isomaltase deficiency via differential urinary disaccharide test or measurement of intestinal disaccharides following small bowel biopsy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	



# SUNOSI (solriamfetol)

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## Products Affected

- SUNOSI ORAL TABLET 150 MG, 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	End-stage renal disease, administration of non-selective monoamine oxidase inhibitors within 14 days of initiation, serious arrhythmias, unstable cardiovascular disease including uncontrolled hypertension
Required Medical Information	Diagnosis of covered use, submission of baseline blood pressure reading.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology and sleep medicine
Coverage Duration	1 year
Other Criteria	PA applies to all. Solriamfetol is not indicated to treat the underlying airway obstruction in obstructive sleep apnea and will not be approved for this use.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SUTENT (sunitinib)

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### Products Affected

- *sunitinib malate*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SYMDEKO (tezacaftor/ivacaftor)

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### Products Affected

- SYMDEKO

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with strong CYP3A inducers
<b>Required Medical Information</b>	Diagnosis of covered use. Documentation that the patient is homozygous for the F508del mutation or has at least one mutation in the CTFR gene responsive to the drug (see section 12.1, table 4 of package insert for full list) provided from an FDA-cleared CF mutation test. Submission of documentation that baseline and follow-up ophthalmologic exams will be performed in pediatric patients starting on therapy.
<b>Age Restrictions</b>	6 years of age or older
<b>Prescriber Restrictions</b>	Restricted to pulmonology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to new starts only.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## SYMLIN (pramlintide)

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### Products Affected

- SYMLINPEN 120 SUBCUTANEOUS SOLUTION PEN-INJECTOR
- SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR

PA Criteria	Criteria Details
Exclusion Criteria	Gastroparesis
Required Medical Information	Diagnosis of covered use, confirmation of current use of a mealtime insulin.
Age Restrictions	
Prescriber Restrictions	PA not required for endocrinology
Coverage Duration	1 year
Other Criteria	PA applies to all except when prescribed by endocrinology.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SYMPROIC (naldemedine)

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### Products Affected

- SYMPROIC

PA Criteria	Criteria Details
Exclusion Criteria	Known or suspected gastrointestinal obstruction or increased risk of recurrent obstruction, severe hepatic impairment (Child-Pugh class C)
Required Medical Information	Diagnosis of covered use, documentation patient has been using opioids at a morphine equivalent dose of at least 30 mg daily for at least 4 weeks prior to initiation, provider must attest that if opioid medication is stopped for any reason, naldemedine will be discontinued.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

## SYNAREL (nafarelin)

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### Products Affected

- SYNAREL

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy/breast-feeding, undiagnosed abnormal vaginal bleeding
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	For endometriosis, 6 months. For all other diagnoses, 1 year.
Other Criteria	PA applies to all. Re-treatment for endometriosis is not recommended because safety data are not available.
Indications	All Medically-accepted Indications.
Off Label Uses	

# SYNRIBO (omacetaxine)

## Products Affected

- SYNRIBO

PA Criteria	Criteria Details
Exclusion Criteria	Poor glycemic control
Required Medical Information	Diagnosis of covered use, submission of prior therapies tried and failed, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TABRECTA (capmatinib)

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## Products Affected

- TABRECTA ORAL TABLET 150 MG, 200 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of MET exon 14 skipping mutation, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# TAFAMIDIS

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## Products Affected

- VYNDAMAX
- VYNDAQEL

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TAFINLAR (dabrafenib)

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## Products Affected

- TAFINLAR

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP2C8 or CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of BRAF V600E or V600K mutation. For non-small cell lung cancer, thyroid cancer, or unresectable/metastatic melanoma with a BRAF V600K mutation, attestation that therapy will be used in combination with trametinib.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TAGRISSO (osimertinib)

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## Products Affected

- TAGRISSO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, confirmation of the presence of required genetic mutations/deletions as detected by an FDA-approved test, pregnancy status for female patients of childbearing potential. For EGFR T790M mutation-positive NSCLC, documentation that the patient has progressed on or after EGFR TKI therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	Initially 3 months, then 1 year
Other Criteria	PA applies to new starts only. Continuation of approval requires affirmation of absence of unacceptable toxicities.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TALZENNA (talazoparib)

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## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of FDA-approved companion test results showing patient is a candidate for therapy and pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TARGRETIN (bexarotene) GEL

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## Products Affected

- *bexarotene external*

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy
Required Medical Information	Diagnosis of covered use, submission of previous therapies.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to dermatology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All FDA-approved Indications.
Off Label Uses	

# TARPEYO (budesonide)

## Products Affected

- TARPEYO

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Severe (Child-Pugh class C) hepatic impairment, eGFR less than 35 mL/min/1.73 m <sup>2</sup>
<b>Required Medical Information</b>	Diagnosis of covered use, primary IgA nephropathy confirmed by biopsy, submission of 24-hour urine protein of at least 1 gram/day or 24-hour urine protein-to-creatinine ratio of at least 1.5 g/g, submission of eGFR and liver function testing or Child-Pugh class, attestation patient is stable on a maximally-tolerated renin-angiotensin system antagonist (ACE inhibitor or ARB), documentation patient has progressed on at least one immunosuppressant (e.g., azathioprine, mycophenolate, etc.).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to immunology and nephrology
<b>Coverage Duration</b>	41 weeks
<b>Other Criteria</b>	PA applies to all. Approval for additional 41-week courses requires documentation of clinically relevant response to therapy, including, but not limited to stabilization or improvement of urine protein-to-creatinine ratio or eGFR.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# TASIGNA (nilotinib)

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## Products Affected

- TASIGNA

PA Criteria	Criteria Details
Exclusion Criteria	Uncorrected hypokalemia, uncorrected hypomagnesemia, long QT syndrome, coadministration with drugs that prolong the QT interval or strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of baseline ECG, Philadelphia chromosome (Ph) status, potassium and magnesium levels.
Age Restrictions	
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TAVALISSE (fostamatinib)

## Products Affected

- TAVALISSE ORAL TABLET 100 MG, 150 MG

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with strong CYP3A inducers
<b>Required Medical Information</b>	Diagnosis of covered use, submission of platelet count, documentation patient had an insufficient response to prior treatment (including at least one of the following: corticosteroids, immunoglobulins, splenectomy, and/or a thrombopoietin receptor agonist).
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to hematology
<b>Coverage Duration</b>	Initially 12 weeks, then 1 year
<b>Other Criteria</b>	PA applies to all. Documentation of an improvement in platelet count will be required for initial reauthorization after the first 12 weeks. Maintenance of a clinical benefit will be required for subsequent annual reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



# TAVNEOS (avacopan)

## Products Affected

- TAVNEOS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Coadministration with moderate or strong CYP3A4 inducers, active serious infection, chronic active hepatitis B, untreated hepatitis C, uncontrolled autoimmune hepatitis, cirrhosis
<b>Required Medical Information</b>	Diagnosis of covered use (GPA or MPA variant of ANCA-associated vasculitis) and confirmation patient is using rituximab, cyclophosphamide/azathioprine, or another compendium-supported therapy for the treatment of ANCA-associated vasculitis, along with glucocorticoids.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to immunology, nephrology, pulmonology, and rheumatology
<b>Coverage Duration</b>	Initially 6 months, then 1 year
<b>Other Criteria</b>	PA applies to all. Reauthorization requires documentation of clinically relevant response to therapy, including but not limited to disease remission defined using changes in Birmingham Vasculitis Activity Score, a documented reduction in maintenance glucocorticoid dose, or improved or sustained renal function.
<b>Indications</b>	All FDA-approved Indications.
<b>Off Label Uses</b>	

# TAZVERIK (tazemetostat)

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## Products Affected

- TAZVERIK

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inhibitors or moderate or strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, pregnancy status for female patients of childbearing potential.
Age Restrictions	16 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	Initially 16 weeks, then 1 year
Other Criteria	PA applies to new starts only. Continuation of therapy requires (a) documentation of a positive clinical response and (b) attestation no known secondary malignancies have developed.
Indications	All Medically-accepted Indications.
Off Label Uses	

