



**DEPT. OF HEALTH AND HUMAN SERVICES** 

# Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

For the most up to date list of covered drugs consult the Drug Lookup on the Nebraska Medicaid Website at <a href="https://druglookup.fhsc.com/druglookupweb/?client=nestate">https://druglookup.fhsc.com/druglookupweb/?client=nestate</a>

- PDMP Check Requirements- Nebraska Medicaid providers are required to check the prescription drug history in the statewide PDMP before prescribing CII controlled substances to certain Medicaid beneficiaries. (Exemption to this requirement are for beneficiaries receiving cancer treatment, hospice/palliative care, or in long-term care facilities). If not able to check the PDMP, then provider is required to document good faith effort, including reasons why unable to conduct the check and may be required to submit documentation to the State upon request.
  - PDMP check requirements are under Section 5042 of the SUPPORT for Patients and Communities Act, consistent with section 1944 of the Social Security Act [42 U.S.C. 1396w-3a], beginning October 1, 2021.
- **Opioids** The maximum opioid dose covered will decrease from 120 Morphine Milligram Equivalents (MME) per day to 90 Morphine Milligram Equivalents (MME) per day. (beginning December 1, 2020)

#### **Non-Preferred Drug Coverage**

Class and drug-specific therapeutic trial and failure requirements are found within this document. Examples of non-preferred exception criteria include:

- Adverse reaction to preferred drugs
- Allergy to preferred drugs
- Contraindication to preferred drugs
- Documentation of inability to swallow solid dosage forms

Specific Class Prior Authorization forms can be found within the PDL class listings and at: https://nebraska.fhsc.com/priorauth/paforms.asp

- Asthma Immunomodulator PA Form
- Buprenorphine Products PA Form
- Buprenorphine Products Informed Consent
- Growth Hormone PA Form
- HAE Treatments PA Form
- Hepatitis C PA Form

For all other class medically-necessary coverage, quantity, and high dose requests use the following:

Documentation of Medical Necessity PA Form

#### with Prior Authorization Criteria

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https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf

ACNE AGENTS, TOPICAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benzoyl peroxide (BPO) WASH, LOTION clindamycin/BPO (generic Duac) clindamycin phosphate PLEDGET clindamycin phosphate SOLUTION DIFFERIN LOTION, CREAM, Rx-GEL (adapalene) DIFFERIN GEL (adapalene) OTC erythromycin GEL erythromycin-BPO (generic for Benzamycin) RETIN-A (tretinoin) <sup>AL</sup> CREAM, GEL	adapalene (generic differin) adapalene/BPO (generic Epiduo) adapalene/BPO (generic Epiduo Forte) AKLIEF (trifarotene) AL ALTRENO (tretinoin) AL AMZEEQ (minocycline) ARAZLO (tazarotene) ARAZLO (tazarotene) ARAZLO (tazarotene) ARAZLO (tazarotene) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) AZELEX (azelaic acid) BENZACLIN PUMP	Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

#### with Prior Authorization Criteria

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#### **ALZHEIMER'S AGENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHOLINESTERA donepezil (generic for Aricept)	ASE INHIBITORS donepezil 23 (generic for Aricept 23)	<ul> <li>Non-preferred agents will be approved for patients who have failed a 120-day trial of ONE</li> </ul>
donepezil ODT (generic for Aricept ODT)	galantamine (generic for Razadyne) SOLUTION, TABLET	preferred agent within this drug class within the last 6 months
EXELON Transdermal (rivastigmine)	galantamine ER (generic for Razadyne ER) rivastigmine (generic for Exelon)	<ul> <li>OR</li> <li>Current, stabilized therapy of the non-preferred agent within the previous 45 days</li> </ul>
NMDA RECEPTO	OR ANTAGONIST	Drug-specific criteria:
memantine (generic for Namenda)	memantine ER (generic for Namenda XR) memantine <b>SOLUTION</b> (generic for Namenda) NAMENDA (memantine) NAMZARIC (memantine/donepezil)	Donepezil 23: Requires donepezil 10mg/day for at least 3 months AND clinical reason as to why 5mg or 10mg tablets can't be used (to deliver 20mg or 25mg)

#### with Prior Authorization Criteria

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#### **ANALGESICS, OPIOID LONG-ACTING**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) <sup>QL</sup> PATCH fentanyl 25, 50, 75, 100 mcg PATCH <sup>QL</sup> morphine ER TABLET (generic MS Contin, Oramorph SR) OXYCONTIN <sup>CL</sup> (oxycodone ER) tramadol ER (generic Ultram ER) <sup>CL</sup>	ARYMO ER (morphine sulfate) <sup>QL</sup> BELBUCA (buprenorphine) <sup>QL</sup> BUCCAL  buprenorphine BUCCAL (generic for Belbuca) <sup>AL,NR,QL</sup> buprenorphine PATCH (generic Butrans) <sup>QL</sup> EMBEDA (morphine sulfate/naltrexone)  DURAGESIC MATRIX (fentanyl) <sup>QL</sup> fentanyl 37.5, 62.5, 87.5 mcg PATCH <sup>QL</sup> hydrocodone ER (generic for Hysingla ER) NR, QL  hydrocodone bitartrate ER (generic for Zohydro ER)  hydromorphone ER (generic for Exalgo) <sup>CL</sup> HYSINGLA ER (hydrocodone ER)  KADIAN (morphine ER)  methadone TABLET <sup>CL</sup> methadone ORAL SYR <sup>CL,NR</sup> MORPHABOND ER (morphine sulfate)  morphine ER (generic for Avinza, Kadian) CAPSULE  NUCYNTA ER (tapentadol) <sup>CL</sup> oxycodone ER (generic Oxycontin)  oxymorphone ER (generic Opana ER)  tramadol ER (generic Conzip) <sup>CL</sup>	The Center for Disease Control (CDC) does not recommend long acting opioids when beginning opioid treatment.  • Preferred agents require previous use of a long acting opioid or documentation of a trial on a short acting agent within 90 days  • Non-preferred agents will be approved with failure on, or intolerance to TWO preferred agents within this drug class  Drug-specific criteria:  • Methadone: Will only be approved for use in pain control or end of life care. Trial of preferred agent not required for end of life care  • Oxycontin®: Pain contract required for maximum quantity authorization

#### with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting **ANALGESICS, OPIOID SHORT-ACTING**  $^{\mathrm{QL}}$ 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
TABLET codeine TABLET hydrocodone/APAP SOLUTION,     TABLET hydrocodone/ibuprofen hydromorphone TABLET morphine CONC SOLUTION,     SOLUTION, TABLET oxycodone TABLET, SOLUTION oxycodone/APAP Tramadol 50 TABLETAL (generic     Ultram) tramadol/APAP (generic Ultracet)	APADAZ (benzhydrocodone/APAP) <sup>CL</sup> benzhydrocodone/APAP (generic Apadaz <sup>-CL</sup> butalbital/caffeine/APAP/codeine butalbital compound w/codeine   (butalbital/ASA/caffeine/codeine) carisoprodol compound-codeine   (carisoprodol/ASA/codeine) dihydrocodeine/APAP/caffeine dihydrocodeine/APAP/caffeine dihydrocodeine/aspirin/caffeine FIORINAL/CODEINE (butalbital/	<ul> <li>Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the last 12 months</li> <li>Note: for short acting opiate tablets and capsules there is a maximum quantity limit of #150 per 30 days.</li> <li>Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will consist of -prescriptions limited to a 7 day supply, AND -initial opiate prescription fill limited to maximum of 50 Morphine Milligram Equivalents (MME) per day</li></ul>