## TEGSEDI (inotersen)

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### Products Affected

- TEGSEDI

PA Criteria	Criteria Details
Exclusion Criteria	Platelet count less than $100 \times 10^9$ L
Required Medical Information	Diagnosis of covered use, submission of platelet count.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated platelet count since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TEPMETKO (tepotinib)

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## Products Affected

- TEPMETKO

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers or dual strong CYP3A4/P-glycoprotein inhibitors
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of MET exon 14 skipping mutation, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TERIPARATIDE

## Products Affected

- FORTEO SUBCUTANEOUS SOLUTION 600 MCG/2.4ML
- FORTEO SUBCUTANEOUS SOLUTION PEN-INJECTOR
- TERIPARATIDE (RECOMBINANT)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, postmenopausal status, submission of serum calcium level, documentation that other treatment options have failed (or are contraindicated), submission of a value, condition, or past medical history that assesses fracture risk (e.g., DEXA scan results or prior fracture), submission of number of total months of all prior use of parathyroid hormone analogs and parathyroid hormone related peptides.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated serum calcium since the previous authorization will be required for reauthorization.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TESTOSTERONE REPLACEMENT PRODUCTS

## Products Affected

- ANDRODERM TRANSDERMAL PATCH 24 HOUR (1%)
- TESTOSTERONE TRANSDERMAL GEL 10 MG/ACT (2%) • *testosterone transdermal solution*
- *testosterone transdermal gel 12.5 mg/act (1%), 20.25 mg/1.25gm (1.62%), 20.25 mg/act (1.62%), 25 mg/2.5gm (1%), 40.5 mg/2.5gm (1.62%), 50 mg/5gm*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of serum testosterone level, documentation that patient has been evaluated for the presence of breast and prostate cancer prior to initiation of therapy.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Documentation of clinically relevant response to therapy (including, but not limited to submission of updated serum testosterone level) will be required for subsequent reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TIBSOVO (ivosidenib)

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## Products Affected

- TIBSOVO

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of IDH1 mutation.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TOBI PODHALER (tobramycin)

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## Products Affected

- TOBI PODHALER

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	6 years of age or older
Prescriber Restrictions	
Coverage Duration	1 month
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# TOLVAPTAN (HYPONATREMIA)

## Products Affected

- SAMSCA ORAL TABLET 15 MG
- *tolvaptan*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	History of signs or symptoms of significant liver impairment or injury, need to raise serum sodium acutely, inability to sense or respond to thirst, hypovolemia, anuria, coadministration with strong CYP3A inhibitors or inducers or desmopressin
<b>Required Medical Information</b>	Diagnosis of covered use, submission of serum sodium.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	30 days
<b>Other Criteria</b>	PA applies to all. Treatment should be initiated in a setting where serum sodium can be monitored closely. Treatment is limited to 30 days to prevent liver injury. This formulation of tolvaptan will not be approved for autosomal dominant polycystic kidney disease (ADPKD) because the tolvaptan formulation approved for ADPKD has a mandatory REMS program.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# TOPICAL ONYCHOMYCOSIS TREATMENTS

## Products Affected

- JUBLIA
- *tavaborole*

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of culture-proven <i>Trichophyton rubrum</i> or <i>Trichophyton mentagrophytes</i> infection, documentation patient has (1) tried and failed to respond to or tolerate oral terbinafine therapy or a documented contraindication to its use exists, and (2) tried and failed therapy with topical ciclopirox nail solution.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	48 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TOPICAL PSORIASIS TREATMENTS

## Products Affected

- VTAMA
- ZORYVE

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of percent body surface area affected (with a requirement BSA affected is less than or equal to 20 percent), documentation patient either (1) has tried and failed, had an incomplete response to, had an intolerance to, or has contraindications to at least one Class/Group 3 high potency or stronger topical corticosteroid and at least one of the following other topical agents: (a) a vitamin D analog such as calcipotriene or calcitriol, (b) tazarotene, or (c) a topical calcineurin inhibitor, or (2) patient is currently using a systemic medication (biologic or otherwise) to manage psoriasis.
Age Restrictions	For Vtama, 18 years of age or older. For Zoryve, 12 years of age or older.
Prescriber Restrictions	For Vtama, restricted to dermatology. For Zoryve, PA not required for dermatology.
Coverage Duration	1 year
Other Criteria	PA applies to all. Documentation of a positive response to therapy will be required for reauthorization.
Indications	All FDA-approved Indications.
Off Label Uses	

# TRIKAFTA (elexacaftor/tezacaftor/ivacaftor)

## Products Affected

- TRIKAFTA ORAL TABLET THERAPY PACK 100-50-75 & 150 MG, 50-25-37.5 & 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, documentation that the patient has at least one mutation in the CFTR gene responsive to the drug (see section 12.1, table 4 of package insert for full list) or a mutation that is responsive based on in vitro data provided from an FDA-cleared CF mutation test, submission of documentation that baseline and follow-up ophthalmologic exams will be performed in pediatric patients starting on therapy.
Age Restrictions	
Prescriber Restrictions	Restricted to pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TRUSELTIQ (infigratinib)

## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers or inhibitors, coadministration with proton pump inhibitors
Required Medical Information	Diagnosis of covered use, submission of test result confirming presence of FGFR2 fusion or rearrangement, submission of previous systemic treatment(s) tried.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TUKYSA (tucatinib)

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## Products Affected

- TUKYSA ORAL TABLET 150 MG, 50 MG

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy, coadministration with strong CYP3A inducers, strong CYP2C8 inhibitors, or moderate CYP2C8 inducers
Required Medical Information	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HER2-positive, submission of previous systemic treatment including prior HER2-directed therapy, pregnancy status for female patients of childbearing potential, confirmation that the treatment regimen will include concomitant use of capecitabine and trastuzumab.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TURALIO (pexidartinib)

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## Products Affected

- TURALIO

PA Criteria	Criteria Details
Exclusion Criteria	Active liver or biliary tract disease (including increased ALP), pre-existing increased serum transaminases, total or direct bilirubin greater than the upper limit of normal, coadministration with other hepatotoxic medications, strong CYP3A inducers, or proton pump inhibitors
Required Medical Information	Diagnosis of covered use (and surgical intervention is not possible or practical), submission of serum transaminases, total and direct bilirubin, and ALP.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# TYMLOS (abaloparatide)

## Products Affected

- TYMLOS

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Male patients, female patients of childbearing potential, pre-existing hypercalcemia, underlying hypercalcemic disorder (such as primary hyperparathyroidism), patients with an increased risk of osteosarcoma (such as those with Paget's disease)
<b>Required Medical Information</b>	Diagnosis of covered use where "high risk for fracture" is defined as (1) a history of osteoporotic fracture or (2) multiple risk factors for fracture or (3) patients who have failed or are intolerant of other available osteoporosis therapies, submission of baseline serum calcium, postmenopausal status, documentation that at least one bisphosphonate was tried and failed (or a bisphosphonate is contraindicated), submission of a value, condition, or past medical history that assesses fracture risk (e.g., DEXA scan results or prior fracture), submission of number of total months of all prior use of parathyroid hormone analogs and parathyroid hormone related peptides.
<b>Age Restrictions</b>	
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	2 years maximum dependent on patient's prior use of all PTH analogs and PTH-related peptides
<b>Other Criteria</b>	PA applies to all. Use of parathyroid hormone analogs and/or parathyroid hormone related peptides for more than 2 years during a patient's lifetime is not recommended and requests for therapy with any of these agents for a combined total of greater than 2 years will not be approved.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	



## UPNEEQ (oxymetazoline)

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### Products Affected

- UPNEEQ

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to ophthalmologic surgery
Coverage Duration	Initially 90 days, then 1 year
Other Criteria	PA applies to all. Submission of clinically significant response to therapy will be required for reauthorization. Not FDA-approved for cosmetic use and therefore uses outside of acquired blepharoptosis will not be approved.
Indications	All FDA-approved Indications.
Off Label Uses	