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# ANALGESICS, OPIOID SHORT-ACTINGQL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NA	NASAL	
	butorphanol <b>SPRAY</b> <sup>QL</sup> LAZANDA (fentanyl citrate)	
BUCCAL/TRANSMUCOSAL <sup>CL</sup>		Drug-specific criteria: _• Abstral®/Actiq®/Fentora®/
	ABSTRAL (fentanyl) <sup>CL</sup> fentanyl <b>TRANSMUCOSAL</b> (generic Actiq) <sup>CL</sup> FENTORA (fentanyl) <sup>CL</sup>	Onsolis (fentanyl): Approved only for diagnosis of cancer AND current use of long-acting opiate

#### ANDROGENIC AGENTS (Topical)CL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NDROGEL (testosterone) <b>PUMP</b> <sup>CL</sup>	ANDRODERM (testosterone) <sup>CL</sup> NATESTO (testosterone) <sup>CL</sup> testosterone PACKET (generic Androgel) <sup>CL</sup> testosterone PUMP (generic Androgel) <sup>CL</sup> testosterone GEL, PACKET, PUMP (generic Vogelxo) testosterone (generic Axiron) testosterone (generic Fortesta) testosterone (generic Testim)	<ul> <li>Preferred agents approved for diagnosis of Primary hypogonadism (congenital or acquired) or Hypogonadotropic hypogonadism. Off label use for the following will be considered with documentation of necessity: female to male transsexual – gender dysphoria, weight gain, male osteoporosis, delayed puberty in males, corticosteroid-induced hypogonadism and osteoporosis, inoperable carcinoma of the breast, postpartum breast pain and engorgement, and menopause</li> <li>In addition, non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the la 6 months</li> <li>Drug-specific criteria:         <ul> <li>Androderm®/Androgel®: Approved for Males only</li> <li>Natesto®: Approved for Males on with diagnosis of: Primary hypogonadism (congenital or acquired)</li> </ul> </li> </ul>

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#### **ANGIOTENSIN MODULATORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benazepril (generic Lotensin) enalapril (generic Vasotec) fosinopril (generic Monopril) lisinopril (generic Prinivil, Zestril) quinapril (generic Accupril) ramipril (generic Altace)	captopril (generic Capoten) EPANED (enalapril) <sup>CL</sup> ORAL SOLUTION enalapril (generic for Epaned) <sup>CL</sup> ORAL SOLUTION moexepril (generic Univasc) perindopril (generic Aceon) QBRELIS (lisinopril) <sup>CL</sup> ORAL SOLUTION trandolapril (generic Mavik)  JRETIC COMBINATIONS captopril/HCTZ (generic Capozide) moexipril/HCTZ (generic Uniretic)	<ul> <li>Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 12 months</li> <li>Non-preferred combination products may be covered as individual prescriptions without prior authorization</li> <li>Drug-specific criteria:         <ul> <li>Epaned® and Qbrelis® Oral Solution: Clinical reason why oral tablet is not appropriate</li> </ul> </li> </ul>
ANGIOTENSIN RI	ECEPTOR BLOCKERS	
irbesartan (generic Avapro) losartan (generic Cozaar) olmesartan (generic Benicar) valsartan (generic Diovan)	candesartan (generic Atacand) EDARBI (azilsartan) eprosartan (generic Teveten) telmisartan (generic Micardis)	

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#### **ANGIOTENSIN MODULATORS (Continued)**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANGIOTENSIN RECEPTOR BLO	ANGIOTENSIN RECEPTOR BLOCKER/DIURETIC COMBINATIONS	
irbesartan/HCTZ (generic Avalide) losartan/HCTZ (generic Hyzaar) olmesartan/HCTZ (generic Benicar- HCT) valsartan/HCTZ (generic Diovan-HCT)	candesartan/HCTZ (generic Atacand- HCT) EDARBYCLOR (azilsartan/ chlorthalidone) telmisartan/HCTZ (generic Micardis- HCT)	<ul> <li>approved for patients who have failed TWO preferred agents within this drug class within the last 12 months</li> <li>Non-preferred combination products may be covered as individual prescriptions without prior authorization</li> </ul>
	MODULATOR/ OCKER COMBINATIONS	<ul> <li>Angiotensin Modulator/Calcium</li> <li>Channel Blocker Combinations:</li> <li>Combination agents may be</li> </ul>
amlodipine/benazepril (generic Lotrel) amlodipine/olmesartan (generic Azor) amlodipine/valsartan (generic Exforge)	amlodipine/olmesartan/HCTZ (generic Tribenzor) amlodipine/telmisartan (generic Twynsta) amlodipine/valsartan/HCTZ (generic Exforge HCT) PRESTALIA (perindopril/amlodipine) trandolapril/verapamil (generic Tarka)	approved if there has been a trial and failure of preferred agent
DIRECT DENI	N INHIBITORS	<ul> <li>Direct Renin Inhibitors/Direct Renin Inhibitor Combinations:</li> </ul>
DIRECT RENI	aliskiren (generic Tekturna) <sup>QL</sup>	<ul> <li>May be approved with history of TWO preferred ACE Inhibitors or Angiotensin Receptor Blockers</li> </ul>
DIRECT RENIN INHIB	ITOR COMBINATIONS	within the last 12 months
	TEKTURNA/HCT (aliskiren/HCTZ)	Drug Specific Criteria
NEPRILYSIN INHIBITOR COMBINATION		Entresto: May be approved with a diagnosis of heart failure
ENTRESTO (sacubitril/valsartan) <sup>AL,QL</sup>		AND ≥ 18 years old
ANGIOTENSIN RECEPTOR BLOCKE	ER/BETA-BLOCKER COMBINATIONS	
	BYVALSON (nevibolol/valsartan)	

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#### **ANTHELMINTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
albendazole (generic for Albenza) BILTRICIDE (praziquantel) ivermectin (generic for Stromectol)	ALBENZA (albendazole) EMVERM (mebendazole) <sup>CL</sup> praziquantel (generic for Biltricide) STROMECTOL (ivermectin)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> <li>Drug-specific criteria:</li> <li>Emverm: Approval will be considered for indications not covered by preferred agents</li> </ul>

#### ANTI-ALLERGENS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ORALAIR (sweet vernal/orchard/rye/timothy/kentucky blue grass mixed pollen allergen extract)  PALFORZIA AL.CL (peanut allergen powder-dnfp)	ORALAIR  Confirmed by positive skin test or in vitro testing for pollenspecific IgE antibodies for Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass Mixed Pollens.  For use in patients 10 through 65 years of age.  PALFORZIA  Confirmed diagnosis of peanuallergy by allergist  For use in patients ages 4 to 17; it may be continued in patients 18 years and older with documentation of previous use within the past 90 days  Initial dose and increase titration doses should be given in a healthcare setting  Should not be used in patients with uncontrolled asthma or concurrently on a NSAID

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#### ANTIBIOTICS, GASTROINTESTINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FIRVANQ (vancomycin) SOLUTION netronidazole TABLET neomycin inidazole (generic Tindamax) <sup>CL</sup>	DIFICID (fidaxomicin) CL TABLET, SUSPNR FLAGYL ER (metronidazole)CL MetronidazoleCL CAPSULE nitazoxanide (generic Alinia) TABLETAL, CL, QL paromomycin SOLOSEC (secnidazole) vancomycin CAPSULE (generic Vancocin)CL XIFAXAN (rifaximin)CL	<ul> <li>Note: Although azithromycin, ciprofloxacin and trimethoprim/ sulfmethoxazole are not included in this review, they are available without prior authorization</li> <li>Drug-specific criteria:</li> <li>Alinia®: Trial and failure with metronidazo is required for a diagnosis of giardiasis</li> <li>Dificid®: Trial and failure with oral vancomycin is required for a diagnosis of difficile diarrhea (pseudomembranous colitis)</li> <li>Flagyl ER®: Trial and failure with metronidazole is required</li> <li>Flagyl®/Metronidazole 375mg capsules and Flagyl ER®/ Metronidazole 750mg Etabs: Clinical reason why the generic regular-release cannot be used</li> <li>tinidazole:         Approvable diagnoses include: Giardia         Amebiasis intestinal or liver abscess         Bacterial vaginosis or trichomoniasis     </li> <li>vancomycin capsules: Requires patient specific documentation of why the Firvanq/vancomycin solution is not appropriate for patient</li> <li>Xifaxan®: Approvable diagnoses include: Travelers's diarrhea resistant to quinolone Hepatic encephalopathy with treatment failure of lactulose or neomycin Diarrhea-Predominant IBS (IBS-D) 550mg strength only with treatment failure of Lomotil® AND Imodium®</li> </ul>

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#### **ANTIBIOTICS, INHALED**

Preferred Agents <sup>CL</sup>	Non-Preferred Agents	Prior Authorization/Class Criteria
BETHKIS (tobramycin) <sup>CL</sup> KITABIS PAK (tobramycin) <sup>CL</sup> TOBI-PODHALER (tobramycin) <sup>CL,QL</sup>	ARIKAYCE (amikacin liposomal inh) <sup>CL</sup> SUSPENSION CAYSTON (aztreonam lysine) <sup>QL,CL</sup> tobramycin (generic for Bethkis) tobramycin (generic Tobi) <sup>CL</sup>	

#### **ANTIBIOTICS, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
bacitracin <b>OINTMENT</b> bacitracin/polymyxin (generic Polysporin) mupirocin <b>OINTMENT</b> (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/ pramoxine	CENTANY (mupirocin) gentamicin OINTMENT, CREAM mupirocin CREAM (generic Bactroban) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class within the last 12 months</li> <li>Drug-specific criteria:</li> <li>Mupirocin® Cream: Clinical reason the ointment cannot be used</li> </ul>

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### ANTIBIOTICS, VAGINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CLEOCIN <b>OVULES</b> (clindamycin) clindamycin <b>CREAM</b> (generic Cleocin) CLINDESSE (clindamycin) NUVESSA (metronidazole) VANDAZOLE (metronidazole)	CLEOCIN <b>CREAM</b> (clindamycin) METROGEL (metronidazole) metronidazole, vaginal	<ul> <li>Non-preferred agents will be approved for patients who have failed a therapeutic trial (duration = 3 days) with ONE preferred agent within this drug class within the last 6 months</li> </ul>

#### **ANTICOAGULANTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg <sup>CL,QL</sup> XARELTO DOSE PACK (rivaroxaban)	BEVYXXA (betrixaban) <sup>QL</sup> fondaparinux (generic Arixtra) FRAGMIN (dalteparin) SAVAYSA (edoxaban) <sup>QL</sup> XARELTO (rivaroxaban) <sup>NR</sup> SUSP	<ul> <li>Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 12 months</li> <li>Drug-specific criteria:</li> <li>Coumadin®: Clinical reason generic warfarin cannot be used</li> <li>Savaysa®: Approved diagnoses include:         Stroke and systemic embolism (SE) risk reduction in nonvalvular atrial fibrillation (NVAF) OR</li></ul>

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#### **ANTIEMETICS/ANTIVERTIGO AGENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
dronabinol (generic Marinol) <sup>AL</sup>	CESAMET (nabilone)	<ul> <li>Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the same</li> </ul>
5HT3 RECEPTO	OR BLOCKERS	group
ondansetron (generic Zofran/Zofran ODT) <sup>QL</sup>	ANZEMET (dolasetron) granisetron (generic Kytril) SANCUSO (granisetron) <sup>CL</sup> ZUPLENZ (ondansetron)	Drug-specific criteria:  • Akynzeo®/Varubi®: Approved for Moderately/Highly emetogenic chemotherapy with dexamethasone and a 5-HT3 antagonist
NK-1 RECEPTO	R ANTAGONIST	Regimens include: AC combination     (Doxorubicin or Epirubicin with
EMEND (aprepitant) CAPSULE, CAPSULE PACKQL	aprepitant (generic Emend) QL,CL AKYNZEO (netupitant/palonosetron)CL VARUBI (rolapitant) <b>TABLET</b> CL	Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine, Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine, Dacarbazine, Dactinomycin, Daunorubicin,
TRADITIONAL	ANTIEMETICS	Epirubicin, Etoposide,
DICLEGIS (doxylamine/pyridoxine) <sup>CL,QL</sup> dimenhydrinate (generic Dramamine) OTC meclizine (generic Antivert) metoclopramide (generic Reglan) phosphoric acid/dextrose/fructose	BONJESTA (doxylamine/pyridoxine)·CL,QL COMPRO (prochlorperazine) doxylamine/pyridoxine (generic Diclegis)CL,QL metoclopramide ODT (generic Metozolv ODT) prochlorperazine SUPPOSITORY (generic Compazine) promethazine SUPPOSITORY 50mg scopolamine TRANSDERMAL trimethobenzamide TABLET (generic Tigan)	<ul> <li>Hexamethylmelamine, Idarubicin, Ifosfamide, Imatinib, Interferon α, Irinotecan, Mechlorethamine, Melphalan, Methotrexate, Oxaliplatin, Procarbazine, Streptozotocin, Temozolomide</li> <li>Diclegis®/Bonjesta: Approved only for treatment of nausea and vomiting of pregnancy</li> <li>Metozolv ODT®: Documentation of inability to swallow or Clinical reason oral liquid cannot be used</li> <li>Sancuso®/Zuplenz®: Documentation of oral dosage form intolerance</li> </ul>

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#### ANTIFUNGALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche) fluconazole SUSPENSION, TABLET (generic Diflucan) griseofulvin SUSPENSION griseofulvin microsized TABLET nystatin SUSPENSION, TABLET terbinafine (generic Lamisil)	BREXAFEMME (ibrexafungerp) <sup>QL,NR</sup> CRESEMBA (isavuconazonium) <sup>CL</sup> flucytosine (generic Ancobon) <sup>CL</sup> griseofulvin ultramicrosize (generic GRIS-PEG) itraconazole (generic Sporanox) <sup>CL</sup> ketoconazole (generic Nizoral) nystatin <b>POWDER</b> ONMEL (itraconazole) posaconazole (generic Noxafil) <sup>AL,CL</sup> TOLSURA (itraconazole) <sup>CL</sup> voriconazole (generic VFEND) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class</li> <li>Drug-specific criteria:</li> <li>Cresemba®: Approved for diagnosis of invasive aspergillosis or invasive mucomycosis</li> <li>Flucytosine: Approved for diagnosis of:         <ul> <li>Candida: Septicemia, endocarditis, UTIs</li> <li>Cryptococcus: Meningitis, pulmonary infections</li> </ul> </li> <li>Noxafil®: No trial for diagnosis of Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropenic hematologic malignancies, Graft vs. Host disease(GVHD), Immunosuppression secondary to hematopoietic stem cell transplant</li> <li>Noxafil® Suspension:</li></ul>