## UPTRAVI (selexipag)

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### Products Affected

- UPTRAVI ORAL TABLET 1000 MCG, 1200 MCG, 1400 MCG, 1600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG
- UPTRAVI ORAL TABLET THERAPY PACK

PA Criteria	Criteria Details
Exclusion Criteria	Severe (Child-Pugh class C) hepatic impairment
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology and pulmonology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VALCHLOR (mechlorethamine)

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## Products Affected

- VALCHLOR

PA Criteria	Criteria Details
Exclusion Criteria	Use as initial therapy
Required Medical Information	Diagnosis of covered use, submission of previous skin-directed therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to dermatology and oncology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to new starts only. Submission of clinically significant response to therapy will be required for reauthorization.
Indications	All Medically-accepted Indications.
Off Label Uses	

## VECAMEYL (mecamylamine)

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### Products Affected

- VECAMEYL

PA Criteria	Criteria Details
Exclusion Criteria	Mild, moderate, labile hypertension, coronary insufficiency or history of recent myocardial infarction, uremia, glaucoma, organic pyloric stenosis
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology and neurology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VEMLIDY (tenofovir alafenamide)

## Products Affected

- VEMILIDY

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	End-stage renal disease patients not receiving chronic hemodialysis, decompensated (Child-Pugh class B or C) hepatic impairment
<b>Required Medical Information</b>	Diagnosis of covered use, submission of creatinine clearance (or serum creatinine, actual body weight, and height to calculate estimated creatinine clearance), confirmation of HIV test and that drug will not be used by itself in the case of HIV co-infection.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	PA not required for gastroenterology or infectious diseases
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all except gastroenterology or infectious diseases. Updated creatinine clearance since the previous authorization will be required for subsequent annual reauthorizations.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# VENCLEXTA (venetoclax)

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## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A inducers. For CLL/SLL, coadministration with strong CYP3A inhibitors at treatment initiation and initial dosage titration.
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VENTAVIS (iloprost)

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## Products Affected

- VENTAVIS

PA Criteria	Criteria Details
Exclusion Criteria	Systolic blood pressure below 85 mmHg
Required Medical Information	Diagnosis of covered use, submission of baseline systolic blood pressure.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to cardiology and pulmonology
Coverage Duration	1 year
Other Criteria	This medication is covered as a Part B benefit except for enrollees residing in a long-term care facility. PA applies to new starts only when covered as a Part D benefit.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VERQUVO (vericiguat)

## Products Affected

- VERQUVO ORAL TABLET 10 MG, 2.5 MG, 5 MG

PA Criteria	Criteria Details
Exclusion Criteria	Concomitant use of another soluble guanylate cyclase (sGC) stimulator or a phosphodiesterase-5 (PDE-5) inhibitor
Required Medical Information	Diagnosis, including either hospitalization for heart failure with reduced ejection fraction (HFrEF) within the previous 6 months or outpatient IV diuretic use within the previous 3 months, submission of left ventricular ejection fraction and pregnancy status for female patients of childbearing potential. Prescribers are also required to submit current regimen for the treatment of HFrEF, which must include (1) a renin-angiotensin system (RAS) inhibitor (ACE inhibitor, ARB, or sacubitril/valsartan), (2) a beta-blocker (BB), and (3) a mineralocorticoid receptor antagonist (MRA), each at maximally-tolerated doses. If any of these three therapies are not currently being used, prescriber is required to submit documentation as to why (e.g., contraindications, intolerances, etc.). Using the recommended dose of each therapeutic component to treat HFrEF is required. If the doses of any of these three components have not been optimized to the recommended dose to treat HFrEF, the prescriber is required to submit documentation as to why (e.g., intolerances, physiologic parameters, etc.). If the patient is using a BB not indicated for HFrEF, the patient will be required to switch to one of the three FDA-approved BBs for HFrEF (bisoprolol, carvedilol, or metoprolol succinate).
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All FDA-approved Indications.
Off Label Uses	



# VERZENIO (abemaciclib)

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## Products Affected

- VERZENIO

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with moderate or strong CYP3A4 inducers or ketoconazole
Required Medical Information	Diagnosis of covered use, submission of genetic tumor testing showing that the primary tumor type is HR-positive, HER2-negative, submission of baseline liver function tests and CBC, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VIBERZI (eluxadoline)

## Products Affected

- VIBERZI

PA Criteria	Criteria Details
Exclusion Criteria	Prior cholecystectomy, known or suspected biliary duct obstruction, known or suspected sphincter of Oddi disease or dysfunction, alcoholism, alcohol abuse, alcohol addiction, or patients who drink more than 3 alcoholic beverages/day, history of pancreatitis, structural diseases of pancreas including known or suspected pancreatic duct obstruction, severe hepatic impairment (Child-Pugh class C), severe constipation or sequelae from constipation, known or suspected mechanical gastrointestinal obstruction
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to gastroenterology
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VIJOICE (alpelisib)

## Products Affected

- *vijoice oral tablet therapy pack 125 mg, 200 & 50 mg, 50 mg*

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A4 inducers
Required Medical Information	Diagnosis of covered use including at least one target lesion on imaging with requesting provider attestation patient has severe or life-threatening disease, submission of test confirming presence of mutation in PIK3CA gene, confirmation of negative pregnancy status for female patients of childbearing potential or attestation from physician patient is not pregnant and will be using a highly effective method of contraception.
Age Restrictions	
Prescriber Restrictions	Restricted to specialists in genetic diseases or inborn errors of metabolism
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to all. Submission of objective documentation of a clinical benefit (e.g., reductions in target lesion size, pain, vascular malformations, limb enlargements, etc.) in the absence of unacceptable toxicity will be required for subsequent reauthorizations.
Indications	All FDA-approved Indications.
Off Label Uses	

# VITRAKVI (larotrectinib)

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## Products Affected

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

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of evidence the solid tumor has a neurotrophic receptor tyrosine kinase (NTRK) gene fusion without a known acquired resistance mutation, pregnancy status for female patients of childbearing potential.
Age Restrictions	
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VIVJOA (oteseconazole)

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## Products Affected

- VIVJOA

PA Criteria	Criteria Details
Exclusion Criteria	Women of reproductive potential
Required Medical Information	Diagnosis of covered use, including attestation patient has had at least three episodes of vulvovaginal candidiasis in the previous 12 months, attestation patient is either (a) postmenopausal or (b) infertile.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	12 weeks
Other Criteria	PA applies to all.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VIZIMPRO (dacomitinib)

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## Products Affected

- VIZIMPRO ORAL TABLET 15 MG, 30 MG, 45 MG

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with a proton pump inhibitor
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of EGFR exon 19 deletion or exon 21 L858R substitution mutation, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VMAT2 INHIBITORS

## Products Affected

- AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG
- INGREZZA ORAL CAPSULE 40 MG, 60 MG, 80 MG
- INGREZZA ORAL CAPSULE THERAPY PACK
- *tetrabenazine*

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Congenital long QT syndrome or a history of cardiac arrhythmia associated with a prolonged QT interval, coadministration with monoamine oxidase inhibitors. For tetrabenazine and Austedo, actively suicidal or untreated/undertreated depression, hepatic impairment. For Ingrezza, coadministration with strong CYP3A4 inducers.
<b>Required Medical Information</b>	Diagnosis of covered use.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	Restricted to neurology and psychiatry
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

## VONJO (pacritinib)

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### Products Affected

- VONJO

PA Criteria	Criteria Details
Exclusion Criteria	Moderate or severe (Child-Pugh class B or C) hepatic impairment, estimated glomerular filtration rate (eGFR) less than 30 mL/min, QTc interval greater than 480 msec at baseline, coadministration with strong CYP3A4 inducers or strong CYP3A4 inhibitors
Required Medical Information	Diagnosis of covered use, submission of platelet count, eGFR, and QTc interval, documentation from a physical exam patient has splenomegaly.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology or oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



# VOSEVI (sofosbuvir, velpatasvir, voxilaprevir)

## Products Affected

- VOSEVI

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Moderate or severe hepatic impairment, coadministration with rifampin or drugs that are strong P-glycoprotein inducers or moderate to strong CYP2B6, CYP2C8, or CYP3A4 inducers
<b>Required Medical Information</b>	Diagnosis of covered use, laboratory confirmation of hepatitis C virus (HCV) and genotype, documentation of whether cirrhosis is present or not and whether or not it is compensated or decompensated, submission of previous treatment regimen, confirmation a test for HBV infection (HBsAg and anti-HBc) was completed.
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restrictions</b>	
<b>Coverage Duration</b>	12 weeks
<b>Other Criteria</b>	PA applies to all.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# VOTRIENT (pazopanib)

## Products Affected

- VOTRIENT

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, uncontrolled hypertension, uncorrected hypokalemia, hypocalcemia, or hypomagnesemia, coadministration with strong CYP3A4 inducers or drugs that can prolong the QT interval
Required Medical Information	Diagnosis of covered use, submission of baseline blood pressure, ALT, AST, bilirubin, and serum potassium, calcium, and magnesium, pregnancy status for female patients of childbearing potential. For soft tissue sarcoma, submission of previous chemotherapy regimen(s).
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# VRAYLAR (cariprazine)

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## Products Affected

- VRAYLAR ORAL CAPSULE
- VRAYLAR ORAL CAPSULE THERAPY PACK

PA Criteria	Criteria Details
Exclusion Criteria	Dementia-related psychosis, severe hepatic impairment, severe renal impairment (creatinine clearance less than 30 mL/min), coadministration with CYP3A4 inducers
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology and psychiatry
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## WAKIX (pitolisant)

### Products Affected

- WAKIX ORAL TABLET 17.8 MG, 4.45 MG

PA Criteria	Criteria Details
Exclusion Criteria	Severe hepatic impairment, end-stage renal disease, known QT interval prolongation, symptomatic bradycardia, uncorrected hypokalemia or hypomagnesemia, coadministration with medications that prolong the QT interval
Required Medical Information	Diagnosis of covered use, submission of serum potassium and magnesium.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology and sleep medicine
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated serum potassium and magnesium since the previous authorization will be required for subsequent annual reauthorizations. For excessive daytime sleepiness associated with narcolepsy, pitolisant will be authorized only if the patient previously tried and had an inadequate clinical response or an intolerance to armodafinil or modafinil.
Indications	All Medically-accepted Indications.
Off Label Uses	

## WELIREG (belzutifan)

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### Products Affected

- WELIREG

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use including confirmation patient does not require immediate surgery, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# WHITE BLOOD CELL STIMULATORS

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## Products Affected

- NIVESTYM
- ZARXIO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	For approval of Nivestym, the patient must have tried and failed to have an adequate response to or had an intolerance to Zarxio. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# XALKORI (crizotinib)

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## Products Affected

- XALKORI

PA Criteria	Criteria Details
Exclusion Criteria	Congenital long QT syndrome, coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of results showing tumor is ALK or ROS1-positive as detected by an FDA-approved test, pregnancy status for female patients of childbearing potential.
Age Restrictions	For ALK-positive systemic anaplastic large cell lymphoma only, 1 year of age to 21 years of age
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## XATMEP (methotrexate oral solution)

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### Products Affected

- XATMEP

PA Criteria	Criteria Details
Exclusion Criteria	Pregnancy (for polyarticular juvenile idiopathic arthritis [pJIA] indication only)
Required Medical Information	Diagnosis of covered use. For acute lymphoblastic leukemia, confirmation that medication is being used as a component of a combination chemotherapy maintenance regimen.
Age Restrictions	2 years of age through 18 years of age
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



## XCOPRI (cenobamate)

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### Products Affected

- XCOPRI (250 MG DAILY DOSE)
- XCOPRI (350 MG DAILY DOSE)
- XCOPRI ORAL TABLET 100 MG, 150 MG, 200 MG, 50 MG
- XCOPRI ORAL TABLET THERAPY PACK

PA Criteria	Criteria Details
Exclusion Criteria	Familial short QT syndrome, patients with end-stage renal disease (creatinine clearance less than 15 mL/min) undergoing dialysis, severe hepatic impairment
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to neurology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## XERMELO (telotristat)

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### Products Affected

- XERMELO

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation patient has been on at least 12 weeks of prior somatostatin analog therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	12 weeks
Other Criteria	PA applies to all. Continuation of therapy requires that symptoms have stabilized or improved and that the patient has not experienced episodes of severe constipation.
Indications	All Medically-accepted Indications.
Off Label Uses	

## XGEVA (denosumab)

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### Products Affected

- XGEVA

PA Criteria	Criteria Details
Exclusion Criteria	Hypocalcemia
Required Medical Information	Diagnosis of covered use, submission of serum calcium level, pregnancy status for female patients of childbearing potential.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# XOLAIR (omalizumab)

## Products Affected

- XOLAIR

PA Criteria	Criteria Details
<b>Exclusion Criteria</b>	Patients whose pre-treatment serum IgE level and body weight place them in the "insufficient data to recommend a dose" category based on dosing charts in the prescribing information
<b>Required Medical Information</b>	Diagnosis of covered use. For asthma, documentation that patient's symptoms are poorly controlled with inhaled corticosteroids, submission of patient's current body weight, pre-treatment serum IgE level, pulmonary function test results including FEV1, positive skin test result or demonstrated in vitro reactivity (RAST test) to a perennial aeroallergen, frequency of inhaled short-acting beta2-agonist therapy, frequency of daily and nighttime symptoms and exacerbations, and effect of exacerbations on activity. For chronic idiopathic urticaria, documentation that the patient continues to experience severe itching and hives despite the use of an H1 antihistamine at an approved dose. For nasal polyps, documentation that patient's symptoms are poorly controlled with intranasal corticosteroids and current intranasal corticosteroid therapy.
<b>Age Restrictions</b>	6 years of age or older
<b>Prescriber Restrictions</b>	Restricted to allergy, dermatology, immunology, otolaryngology/otorhinolaryngology, and pulmonology
<b>Coverage Duration</b>	1 year
<b>Other Criteria</b>	PA applies to all. Submission of objective documentation of symptomatic improvement and updated patient weight will be required for subsequent annual reauthorizations. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
<b>Indications</b>	All Medically-accepted Indications.
<b>Off Label Uses</b>	

# XOSPATA (gilteritinib)

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## Products Affected

- XOSPATA

PA Criteria	Criteria Details
Exclusion Criteria	Uncorrected hypokalemia or hypomagnesemia, coadministration with dual strong CYP3A/P-glycoprotein inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of FLT3 mutation, serum calcium and magnesium, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

# XPOVIO (selinexor)

## Products Affected

- XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 50 MG
- XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG
- XPOVIO (40 MG TWICE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG
- XPOVIO (60 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 60 MG
- XPOVIO (60 MG TWICE WEEKLY)
- XPOVIO (80 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG
- XPOVIO (80 MG TWICE WEEKLY)

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation of treatment failure with or intolerance to all prior therapies to match the indication, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All FDA-approved Indications.
Off Label Uses	

## XURIDEN (uridine triacetate)

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### Products Affected

- XURIDEN

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, submission of baseline CBC including neutrophil count and mean corpuscular volume, baseline urine orotic acid level.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated urine orotic acid level and CBC including neutrophil count and mean corpuscular volume since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ZEJULA (niraparib)

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### Products Affected

- ZEJULA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use, documentation of treatment failure with or intolerance to all prior therapies to match the indication, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	