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#### **ANTIFUNGALS, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
· · · · · · · · · · · · · · · · · · ·	ALEVAZOL (clotrimazole) OTC ciclopirox CREAM, GEL, SUSPENSION   (generic Ciclodan, Loprox) ciclopirox NAIL LACQUER (generic   Penlac) ciclopirox SHAMPOO (generic Loprox) clotrimazole SOLUTION RX (generic   Lotrimin) DESENEX POWDER OTC (miconazole) econazole (generic Spectazole) ERTACZO (sertaconazole) EXELDERM (sulconazole) FUNGOID OTC JUBLIA (efinaconazole) ketoconazole FOAM (generic Extina,   Ketodan) LAMISIL AT GEL, SPRAY (terbinafine)   OTC LOPROX (ciclopirox) SUSPENSION,   SHAMPOO, CREAM LOTRIMIN AF CREAM OTC   (clotrimazole) LOTRIMIN ULTRA (butenafine) luliconazole (generic Luzu) MENTAX (butenafine) miconazole OTC OINTMENT, SPRAY miconazole/zinc oxide/petrolatum (generic   Vusion) naftifine CREAM, GEL (generic Naftin) oxiconazole (generic Bensal HP) tavaborole SOLUTION (generic   Kerydin) <sup>NR</sup> tolnaftate SPRAY, OTC	Prior Authorization/Class Criteria  Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months  Drug-specific criteria: Extina: Requires trial and failure or contraindication to other ketoconazole forms Jublia: Approved diagnoses includ Onychomycosis of the toenails due to T.rubrum OR T. Mentagrophytes ciclopirox nail lacquer: No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine
	COID COMBINATIONS clotrimazole/betamethasone LOTION (generic Lotrisone)	

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#### with Prior Authorization Criteria

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#### **ANTIHISTAMINES, MINIMALLY SEDATING**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
cetirizine TABLET, SOLUTION (Rx only) (generic for Zyrtec) loratadine TABLET, SOLUTION (generic for Claritin) levocetirizine TABLET (generic for Xyzal)	cetirizine CHEWABLE (generic for Zyrtec) cetirizine SOLUTION (OTC) desloratadine (generic for Clarinex) desloratadine ODT (generic for Clarinex Reditabs) fexofenadine (generic for Allegra) fexofenadine 180mg (generic for Allegra 180mg) <sup>QL</sup> levocetirizine (generic for Xyzal) SOLUTION loratadine CAPSULE, CHEWABLE, ODT (generic for Claritin Reditabs)	<ul> <li>Non-preferred agents will be approved for patients who have failed TWO preferred agents within this drug class</li> <li>Combination products not covered – individual products may be covered</li> </ul>

#### **ANTIHYPERTENSIVES. SYMPATHOLYTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CATAPRES-TTS (clonidine) clonidine <b>TABLET</b> (generic for Catapres) guanfacine (generic for Tenex) methyldopa	clonidine <b>TRANSDERMAL</b> methyldopa/hydrochlorothiazide	Non-preferred agents will be approved for patients who have failed a 30-day trial with ONE preferred agent within this drug class

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#### **ANTIHYPERURICEMICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
allopurinol (generic for Zyloprim) MITIGARE (colchicine) probenecid probenecid/colchicine (generic for Col- Probenecid)	colchicine <b>TABLET</b> (generic for Colcrys) <sup>CL</sup> colchicine <b>CAPSULE</b> (generic for Mitigare) febuxostat (generic for Uloric) <sup>CL</sup> <i>GLOPERBA</i> <b>SOLN</b> (colchicine) <sup>CL,QL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> <li>colchicine tablet®: Approved without trial for familial Mediterranean fever OR pericarditis</li> <li>Gloperba: Approved for documented swallowing disorder</li> <li>Uloric®: Clinical reason why allopurinol cannot be used</li> </ul>

#### with Prior Authorization Criteria

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#### ANTIMIGRAINE AGENTS. OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
JOVY (fremanezumab-vfrm) CL, QL PEN, Autoinjector, Autoinjector 3-packNR  JOVY (fremanezumab-vfrm) Autoinjector 3-packCL,NR,QL  MGALITY 120 mg/mL (galcanezumab-gnlm) CL, QL PEN, SYRINGE  BRELVY (ubrogepant)AL,CL, QL  TABLET	AIMOVIG (erenumab-aooe) CL,QL CAFERGOT (ergotamine/caffeine) CAMBIA (diclofenac potassium) dihydroergotamine mesylate NASAL ELYXYB (celecoxib)AL,NR,QL SOLN EMGALITY 100 mg (galcanezumabgnlm) CL,QL SYRINGE ERGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL NURTEC ODT (rimegepant)AL,CL,QL QULIPTA (atogepant)AL,NR,QL REYVOW (lasmiditan)AL, CL,QL TABLET TRUDHESA (dihydroergotamine mesylate)AL,NR,QL NASAL	<ul> <li>All acute treatment agents will be approved for patients who have a failed trial or contraindication of a triptan.</li> <li>In addition, all non-preferred agents will require a failed trial or contraindication of a preferred agent of the same indication</li> <li>Drug-specific criteria:</li> <li>Cambia®: Requires diagnosis of migraine and documentation of why solid dosing forms not appropriate</li> <li>Emgality 120mg is recommended dosing for Migraine, Emgaility 100mg is recommended dosing for Episodic Cluster Headache</li> <li>Aimovig, Ajovy and Emgality 120mg: Require ≥ 4 migraines per month for ≥ 3 months and has tried and failed a ≥ 1 month trial of two medications listed in the 2012 American Academy of Neurology/American Headache Society guidelines (examples include: antidepressants (amitriptyline, venlafaxine), Beta blockers (propranolol, metroprolol, timolol, atenolol), anti-epileptics (valproate, topiramate), ACE/ARB (lisinopril, candesartan)</li> <li>In addition, Aimovig requires a trial of Emgality 120mg or Ajovy or clinical, patient specific reason that a preferred agent</li> </ul>

#### with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting ANTIMIGRAINE AGENTS, TRIPTANSQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORAL		Non-preferred agents will be
rizatriptan (generic Maxalt) rizatriptan ODT (generic Maxalt MLT) sumatriptan  NA IMITREX (sumatriptan)	almotriptan (generic Axert) eletriptan (generic Relpax) frovatriptan (generic Frova) IMITREX (sumatriptan) naratriptan (generic Amerge) RELPAX (eletriptan) <sup>QL</sup> sumatriptan/naproxen (generic Treximet) zolmitriptan (generic Zomig/Zomig ZMT)  SSAL  ONZETRA XSAIL (sumatriptan) sumatriptan (generic Imitrex Nasal) TOSYMRA (sumatriptan) zolmitriptan (generic for Zomig) ZOMIG (zolmitriptan)	approved for patients who have failed ALL preferred agents within this drug class  Drug-specific criteria:  • Sumavel® Dosepro: Requires clinical reason sumatriptan injection cannot be used  • Onzetra, Zembrace: approved for patients who have failed ALL preferred agents
INJE	CTABLE	
sumatriptan KIT, SYRINGE, VIAL	IMITREX (sumatriptan) <b>INJECTION</b> SUMAVEL DOSEPRO (sumatriptan) ZEMBRACE SYMTOUCH (sumatriptan)	

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# ANTIPARASITICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NATROBA (spinosad) permethrin 1% OTC (generic Nix) permethrin 5% RX (generic Elimite) pyrethrin/piperonyl butoxide (generic RID, A-200)	CROTAN (crotamiton) LOTION EURAX (crotamiton) CREAM, LOTION ivermectin (generic Sklice) LOTION NR lindane malathion (generic Ovide) spinosad (generic Natroba) VANALICE (piperonyl butoxide/pyrethrins)	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