## ZERBAXA (ceftolozane/tazobactam)

### Products Affected

- ZERBAXA

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use. For complicated intra-abdominal infections, confirmation patient will receive concurrent metronidazole therapy.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	For UTI including pyelonephritis, 7 days. For all other FDA-approved indications, 14 days.
Other Criteria	PA applies to all. A description of how the drug will be used and/or obtained may be required to determine whether it will be covered under Medicare Part B or Part D. If the medication will be self-administered after proper training or the provider agrees to have the medication filled at a pharmacy and delivered to the office by the patient for provider administration, it may be covered under Part D. If these conditions are not satisfied this medication may be covered under Part B.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ZILEUTON ER

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## Products Affected

- ZILEUTON ER

PA Criteria	Criteria Details
Exclusion Criteria	Active liver disease or persistent hepatic function elevation enzyme greater than or equal to 3 times the upper limit of normal
Required Medical Information	Diagnosis of covered use, submission of hepatic function enzymes and serum bilirubin.
Age Restrictions	12 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all. Updated hepatic function enzymes and serum bilirubin since the previous authorization will be required for subsequent annual reauthorizations.
Indications	All Medically-accepted Indications.
Off Label Uses	

# ZOKINVY (lonafarnib)

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## Products Affected

- ZOKINVY ORAL CAPSULE 50 MG, 75 MG

PA Criteria	Criteria Details
Exclusion Criteria	Body surface area less than 0.39 m <sup>2</sup>
Required Medical Information	Diagnosis of covered use including results of genetic testing supporting diagnosis, pregnancy status for female patients of childbearing potential.
Age Restrictions	
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to all.
Indications	All FDA-approved Indications.
Off Label Uses	

# ZONTIVITY (vorapaxar)

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## Products Affected

- ZONTIVITY

PA Criteria	Criteria Details
Exclusion Criteria	History of stroke, transient ischemic attack, or intracranial hemorrhage, active pathological bleeding, severe hepatic impairment, coadministration with strong CYP3A inhibitors or inducers
Required Medical Information	Diagnosis of covered use, confirmation that patient has not had prior stroke, transient ischemic attack, or intracranial hemorrhage, documentation of concurrent use with aspirin and/or clopidogrel.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ZORBTIVE (somatropin)

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### Products Affected

- ZORBTIVE

PA Criteria	Criteria Details
Exclusion Criteria	Active malignancy, acute critical illness, active proliferative or severe non-proliferative diabetic retinopathy
Required Medical Information	Diagnosis of covered use.
Age Restrictions	18 years of age or older
Prescriber Restrictions	
Coverage Duration	4 weeks
Other Criteria	PA applies to all. Zorbtive is indicated only for the treatment of short bowel syndrome and uses outside of this indication will not be approved.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ZTALMY (ganaxolone)

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### Products Affected

- ZTALMY

PA Criteria	Criteria Details
Exclusion Criteria	
Required Medical Information	Diagnosis of covered use confirmed by genetic testing including either (a) a CDKL5 gene that is pathogenic or likely to be pathogenic or (b) CDKL5 deficiency, documentation of failure of at least two previous anticonvulsant therapies, submission of baseline monthly major motor seizure (defined as bilateral tonic, generalized tonic-clonic, bilateral clonic, atonic, or focal to bilateral tonic-clonic seizure) frequency.
Age Restrictions	
Prescriber Restrictions	Restricted to neurology
Coverage Duration	Initially 6 months, then 1 year
Other Criteria	PA applies to new starts only. Submission of documentation demonstrating a sustained reduction in monthly major motor seizure frequency is required for reauthorization.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ZYDELIG (idelalisib)

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### Products Affected

- ZYDELIG

PA Criteria	Criteria Details
Exclusion Criteria	History of serious hypersensitivity reactions, including toxic epidermal necrolysis with any drug, coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to hematology and oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	

## ZYKADIA (ceritinib)

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### Products Affected

- ZYKADIA ORAL TABLET

PA Criteria	Criteria Details
Exclusion Criteria	Coadministration with strong CYP3A inducers
Required Medical Information	Diagnosis of covered use, submission of FDA-approved test confirming presence of ALK-positive tumor, pregnancy status for female patients of childbearing potential.
Age Restrictions	18 years of age or older
Prescriber Restrictions	Restricted to oncology
Coverage Duration	1 year
Other Criteria	PA applies to new starts only.
Indications	All Medically-accepted Indications.
Off Label Uses	





## Index

ABILIFY MYCITE MAINTENANCE KIT ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG.....	2	BUTALBITAL-ACETAMINOPHEN ORAL TABLET 50-300 MG, 50-325 MG.....	28
ABILIFY MYCITE ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG.....	2	BUTALBITAL-APAP-CAFF-COD.....	28
ABILIFY MYCITE STARTER KIT ORAL TABLET 10 MG, 15 MG, 2 MG, 20 MG, 30 MG, 5 MG.....	2	<i>butalbital-apap-caffeine oral capsule</i> .....	28
<i>abiraterone acetate oral tablet 250 mg</i> .....	187	<i>butalbital-apap-caffeine oral tablet 50-325-40 mg</i> ....	28
ACTEMRA ACTPEN.....	22	<i>butalbital-asa-caff-codeine</i> .....	28
ACTEMRA SUBCUTANEOUS.....	22	<i>butalbital-aspirin-caffeine oral capsule</i> .....	28
ACTHAR.....	3	BYLVAY.....	29
ACTIMMUNE.....	4	BYLVAY (PELLETS).....	29
ADEMPAS.....	5	CABLIVI.....	30
AIMOVIG SUBCUTANEOUS SOLUTION AUTO-INJECTOR 140 MG/ML, 70 MG/ML.....	38	CABOMETYX.....	31
AJOVY.....	38	CALQUENCE.....	32
AKYNZEO ORAL.....	6	CAMCEVI.....	88
ALECENSA.....	7	<i>camzyos</i> .....	33
ALUNBRIG.....	9	CAPLYTA ORAL CAPSULE 10.5 MG, 21 MG, 42 MG.....	34
<i>alyq</i> .....	175	CAPRELSA.....	35
<i>ambrisentan</i> .....	69	<i>carbinoxamine maleate oral solution</i> .....	81
AMVUTTRA.....	11	<i>carbinoxamine maleate oral tablet 4 mg</i> .....	81
ANADROL-50.....	12	<i>carglumic acid</i> .....	36
ANDRODERM TRANSDERMAL PATCH 24 HOUR.....	238	CARIMUNE NF INTRAVENOUS SOLUTION RECONSTITUTED 12 GM, 6 GM.....	99
<i>apomorphine hcl subcutaneous</i> .....	173	CERDELGA.....	37
ARALAST NP INTRAVENOUS SOLUTION RECONSTITUTED 1000 MG, 500 MG.....	8	CHENODAL.....	39
ARANESP (ALBUMIN FREE) INJECTION SOLUTION 100 MCG/ML, 200 MCG/ML, 25 MCG/ML, 300 MCG/ML, 40 MCG/ML, 60 MCG/ML.....	13	CHOLBAM.....	40
ARANESP (ALBUMIN FREE) INJECTION SOLUTION PREFILLED SYRINGE.....	13	CIMZIA PREFILLED.....	22
ARCALYST.....	14	CIMZIA STARTER KIT.....	22
ARIKAYCE.....	15	CIMZIA SUBCUTANEOUS KIT 2 X 200 MG.....	22
<i>ascomp-codeine</i> .....	28	<i>clemastine fumarate oral tablet 2.68 mg</i> .....	81
AURYXIA.....	16	COMETRIQ (100 MG DAILY DOSE) ORAL KIT 80 & 20 MG.....	41
AUSTEDO ORAL TABLET 12 MG, 6 MG, 9 MG.....	263	COMETRIQ (140 MG DAILY DOSE) ORAL KIT 3 X 20 MG & 80 MG.....	41
AYVAKIT ORAL TABLET 100 MG, 200 MG, 25 MG, 300 MG, 50 MG.....	17	COMETRIQ (60 MG DAILY DOSE).....	41
BAFIERTAM.....	83	COPIKTRA ORAL CAPSULE 15 MG, 25 MG.....	42
BALVERSA.....	18	CORLANOR.....	43
BENLYSTA SUBCUTANEOUS.....	20	CORTROPHIN.....	44
BESREMI.....	21	COTELLIC.....	45
<i>bexarotene external</i> .....	229	<i>cypheptadine hcl oral</i> .....	81
BIVIGAM INTRAVENOUS SOLUTION 5 GM/50ML.....	99	CYSTADROPS.....	46
<i>bosentan</i> .....	69	CYSTARAN.....	46
BOSULIF.....	23	<i>dalfampridine er</i> .....	47
BRAFTOVI ORAL CAPSULE 75 MG.....	24	DAURISMO ORAL TABLET 100 MG, 25 MG.....	48
BRIVIACT ORAL.....	25	DAYVIGO ORAL TABLET 10 MG, 5 MG.....	49
BRONCHITOL.....	26	<i>deferiasirox granules</i> .....	50
BRUKINSA.....	27	<i>deferiasirox oral tablet</i> .....	50
BUPAP ORAL TABLET 50-300 MG.....	28	<i>deferiasirox oral tablet soluble</i> .....	50
		<i>deferiprone</i> .....	51
		DIACOMIT.....	52
		<i>diclofenac epolamine external</i> .....	56
		<i>diclofenac sodium external gel</i> .....	53, 55
		<i>diclofenac sodium external solution 1.5 %</i> .....	54
		DIFICID.....	57