#### with Prior Authorization Criteria

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#### ANTIPARKINSON'S AGENTS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHO	LINERGICS	Non-preferred agents will be
benztropine (generic for Cogentin) trihexyphenidyl (generic for Artane)	HIBITORS	approved for patients who have failed ONE preferred agents within this drug class
COMITIN		
	entacapone (generic for Comtan) tolcapone (generic for Tasmar)	<ul> <li>Drug-specific criteria:</li> <li>Carbidopa/Levodopa ODT: Approved for documented swallowing disorder</li> <li>COMT Inhibitors: Approved if using</li> </ul>
DOPAMINE	AGONISTS	as add-on therapy with levodopa-
pramipexole (generic for Mirapex) ropinirole (generic for Requip)	bromocriptine (generic for Parlodel) ropinirole ER (generic for Requip ER) <sup>CL</sup> NEUPRO (rotigotine) <sup>CL</sup> pramipexole ER (generic for Mirapex ER) <sup>CL</sup> ropinirole ER (generic for Requip XL) <sup>CL</sup>	<ul> <li>containing drug</li> <li>Gocovri: Required diagnosis of Parkinson's disease and had trial of or is intolerant to amantadine AND must be used as an add-on therapy with levodopa-containing drug</li> <li>Inbrija: Approval upon diagnosis of Parkinson's disease and concurrent treatment with carbidopa/levodopa agent</li> <li>Neupro®:</li> </ul>
MAO-B IN	IHIBITORS	For Parkinsons: Clinical reason required why preferred agent
selegiline CAPSULE, TABLET (generic for Eldepryl)	rasagiline (generic for Azilect) <sup>QL</sup> XADAGO (safinamide) ZELAPAR (selegiline) <sup>CL</sup>	cannot be used  For Restless Leg (RLS): Requires trial OR Contraindication to ropinirole AND pramipexole  Nourianz: Approval upon diagnosis of
OTHER ANTIPAR	KINSON'S DRUGS	Parkinson's disease and concurrent
amantadine CAPSULE, SYRUP TABLET (generic for Symmetrel) carbidopa/levodopa (generic for Sinemet) carbidopa/levodopa ER (generic for Sinemet CR) levodopa/carbidopa/entacapone (generic for Stalevo)	APOKYN (apomorphine) SUB-Q apomorphine (generic for Apokyn) <sup>NR</sup> SUB-Q carbidopa (generic for Lodosyn) carbidopa/levodopa ODT (generic for Parcopa) DHIVY (carbidopa/levodopa) NR,QL DUOPA (carbidopa/levodopa) GOCOVRI (amantadine) <sup>QL</sup> INBRIJA (levodopa) INHALER <sup>CL,QL</sup> KYNMOBI (apomorphine) <sup>QL,</sup> KIT, SUBLINGUAL NOURIANZ (istradefylline) <sup>CL,QL</sup> OSMOLEX ER (amantadine) <sup>QL</sup> RYTARY (carbidopa/levodopa) STALEVO (ledopa/carbidopa/entacapone)	treatment with carbidopa/levodopa agent  Osmolex ER: Required diagnosis of Parkinson's disease or drug-induced extrapyramidal reactions and had trial of or is intolerant to amantadine IR  Pramipexole ER: Required diagnosis of Parkinson's along with preferred agent trial  Ropinerole ER: Required diagnosis of Parkinson's along with preferred agent trial  Zelapar®: Approved for documented swallowing disorder

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#### **ANTIPSORIATICS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acitretin (generic for Soriatane)	methoxsalen (generic for Oxsoralen- Ultra) SORIATANE (acitretin)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial with THE preferred agent within this drug class</li> <li>Trial of acitretin (Pregnancy category X) not required in pregnancy or while attempting or planning pregnancy</li> </ul>

#### ANTIPSORIATICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
alcipotriene CREAM, OINTMENT, SOLUTION,	calcitriol (generic for Vectical) calcipotriene/betamethasone     OINTMENT(generic for Taclonex) calcipotriene/betamethasone SUSP     (generic for Taclonex Scalp) CALCITRENE (calcipotriene) DOVONEX CREAM (calcipotriene) DUOBRII     (halobetasol prop/tazarotene ENSTILAR     (calcipotriene/betamethasone) SORILUX (calcipotriene)	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

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#### **ANTIVIRALS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acyclovir (generic Zovirax) famciclovir (generic Famvir) valacyclovir (generic Valtrex)	acyclovir (generic for Zovirax) <sup>CL</sup> SUSPENSION SITAVIG (acyclovir buccal) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a 10-day trial of ONE preferred agent within the same group</li> </ul>
ANTI-INFLUE oseltamivir (generic Tamiflu) <sup>QL</sup>	rimantadine (generic Flumadine) RELENZA (zanamivir) <sup>QL</sup> TAMIFLU (oseltamivir) <sup>QL</sup> XOFLUZA (baloxavir marboxil) <sup>AL,CL,QL</sup>	<ul> <li>Acyclovir Susp: Prior authorization NOT required for children ≤ 12 years old</li> <li>Sitavig®: Approved for recurrent herpes labialis (cold sores) in immunocompetent adults</li> <li>Xofluza: Requires clinical, patient specific reason that a preferred agent cannot be used</li> </ul>

#### **ANTIVIRALS, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acyclovir <b>OINTMENT</b>	acyclovir CREAM, (generic Zovirax) DENAVIR (penciclovir) XERESE (acyclovir/hydrocortisone)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred ORAL Antiviral agent</li> </ul>

#### ANXIOLYTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
alprazolam <b>TABLET</b> (generic for Xanax) buspirone (generic for Buspar) chlordiazepoxide diazepam <b>TABLET</b> , <b>SOLUTION</b> (generic for Valium) lorazepam <b>INTENSOL</b> , <b>TABLET</b> (generic for Ativan)	alprazolam ER (generic for Xanax XR) alprazolam ODT alprazolam INTENSOL <sup>CL</sup> clorazepate (generic for Tranxene-T) diazepam INTENSOL <sup>CL</sup> lorazepam ORAL SYRINGE <sup>NR</sup> LOREEV XR (lorazepam) <sup>AL.NR</sup> meprobamate oxazepam	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial with TWO preferred agents within this drug class</li> <li>Drug-specific criteria:</li> <li>Diazepam Intensol®: Requires clinical reason why diazepam solution cannot be used</li> <li>Alprazolam Intensol®: Requires trial of diazepam solution OR lorazepam Intensol®</li> </ul>

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#### **BETA BLOCKERS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
•	acebutolol (generic Sectral) betaxolol (generic Kerlone) BYSTOLIC (nebivolol) HEMANGEOL (propranolol) SOLUTION INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) LEVATOL (penbutolol) metoprolol/HCTZ (generic Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide nebivolol (generic Bystolic) <sup>NR</sup> pindolol (generic Viskin) propranolol/HCTZ (generic Inderide) timolol (generic Blocadren) TOPROL XL (metoprolol ER)	<ul> <li>Non-preferred agents will be approved for patients who have failed TWO diagnosis-appropriate preferred agents within this drug class</li> <li>Drug-specific criteria:         <ul> <li>Bystolic®: Only ONE trial is required with Diagnosis of Obstructive Lung Disease</li> <li>Coreg CR®: Requires clinical reason generic IR product cannot be used</li> <li>Hemangeol®: Covered for diagnosis of Proliferating Infantile Hemangioma</li> <li>Sotylize®: Covered for diagnosis of life –threatening ventricular arrhythmias OR maintenance of normal sinus rhythm in highly symptomatic atrial fibrillation/flutter (AFIB/AFL)</li> <li>Requires clinical reason generic sotalol cannot be used</li> </ul> </li> </ul>
BETA- AND ALF	PHA-BLOCKERS	_
carvedilol (generic Coreg) labetalol (generic Trandate)	carvedilol ER <sup>CL</sup> (generic Coreg CR)	
ANTIARR	HYTHMIC	
sotalol (generic Betapace)	SOTYLIZE (sotalol)	

#### **BILE SALTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ursodiol <b>CAPSULE</b> 300mg (generic for Actigall) ursodiol 250mg <b>TABLET</b> (generic for URSO) ursodiol 500mg <b>TABLET</b> (generic for URSO FORTE)	BYLVAY (odevixibat) <sup>NR</sup> CAP, PELLET CHENODAL (chenodiol) CHOLBAM (cholic acid) LIVMARLI (maralixibat) SOLN <sup>AL,NR</sup> OCALIVA (obeticholic acid) RELTONE (ursodiol 200mg,400mg) CAP <sup>NR</sup>	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