<i>digitek oral tablet 250 mcg</i> .....	58	GAMMAPLEX INTRAVENOUS SOLUTION 10	
<i>digox oral tablet 250 mcg</i> .....	58	GM/100ML, 10 GM/200ML, 20 GM/200ML, 20	
<i>digoxin oral tablet 250 mcg</i> .....	58	GM/400ML, 5 GM/100ML, 5 GM/50ML .....	99
<i>diphenhydramine hcl oral elixir</i> .....	81	GAMUNEX-C .....	99
DOPTELET ORAL TABLET 20 MG .....	59	GATTEX .....	85
<i>dronabinol</i> .....	60	GAVRETO .....	86
<i>droxidopa</i> .....	61	GENOTROPIN .....	89
DUOBRII .....	62	GENOTROPIN MINISQUICK .....	89
DUOPA ENTERAL .....	63	GILOTRIF .....	87
DUPIXENT .....	64	GLASSIA .....	8
EGRIFTA SV .....	65	GOCOVRI .....	10
ELIGARD .....	88	HAEGARDA .....	91
EMFLAZA .....	66	HARVONI ORAL PACKET .....	123
EMGALITY .....	38	HARVONI ORAL TABLET 45-200 MG, 90-400 MG .....	123
EMGALITY (300 MG DOSE) .....	38	HETLIOZ .....	92
EMPAVELI .....	67	HUMATROPE .....	89
EMSAM .....	68	<i>hydroxyzine hcl oral tablet</i> .....	81
ENSPRYNG .....	70	<i>hydroxyzine pamoate oral</i> .....	81
EPCLUSA ORAL PACKET .....	210	IBRANCE .....	93
EPCLUSA ORAL TABLET 200-50 MG, 400-100 MG .....	210	<i>icatibant acetate</i> .....	90
EPIDIOLEX .....	71	ICLUSIG .....	94
ERIVEDGE .....	72	IDHIFA .....	95
ERLEADA .....	187	ILARIS SUBCUTANEOUS SOLUTION .....	96
<i>erlotinib hcl</i> .....	73	IMBRUVICA .....	97
ESBRIET ORAL CAPSULE .....	74	IMCIVREE .....	98
<i>eszopiclone</i> .....	204	INBRIJA .....	173
<i>everolimus oral tablet 10 mg, 2.5 mg, 5 mg, 7.5 mg</i> ..	75	INCRELEX .....	138
<i>everolimus oral tablet soluble</i> .....	75	INGREZZA ORAL CAPSULE 40 MG, 60 MG, 80 MG ...	263
EVRYSDI .....	76	INGREZZA ORAL CAPSULE THERAPY PACK .....	263
EXKIVITY .....	77	INLYTA .....	101
<i>ezetimibe-simvastatin oral tablet 10-80 mg</i> .....	207	INQOVI .....	102
FABIOR .....	182	INREBIC .....	103
FASENRA .....	104	INTRON A .....	106
FASENRA PEN .....	104	INVEGA HAFYERA .....	107
<i>fentanyl citrate buccal</i> .....	78	INVEGA TRINZA INTRAMUSCULAR SUSPENSION	
FERRIPROX ORAL SOLUTION .....	51	PREFILLED SYRINGE .....	108
FINTEPLA .....	79	IRESSA .....	109
FIRDAPSE .....	80	ISTURISA .....	110
FIRMAGON (240 MG DOSE) .....	88	<i>itraconazole oral</i> .....	111
FIRMAGON SUBCUTANEOUS SOLUTION		JAKAFI .....	112
RECONSTITUTED 80 MG .....	88	JUBLIA .....	242
FLEBOGAMMA DIF .....	99	JUXTAPID ORAL CAPSULE 10 MG, 20 MG, 30 MG, 5	
FORTEO SUBCUTANEOUS SOLUTION 600		MG .....	113
MCG/2.4ML .....	237	JYNARQUE .....	114
FORTEO SUBCUTANEOUS SOLUTION PEN-INJECTOR	237	KALYDECO .....	115
FOTIVDA .....	82	KERENDIA .....	116
GALAFOLD .....	84	<i>ketoconazole oral</i> .....	117
GAMMAGARD .....	99	KEVEYIS .....	118
GAMMAGARD S/D LESS IGA .....	99	KEVZARA .....	22
GAMMAKED INJECTION SOLUTION 1 GM/10ML, 10		KISQALI (200 MG DOSE) .....	119
GM/100ML, 20 GM/200ML, 5 GM/50ML .....	99	KISQALI (400 MG DOSE) .....	119
		KISQALI (600 MG DOSE) .....	119
		KISQALI FEMARA (400 MG DOSE) .....	119