#### with Prior Authorization Criteria

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#### **BLADDER RELAXANT PREPARATIONS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Oxybutynin IR, ER (generic Ditropan/Ditropan XL) solifenacin (generic Vesicare) TOVIAZ (fesoterodine ER)	darifenacin ER (generic Enablex) GELNIQUE (oxybutynin) GEMTESA (vibegron)AL,NR,QL flavoxate MYRBETRIQ <b>TAB</b> , <b>SUSP</b> AL,NR,QL (mirabegron) OXYTROL (oxybutynin) tolterodine IR, ER (generic Detrol/ Detrol LA) trospium IR, ER (generic Sanctura/ Sanctura XR) VESICARE (solifenacin) VESICARE LS <b>SUSP</b> (solifenacin succinate)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Myrbetriq®: Covered without trial in contraindication to anticholinergic agents</li> </ul>

#### with Prior Authorization Criteria

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#### **BONE RESORPTION SUPRESSION AND RELATED DRUGS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BISPHOSF	BISPHOSPHONATES	
alendronate (generic Fosamax) <b>TABLET</b> ibandronate (generic Boniva) <sup>QL</sup>	alendronate <b>SOLUTION</b> (generic Fosamax) <sup>QL</sup> ATELVIA DR (risedronate)	approved for patients who have failed a trial of ONE preferred agent within the same group
	BINOSTO (alendronate)	Drug-specific criteria:
	etidronate disodium (generic Didronel) FOSAMAX PLUS DQL	individual agents without prior authorization
	risedronate (generic Actonel) <sup>QL</sup>	<ul> <li>Atelvia DR®: Requires clinical reason alendronate cannot be taken on an empty stomach</li> </ul>
	PRESSION AND RELATED DRUGS	Binosto®: Requires clinical reason why alendronate tablets OR Fosamax® solution cannot be used
calcitonin-salmon <b>NASAL</b> raloxifene (generic Evista)	EVISTA (raloxifene) FORTEO (teriparatide) <sup>CL,QL</sup>	Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification
teriparatide (generic Forteo) CL,QL	TYMLOS (abaloparatide)	• Forteo®: Covered for high risk of fracture
		High risk of fracture:
		<ul><li>BMD -3 or worse</li><li>Postmenopausal women with history of</li></ul>
		non-traumatic fractures
		<ul> <li>Postmenopausal women with 2 or more clinical risk factors</li> </ul>
		<ul> <li>Family history of non-traumatic fractures</li> </ul>
		<ul> <li>DXA BMD T-score ≤ -2.5 at any site</li> </ul>
		<ul> <li>Glucocorticoid use ≥ 6 months at</li> <li>7.5 dose of prednisolone equivalent</li> </ul>
		o Rheumatoid Arthritis
		<ul> <li>Postmenopausal women with BMD T- score ≤ -2.5 at any site with any clinical risk factors</li> </ul>
		<ul> <li>More than 2 units of alcohol per day</li> </ul>
		o Current smoker
		<ul> <li>Men with primary or hypogonadal osteoporosis</li> </ul>
		<ul> <li>Osteoporosis associated with sustained systemic glucocorticoid therapy</li> </ul>
		Trial of calcitonin-salmon not required
		<ul> <li>Maximum of 24 months treatment per lifetime</li> </ul>

#### with Prior Authorization Criteria

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#### **BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALPHA B	LOCKERS	Non-preferred agents will be
alfuzosin (generic Uroxatral) doxazosin (generic Cardura) tamsulosin (generic Flomax)	CARDURA XL (doxazosin) silodosin (generic Rapaflo)	approved for patients who have failed a trial of ONE preferred agent within this drug class
terazosin (generic Hytrin)		Drug-specific criteria:
5-ALPHA-REDUCTAS	SE (5AR) INHIBITORS	Alfuzosin/dutasteride/finasteride
dutasteride (generic for Avodart) finasteride (generic for Proscar)	dutasteride/tamsulosin (generic for Jalyn)	<ul> <li>Covered for males only</li> <li>Cardura XL®: Requires clinical reason generic IR form cannot be used</li> <li>Flomax®: Females covered for a 7 day supply with diagnosis of acute kidney stones</li> <li>Jalyn®: Requires clinical reason why individual agents cannot be used</li> </ul>

#### with Prior Authorization Criteria

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#### **BRONCHODILATORS, BETA AGONIST**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROAIR HFA (albuterol) albuterol HFA (generic for ProAir HFA)	albuterol HFA (Proventil HFA, Ventolin HFA) levalbuterol HFA (generic for Xopenex HFA)  PROAIR DIGIHALER (albuterol) PROAIR RESPICLICK (albuterol) PROVENTIL HFA (albuterol)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Xopenex®: Covered for cardiac diagnoses or side effect of tachycardia with albuterol product</li> </ul>
SEREVENT (salmeterol)	ERS – Long Acting  ARCAPTA NEOHALER (indacaterol)  STRIVERDI RESPIMAT (olodaterol)	
albuterol (2.5mg/3ml premix or 2.5mg/0.5ml) albuterol 100 mg/20 mL albuterol low dose (0.63mg/3ml & 1.25mg/3ml)	ATION SOLUTION  arformoterol tartrate (generic Brovana)  BROVANA (arformoterol)  formoterol fumarate (generic Perforomist)  levalbuterol (generic for Xopenex)	
albuterol SYRUP	ORAL albuterol TABLET albuterol ER (generic for Vospire ER)	
	metaproterenol (formerly generic for Alupent) terbutaline (generic for Brethine)	

#### with Prior Authorization Criteria

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#### **CALCIUM CHANNEL BLOCKERS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SHORT-ACTING Dihydropyridines		Non-preferred agents will be approved for patients who have
Non-dihydodiltiazem (generic Cardizem) verapamil (generic Calan/Isoptin) LONG-	isradipine (generic Dynacirc) nicardipine (generic Cardene) nifedipine (generic Procardia) nimodipine (generic Nimotop) NYMALIZE (nimodipine) SOLUTION ropyridines  ACTING pyridines felodipine ER (generic Plendil)	failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:  Nifedipine: May be approved without trial for diagnosis of Preterm Labor or Pregnancy Induced Hypertension (PIH)  Nimodipine: Covered without trial for diagnosis of subarachnoid hemorrhage  Katerzia: May be approved with documented swallowing difficulty
nifedipine ER (generic Procardia XL/ Adalat CC)	KATERZIA (amlodipine) <sup>QL</sup> <b>SUSP</b> nisoldipine (generic Sular)	-
Non-dihyd	ropyridines	_
diltiazem ER (generic Cardizem CD) verapamil ER <b>TABLET</b>	CALAN SR (verapamil) diltiazem ER (generic Cardizem LA) MATZIM LA (diltiazem ER) TIAZAC (diltiazem) verapamil ER CAPSULE verapamil 360mg CAPSULE verapamil ER (generic Verelan PM)	