KISQALI FEMARA (600 MG DOSE) .....	119	<i>miglustat</i> .....	145
KISQALI FEMARA(200 MG DOSE) .....	119	MYALEPT .....	146
KORLYM .....	120	MYCAPSSA .....	147
KOSELUGO .....	121	MYFEMBREE .....	148
KYNMOBI .....	173	MYTESI .....	149
KYNMOBI TITRATION KIT .....	173	NAMZARIC .....	150
<i>lapatinib ditosylate</i> .....	122	NATPARA .....	151
<i>ledipasvir-sofosbuvir</i> .....	123	NAYZILAM .....	105
<i>lenalidomide</i> .....	197	NERLYNX .....	152
LENVIMA (10 MG DAILY DOSE) .....	124	NEULASTA ONPRO .....	176
LENVIMA (12 MG DAILY DOSE) .....	124	NEULASTA SUBCUTANEOUS SOLUTION PREFILLED	
LENVIMA (14 MG DAILY DOSE) .....	124	SYRINGE .....	176
LENVIMA (18 MG DAILY DOSE) .....	124	NEXLETOL .....	19
LENVIMA (20 MG DAILY DOSE) .....	124	NEXLIZET .....	19
LENVIMA (24 MG DAILY DOSE) .....	124	NINLARO .....	154
LENVIMA (4 MG DAILY DOSE) .....	124	<i>nitisinone</i> .....	155
LENVIMA (8 MG DAILY DOSE) .....	124	NITYR .....	155
LEUKINE INJECTION SOLUTION RECONSTITUTED .....	125	NIVESTYM .....	270
<i>leuprolide acetate injection</i> .....	88	NORDITROPIN FLEXPPO SUBCUTANEOUS SOLUTION	
<i>lidocaine external patch 5 %</i> .....	126	PEN-INJECTOR .....	89
LIVMARLI .....	127	NUBEQA .....	187
LIVTENCITY .....	128	NUCALA SUBCUTANEOUS SOLUTION AUTO-	
LONSURF .....	129	INJECTOR .....	104
LORBRENA ORAL TABLET 100 MG, 25 MG .....	130	NUCALA SUBCUTANEOUS SOLUTION PREFILLED	
LUCEMYRA .....	131	SYRINGE 100 MG/ML .....	104
LUMAKRAS .....	132	NUCALA SUBCUTANEOUS SOLUTION	
LUPANETA PACK .....	88	RECONSTITUTED .....	104
LUPKYNIS .....	133	NUEDEXTA .....	156
LUPRON DEPOT (1-MONTH) .....	88	NUPLAZID ORAL CAPSULE .....	157
LUPRON DEPOT (3-MONTH) .....	88	NUPLAZID ORAL TABLET 10 MG .....	157
LUPRON DEPOT (4-MONTH) .....	88	NURTEC .....	38
LUPRON DEPOT (6-MONTH) .....	88	NUTROPIN AQ NUSPIN 10 SUBCUTANEOUS	
LUPRON DEPOT-PED (1-MONTH) INTRAMUSCULAR		SOLUTION PEN-INJECTOR .....	89
KIT 11.25 MG, 15 MG .....	88	NUTROPIN AQ NUSPIN 20 SUBCUTANEOUS	
LUPRON DEPOT-PED (3-MONTH) INTRAMUSCULAR		SOLUTION PEN-INJECTOR .....	89
KIT 30 MG (PED) .....	88	NUTROPIN AQ NUSPIN 5 SUBCUTANEOUS	
LYBALVI .....	134	SOLUTION PEN-INJECTOR .....	89
LYNPARZA ORAL TABLET .....	135	OALIVA ORAL TABLET 10 MG, 5 MG .....	158
MAVENCLAD (10 TABS) .....	136	OCTAGAM INTRAVENOUS SOLUTION 1 GM/20ML,	
MAVENCLAD (4 TABS) .....	136	10 GM/100ML, 10 GM/200ML, 2 GM/20ML, 2.5	
MAVENCLAD (5 TABS) .....	136	GM/50ML, 20 GM/200ML, 25 GM/500ML, 5	
MAVENCLAD (6 TABS) .....	136	GM/100ML, 5 GM/50ML .....	99
MAVENCLAD (7 TABS) .....	136	ODOMZO .....	159
MAVENCLAD (8 TABS) .....	136	OFEV .....	160
MAVENCLAD (9 TABS) .....	136	ONUREG .....	161
MAVYRET .....	137	OPSUMIT .....	69
<i>megestrol acetate oral suspension 40 mg/ml, 400</i>		ORENITRAM .....	162
<i>mg/10ml, 625 mg/5ml</i> .....	139	ORFADIN ORAL CAPSULE 20 MG .....	155
MEKINIST .....	140	ORFADIN ORAL SUSPENSION .....	155
MEKTOVI .....	24	ORGOVYX .....	163
<i>methamphetamine hcl</i> .....	141	ORIAHNN .....	164
METHITEST .....	144	ORILISSA ORAL TABLET 150 MG, 200 MG .....	165
<i>methyltestosterone oral</i> .....	144	ORKAMBI .....	166

ORLADEYO .....	91	PYRUKYND TAPER PACK ORAL TABLET THERAPY	
OSMOLEX ER ORAL TABLET ER 24 HOUR THERAPY		PACK 5 MG, 7 X 20 MG & 7 X 5 MG, 7 X 50 MG & 7 X	
PACK .....	167	20 MG .....	188
OSMOLEX ER ORAL TABLET EXTENDED RELEASE 24		QINLOCK .....	189
HOUR 129 MG, 193 MG, 258 MG .....	167	QULIPTA .....	38
OTEZLA .....	22	QUVIVIQ .....	190
OTREXUP SUBCUTANEOUS SOLUTION AUTO-		RADICAVA ORS .....	191
INJECTOR 10 MG/0.4ML, 12.5 MG/0.4ML, 15		RADICAVA ORS STARTER KIT .....	191
MG/0.4ML, 17.5 MG/0.4ML, 20 MG/0.4ML, 22.5		RASUVO SUBCUTANEOUS SOLUTION AUTO-	
MG/0.4ML, 25 MG/0.4ML .....	142	INJECTOR 10 MG/0.2ML, 12.5 MG/0.25ML, 15	
OXBRYTA .....	168	MG/0.3ML, 17.5 MG/0.35ML, 20 MG/0.4ML, 22.5	
OXERVATE .....	169	MG/0.45ML, 25 MG/0.5ML, 30 MG/0.6ML, 7.5	
PALYNZIQ .....	171	MG/0.15ML .....	142
PANRETIN .....	172	RAVICTI .....	192
PEGASYS PROCLICK SUBCUTANEOUS SOLUTION 135		RECORLEV .....	193
MCG/0.5ML .....	182	REDITREX .....	142
PEGASYS PROCLICK SUBCUTANEOUS SOLUTION		REPATHA .....	174
AUTO-INJECTOR 135 MCG/0.5ML .....	182	REPATHA PUSHTRONEX SYSTEM .....	174
PEGASYS SUBCUTANEOUS SOLUTION .....	182	REPATHA SURECLICK .....	174
PEGASYS SUBCUTANEOUS SOLUTION PREFILLED		RETACRIT INJECTION SOLUTION 10000 UNIT/ML,	
SYRINGE .....	182	2000 UNIT/ML, 20000 UNIT/ML, 3000 UNIT/ML,	
PEG-INTRON REDIPEN SUBCUTANEOUS KIT 50		4000 UNIT/ML, 40000 UNIT/ML .....	194
MCG/0.5ML .....	182	RETEVMO ORAL CAPSULE 40 MG, 80 MG .....	195
PEGINTRON SUBCUTANEOUS KIT 50 MCG/0.5ML ...	182	REVCovi .....	196
PEG-INTRON SUBCUTANEOUS KIT 50 MCG/0.5ML ...	182	REVLIMID .....	197
PEMAZYRE ORAL TABLET 13.5 MG, 4.5 MG, 9 MG ...	177	REZUROCK .....	198
PIQRAY (200 MG DAILY DOSE) .....	178	ROZLYTREK ORAL CAPSULE 100 MG, 200 MG .....	199
PIQRAY (250 MG DAILY DOSE) .....	178	RUBRACA .....	200
PIQRAY (300 MG DAILY DOSE) .....	178	RUCONEST .....	90
<i>pirfenidone oral tablet 267 mg, 801 mg</i> .....	74	RYDAPT .....	201
POMALYST .....	179	<i>sajazir</i> .....	90
PRALUENT SUBCUTANEOUS SOLUTION AUTO-		SAMSCA ORAL TABLET 15 MG .....	241
INJECTOR .....	174	<i>sapropterin dihydrochloride oral packet</i> .....	202
PRETOMANID .....	180	<i>sapropterin dihydrochloride oral tablet</i> .....	202
PREVYMIS ORAL .....	181	SCEMBLIX ORAL TABLET 20 MG, 40 MG .....	203
PRIVIGEN .....	99	SEROSTIM SUBCUTANEOUS SOLUTION	
PROCYSBI .....	183	RECONSTITUTED 4 MG, 5 MG, 6 MG .....	205
PROLASTIN-C INTRAVENOUS SOLUTION		SIGNIFOR .....	206
RECONSTITUTED .....	8	<i>sildenafil citrate oral suspension reconstituted</i> .....	175
PROLIA SUBCUTANEOUS SOLUTION PREFILLED		<i>sildenafil citrate oral tablet 20 mg</i> .....	175
SYRINGE .....	184	SIMPONI SUBCUTANEOUS SOLUTION AUTO-	
PROMACTA ORAL PACKET .....	185	INJECTOR .....	22
PROMACTA ORAL TABLET 12.5 MG, 25 MG, 50 MG,		SIMPONI SUBCUTANEOUS SOLUTION PREFILLED	
75 MG .....	185	SYRINGE .....	22
<i>promethazine hcl oral</i> .....	186	<i>simvastatin oral tablet 80 mg</i> .....	207
PROMETHAZINE HCL RECTAL SUPPOSITORY 12.5		SIRTURO .....	208
MG .....	186	SIVEXTRO .....	209
<i>promethazine hcl rectal suppository 25 mg</i> .....	186	<i>sofosbuvir-velpatasvir</i> .....	210
<i>promethazine vc plain</i> .....	186	SOMATULINE DEPOT .....	211
<i>promethazine-phenylephrine</i> .....	186	SOMAVERT .....	212
<i>promethegan rectal suppository 25 mg, 50 mg</i> .....	186	<i>sorafenib tosylate</i> .....	153
PYRUKYND .....	188	SOVALDI ORAL PACKET .....	213
		SOVALDI ORAL TABLET 200 MG, 400 MG .....	213