#### with Prior Authorization Criteria

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#### CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BETA LACTAM/BETA-LACTAMASE INHIBITOR COMBINATIONS		Non-preferred agents will be
amoxicillin/clavulanate TABLETS, SUSPENSION	amoxicillin/clavulanate CHEWABLE amoxicillin/clavulanate ER (generic Augmentin XR) AUGMENTIN (amoxicillin/clavulanate) SUSPENSION, TABLET	approved for patients who have failed a 3-day trial of ONE preferred agent within the same group
CEPHALOSPORINS	S – First Generation	
cefadroxil CAPSULE, SUSPENSION (generic Duricef) cephalexin CAPSULE, SUSPENSION (generic Keflex)	cefadroxil <b>TABLET</b> (generic Duricef) cephalexin <b>TABLET</b>	
CEPHALOSPORINS -	Second Generation	
cefprozil (generic Cefzil)	cefaclor (generic Ceclor)	
cefuroxime TABLET (generic Ceftin)	CEFTIN (cefuroxime) TABLET, SUSPENSION	
CEPHALOSPORINS -	- Third Generation	
cefdinir (generic Omnicef)	cefixime CAPSULE, SUSPENSION (generic Suprax) cefpodoxime (generic Vantin) SUPRAX CAPSULE, CHEWABLE TAB, SUSPENSION, TABLET (cefixime)	

#### **COLONY STIMULATING FACTORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NEUPOGEN (filgrastim) VIAL	GRANIX (tbo-filgrastim) NEUPOGEN <b>DISP SYR</b> (filgrastim) NIVESTYM <b>SYR,VIAL</b> (filgrastim-aafi) Nyvepria (pegfilgrastim-apgf) RELEUKO (filgrastim-ayow) <sup>NR</sup> <b>SYR,VIAL</b> ZARXIO (filgrastim-sndz) ZIEXTENZO <b>SYR</b> (pegfilgrastim-bmez)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

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#### **CONTRACEPTIVES, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
All reviewed agents are recommended preferred at this time  Only those products for review are listed.  Brand name products may be subject to Maximum Allowable Cost (MAC) pricing or require substitution with a generic equivalent	DOLISHALE (ethinyl estradiol/ levonorgestrel) <sup>NR</sup> NEXTSTELLIS(drospirenone/estetrol) <sup>NR</sup> TAYSOFY (norethindrone/ethinyl estradiol/iron) <sup>NR</sup> TYBLUME (levonorgestrel/ ethinyl estradiol) <sup>NR</sup>	
Specific agents can be looked up using the Drug Look-up Tool at:  https://druglookup.fhsc.com/drug lookupweb/?client=nestate		

#### with Prior Authorization Criteria

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#### COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANORO ELLIPTA (umeclidinium/vilanterol) ATROVENT HFA (ipratropium) COMBIVENT RESPIMAT (albuterol/ipratropium) SPIRIVA (tiotropium) STIOLTO RESPIMAT (tiotropium/olodaterol)	BEVESPI AEROSPHERE (glycopyrolate/formoterol) DUAKLIR PRESSAIR (aclidinium br and formoterol fum) INCRUSE ELIPTA (umeclidnium) SEEBRI NEOHALER (glycopyrolate) SPIRIVA RESPIMAT (tiotropium) TUDORZA PRESSAIR (aclidinium br) UTIBRON NEOHALER (indacaterol/glycopyrolate)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class OR Patient specific documentation of inability to use traditional inhaler device.</li> <li>Drug-specific criteria:         <ul> <li>Daliresp®:</li> <li>Covered for diagnosis of severe COPD associated with chronic bronchitis</li> <li>Requires trial of a bronchodilator Requires documentation of one</li> </ul> </li> </ul>
INHALATIO	N SOLUTION	<ul> <li>exacerbation in last year upon initial review</li> </ul>
albuterol/ipratropium (generic for Duoneb) ipratropium <b>SOLUTION</b> (generic for Atrovent)	LONHALA (glycopyrrolate inhalation soln) YUPELRI (revefenacin)	
ORAL	AGENT	
	DALIRESP (roflumilast) <sup>CL, QL</sup>	

#### **COUGH AND COLD, OPIATE COMBINATION**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	guaifenesin/codeine LIQUID hydrocodone/homatropine SYRUP promethazine/codeine SYRUP promethazine/phenylephrine/codeine SYRUP pseudoephedrine/codeine/ guaifenesin (generic for Lortuss EX, Tusnel C, Virtussin DAC)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE dextromethorphan product</li> <li>All codeine or hydrocodone containing cough and cold combinations are limited to ≥ 18 years of age</li> </ul>

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#### CYSTIC FIBROSIS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BRONCHITOL (mannitol) <sup>AL,CL,QL</sup> KALYDECO PACKET, TABLET (ivacaftor) <sup>QL, AL</sup> ORKAMBI (lumacaftor/ivacaftor) PACKET, TABLET <sup>QL, AL</sup> SYMDEKO (tezacaftor/ivacaftor) <sup>QL, AL</sup> TRIKAFTA (elexacaftor, tezacaftor, ivacaftor) <sup>AL, CL</sup>	<ul> <li>Bronchitol: Approved for diagnosis of CF and documentation that the patient has passed the BRONCHITOL Tolerance Test</li> <li>Kalydeco®: Diagnosis of CF and documentation of the drug-specific, FDA-approved mutation of CFTR gene</li> <li>Orkambi®: Diagnosis of CF and documentation of presence of two copies of the F580del mutation (homozygous) of CFTR gene</li> <li>Symdeko: Diagnosis of CF and documentation of the drug specific, FDA approved mutation of CFTR gene.</li> <li>Trikafta: Diagnosis of CF and documentation of at least one F508del mutation in the CFTR gene</li> </ul>

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#### **CYTOKINE & CAM ANTAGONISTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CART, PEN, SYR, VIAL QL HUMIRA (adalimumab) QL OTEZLA (apremilast) ORAL CL, QL	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIBINQO (abrocitinib)AL,NR,QL CIMZIA (certolizumab pegol)QL COSENTYX (secukinumab) ENSPRYNG (satralizumab-mwge) SUB-Q ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYRINGE KINERET (anakinra) OLUMIANT (baricitinib) TABCL,QL ORENCIA (abatacept) SUB-Q RINVOQ ER (upadacitinib)CL,QL SILIQ (brodalumab) SIMPONI (golimumab) SKYRIZI (risankizamab-rzaa) SYRINGE SKYRIZI PEN (risankizamab-rzaa) CTALTZ (ixekizumab)AL TREMFYA (guselkumab) XELJANZ (tofacitinib) TAB, SOLNCL,QL XELJANZ XR (tofacitinib) TABCL,QL	<ul> <li>Preferred agents will be approved with FDA-approved indication – ICD-10 diagnosis code is required.</li> <li>Non-preferred agents will be approved for FDA-approved indications in patients who have failed a trial of ONE preferred agent within this drug class, or upon diagnosis for non-preferred agent with FDA-approved indication if no preferred agent has FDA approval for diagnosis.</li> <li>Drug-specific criteria:</li> <li>Otezla: Requires a trial of Humira</li> <li>Olumiant: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira)</li> <li>Rinvoq: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira)</li> <li>Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira)</li> <li>Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira).</li> </ul>

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#### **DIURETICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SINGLE-AGEN	IT PRODUCTS	Non-preferred agents will be
amiloride TABLET bumetanide TABLET chlorothiazide TABLET chlorthalidone TABLET (generic Diuril) furosemide SOLUTION, TABLET   (generic Lasix) hydrochlorothiazide CAPSULE,     TABLET (generic Microzide) indapamide TABLET metolazone TABLET spironolactone TABLET (generic     Aldactone) torsemide TABLET	CAROSPIR (spironolactone) SUSPENSION eplerenone TABLET (generic Inspra) ethacrynic acid CAPSULE (generic Edecrin) KERENDIA (finerenone) TABLET NR,QL methyclothiazide TABLET THALITONE (chlorthalidone) TABLETNR triamterene (generic Dyrenium)	approved for patients who have failed a trial of <b>TWO</b> preferred agents within this drug class
COMBINATIO	N PRODUCTS	
amiloride/HCTZ <b>TABLET</b> spironolactone/HCTZ <b>TABLET</b> (generic Aldactazide) triamterene/HCTZ <b>CAPSULE</b> , <b>TABLET</b> (generic Dyazide, Maxzide)		