SPRYCEL ORAL TABLET 100 MG, 140 MG, 20 MG, 50 MG, 70 MG, 80 MG .....	214	TOLSURA .....	111
STIVARGA .....	215	<i>tolvaptan</i> .....	241
SUCRAID .....	216	TRACLEER ORAL TABLET SOLUBLE .....	69
<i>sunitinib malate</i> .....	218	TRELSTAR MIXJECT .....	88
SUNOSI ORAL TABLET 150 MG, 75 MG .....	217	TREMFYA .....	22
SYMDEKO .....	219	TRIKAFTA ORAL TABLET THERAPY PACK 100-50-75 & 150 MG, 50-25-37.5 & 75 MG .....	244
SYMLINPEN 120 SUBCUTANEOUS SOLUTION PEN-INJECTOR .....	220	TRUSELTIQ (100MG DAILY DOSE) .....	245
SYMLINPEN 60 SUBCUTANEOUS SOLUTION PEN-INJECTOR .....	220	TRUSELTIQ (125MG DAILY DOSE) .....	245
SYMPROIC .....	221	TRUSELTIQ (50MG DAILY DOSE) .....	245
SYNAREL .....	222	TRUSELTIQ (75MG DAILY DOSE) .....	245
SYNDROS .....	60	TUKYSA ORAL TABLET 150 MG, 50 MG .....	246
SYNRIBO .....	223	TURALIO .....	247
TABRECTA ORAL TABLET 150 MG, 200 MG .....	224	TYMLOS .....	248
<i>tadalafil (pah)</i> .....	175	UBRELVY .....	38
TAFINLAR .....	226	UDENYCA .....	176
TAGRISSO .....	227	UPNEEQ .....	249
<i>takhzyro subcutaneous solution</i> .....	91	UPTRAVI ORAL TABLET 1000 MCG, 1200 MCG, 1400 MCG, 1600 MCG, 200 MCG, 400 MCG, 600 MCG, 800 MCG .....	250
TAKHZYRO SUBCUTANEOUS SOLUTION PREFILLED SYRINGE .....	91	UPTRAVI ORAL TABLET THERAPY PACK .....	250
TALZENNA ORAL CAPSULE 0.25 MG, 0.5 MG, 0.75 MG, 1 MG .....	228	VABOMERE .....	182
TARPEYO .....	230	VALCHLOR .....	251
TASIGNA .....	231	VALTOCO 10 MG DOSE .....	105
<i>tavorole</i> .....	242	VALTOCO 15 MG DOSE .....	105
TAVALISSE ORAL TABLET 100 MG, 150 MG .....	232	VALTOCO 20 MG DOSE .....	105
TAVNEOS .....	233	VALTOCO 5 MG DOSE .....	105
<i>tazarotene external cream</i> .....	182	VECAMYL .....	252
<i>tazarotene external foam</i> .....	182	VEMLIDY .....	253
TAZORAC EXTERNAL CREAM 0.05 % .....	182	VENCLEXTA ORAL TABLET 10 MG, 100 MG, 50 MG ..	254
TAZORAC EXTERNAL GEL .....	182	VENCLEXTA STARTING PACK .....	254
TAZVERIK .....	234	VENTAVIS .....	255
TEGSEDI .....	235	VERQUVO ORAL TABLET 10 MG, 2.5 MG, 5 MG .....	256
TENCON ORAL TABLET 50-325 MG .....	28	VERZENIO .....	257
TEPMETKO .....	236	VIBERZI .....	258
TERIPARATIDE (RECOMBINANT) .....	237	<i>vijoice oral tablet therapy pack 125 mg, 200 &amp; 50 mg, 50 mg</i> .....	259
<i>testosterone cypionate injection solution 200 mg/ml</i> .....	100	VITRAKVI ORAL CAPSULE 100 MG, 25 MG .....	260
<i>testosterone cypionate intramuscular solution 100 mg/ml, 200 mg/ml</i> .....	100	VITRAKVI ORAL SOLUTION .....	260
<i>testosterone enanthate intramuscular solution</i> .....	100	VIVJOA .....	261
TESTOSTERONE TRANSDERMAL GEL 10 MG/ACT (2%) .....	238	VIZIMPRO ORAL TABLET 15 MG, 30 MG, 45 MG .....	262
<i>testosterone transdermal gel 12.5 mg/act (1%), 20.25 mg/1.25gm (1.62%), 20.25 mg/act (1.62%), 25 mg/2.5gm (1%), 40.5 mg/2.5gm (1.62%), 50 mg/5gm (1%)</i> .....	238	VONJO .....	264
<i>testosterone transdermal solution</i> .....	238	VOSEVI .....	265
<i>tetrabenazine</i> .....	263	VOTRIENT .....	266
TIBSOVO .....	239	VRAYLAR ORAL CAPSULE .....	267
TOBI PODHALER .....	240	VRAYLAR ORAL CAPSULE THERAPY PACK .....	267
		VTAMA .....	243
		VTOL LQ .....	28
		VUMERITY .....	83
		VYNDAMAX .....	225
		VYNDAQEL .....	225
		WAKIX ORAL TABLET 17.8 MG, 4.45 MG .....	268
		WELIREG .....	269

XALKORI .....	271
XATMEP .....	272
XCOPRI (250 MG DAILY DOSE) .....	273
XCOPRI (350 MG DAILY DOSE) .....	273
XCOPRI ORAL TABLET 100 MG, 150 MG, 200 MG, 50 MG .....	273
XCOPRI ORAL TABLET THERAPY PACK .....	273
XERMELO .....	274
XGEVA .....	275
XOLAIR .....	276
XOSPATA .....	277
XPOVIO (100 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 50 MG .....	278
XPOVIO (40 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG .....	278
XPOVIO (40 MG TWICE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG .....	278
XPOVIO (60 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 60 MG .....	278
XPOVIO (60 MG TWICE WEEKLY) .....	278
XPOVIO (80 MG ONCE WEEKLY) ORAL TABLET THERAPY PACK 20 MG, 40 MG .....	278
XPOVIO (80 MG TWICE WEEKLY) .....	278
XTANDI .....	187
XURIDEN .....	279
XYREM .....	170
XYWAV .....	170
<i>zaleplon</i> .....	204
ZARXIO .....	270
ZEBUTAL ORAL CAPSULE 50-325-40 MG .....	28
ZEJULA .....	280
ZELBORAF .....	45
ZEMAIRA .....	8
ZEPOSIA .....	22
ZEPOSIA 7-DAY STARTER PACK .....	22
ZEPOSIA STARTER KIT .....	22
ZERBAXA .....	281
ZILEUTON ER .....	282
ZOKINVY ORAL CAPSULE 50 MG, 75 MG .....	283
<i>zolpidem tartrate oral</i> .....	204
ZONTIVITY .....	284
ZORBTIVE .....	285
ZORYVE .....	243
ZTALMY .....	286
ZYDELIG .....	287
ZYKADIA ORAL TABLET .....	288