#### **ENZYME REPLACEMENT, GAUCHERS DISEASE**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ZAVESCA (miglustat) <sup>CL</sup>	CERDELGA (eliglustat) miglustat (generic Zavesca)	<ul> <li>Non-preferred agents require clinical documentation why the preferred product within this drug class is not appropriate</li> <li>Drug-specific criteria:</li> <li>Zavesca: Approved for mild to moderate type 1 Gaucher disease for whom enzyme replacement therapy is not a therapeutic option</li> </ul>

#### EPINEPHRINE. SELF-INJECTEDQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
epinephrine (AUTHORIZED GENERIC for Epipen/ Epipen Jr.) AUTOINJECTOR	epinephrine (generic for Adrenaclick) epinephrine (generic for Epipen/ Epipen Jr.) AUTOINJECTOR EPIPEN (epinephrine) AUTOINJ EPIPEN JR. (epinephrine) AUTOINJ SYMJEPI (epinephrine) PFS	Non-preferred agents require clinical documentation why the preferred product within this drug class is not appropriate  Brand name product may be authorized in event of documented national shortage of generic product.

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

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#### **ERYTHROPOIESIS STIMULATING PROTEINS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
RETACRIT (EPOETIN ALFA- EPBX)	EPOGEN (rHuEPO) PROCRIT (rHuEPO)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

#### FLUOROQUINOLONES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ciprofloxacin <b>TABLET</b> (generic Cipro) levofloxacin <b>TABLET</b> (generic Levaquin)	BAXDELA (delafloxacin) ciprofloxacin ER ciprofloxacin SUSPENSION (generic Cipro) levofloxacin SOLUTION moxifloxacin (generic Avelox) ofloxacin	<ul> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Baxdela: Coverable with documented intolerance or failure of preferred MRSA agents (clindamycin, doxycycline, linezolid, sulfamethoxazole/trimethoprim)</li> <li>Ciprofloxacin/Levofloxacin Suspension: Coverable with documented swallowing disorders</li> <li>Ofloxacin: Trial of preferred not required for diagnoses of Pelvic Inflammatory Disease OR Acute Epididymitis (nongonorrhea)</li> </ul>

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#### **GI MOTILITY, CHRONIC**

Preferred Agents Non-Preferred Agents	Prior Authorization/Class Criteria
LINZESS (linaclotide) <sup>QL</sup> MOVANTIK (naloxegol oxalate) <sup>QL</sup> BSRELA (tenapanor) <sup>AL,NR,QL</sup> lubiprostone (generic Amitiza) <sup>AL,QL</sup> MOTEGRITY (prucalopride succinate) RELISTOR (methylnaltrexone) TABLET <sup>QL</sup> SYMPROIC (naldemedine) TRULANCE (plecanatide) <sup>QL</sup> VIBERZI (eluxodoline)  • • • • •	Non-preferred agents will be approved for patients who have failed a 30-day trial of ONE preferred agent within this drug class  Ag-specific criteria:  Lotronex®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate  Relistor®: Covered for diagnosis of opioid-induced constipation in adults with chronic, non-cancer pain after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.) and failure of Movantik  Symproic: Covered for diagnosis of opioid-induced constipation in adult patients with chronic non-cancer pain after trial on at least TWO OTC laxatives and failure of Movantik  Trulance®: Covered for diagnosis of either chronic idiopathic constipation or IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.)  Viberzi®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate

#### **GLUCAGON AGENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) <sup>AL,QL</sup> <b>NASAL</b> GLUCAGON EMERGENCY (glucagon) <sup>QL</sup> <b>INJ KIT</b> (Lilly) glucagon <sup>QL</sup> <b>INJECTION</b> PROGLYCEM (diazoxide) <b>SUSP</b>	diazoxide SUSP (generic Proglycem) GLUCAGON EMERGENCY (glucagon) <sup>QL</sup> INJ KIT (Fresenius) GVOKE (glucagon) <sup>AL,QL</sup> KIT <sup>NR</sup> , PEN, SYRINGE, VIAL <sup>NR</sup> ZEGALOGUE (dasiglucagon) <sup>AL,NR, QL</sup> AUTO-INJECTOR, SYRINGE	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

#### with Prior Authorization Criteria

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#### **GLUCOCORTICOIDS, INHALED**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GLUCOCO	RTICOIDS	Non-preferred agents within the
GLUCOCO  ASMANEX (mometasone) <sup>QL,AL</sup> FLOVENT HFA (fluticasone) PULMICORT FLEXHALER (budesonide)  GLUCOCORTICOID/BRONCH  ADVAIR DISKUS (fluticasone/ salmeterol) <sup>QL</sup>	AEROSPAN (flunisolide) ALVESCO (ciclesonide) <sup>AL,CL</sup> ARMONAIR DIGIHALER (fluticasone) <sup>AL,QL</sup> ARMONAIR RESPICLICK (fluticasone) <sup>AL</sup> ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) <sup>CL,AL,QL</sup> FLOVENT DISKUS (fluticasone) QVAR (beclomethasone) QVAR Redihaler (beclomethasone)	<ul> <li>Non-preferred agents within the Glucocorticoids and Glucocorticoid/Bronchodilator Combo groups will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months</li> <li>Drug-specific criteria:</li> <li>budesonide respules: Covered without PA for age ≤ 8 years OR for diagnosis of eosinophilic esophagitis in patients ≥ 9 years, by GI biopsy or upper endoscopy. For other indications, must have failed a trial of two preferred agents within this drug class, within the last 6 months.</li> </ul>
ADVAIR HFA (fluticasone/salmeterol) <sup>QL</sup> DULERA (mometasone/formoterol) SYMBICORT (budesonide/ formoterol)	BREO ELLIPTA (fluticasone/vilanterol)  BREZTRI (budesonide/formoterol/ glycopyrrolate) <sup>QL</sup> Budesonide/formoterol (generic for Symbicort)  fluticasone/salmeterol (generic for Advair Diskus) <sup>QL</sup> fluticasone/salmeterol (generic for Airduo Respiclick)  TRELEGY ELLIPTA (fluticasone/ umeclidinium/vilanterol)  WIXELA INHUB (generic for Advair Diskus) <sup>QL</sup>	

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#### **GLUCOCORTICOIDS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORTEF (hydrocortisone) GRANULES <sup>AL</sup> CORTEF (hydrocortisone) cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSPENSION, TABLET <sup>CL</sup> ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg, 32mg ORTIKOS ER (budesonide) <sup>AL,QL</sup> PEDIAPRED (prednisolone sodium phosphate) prednisolone sodium phosphate (generic for Millipred/Veripred) prednisolone sodium phosphate ODT prednisone SOLUTION prednisone INTENSOL RAYOS DR (prednisone) TABLET TARPEYO (budesonide) <sup>NR</sup> CAPSULE	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> <li>Drug-specific criteria:         <ul> <li>Emflaza: Indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients 2 years of age and older</li> <li>Intensol Products: Patient specific documentation of why the less concentrated solution is not appropriate for the patient</li> </ul> </li> </ul>

#### **GROWTH HORMONES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GENOTROPIN (somatropin)	HUMATROPE (somatropin)	Growth Hormone PA Form
NORDITROPIN (somatropin)	NUTROPIN AQ (somatropin)	<b>Growth Hormone Criteria</b>
	OMNITROPE (somatropin)	
	SAIZEN (somatropin)	
	SEROSTIM (somatropin)	
	SKYTROFA (lonapegsomatropin-tcgd) <sup>NR</sup>	
	ZOMACTON (somatropin)	
	ZORBTIVE (somatropin)	

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#### H. PYLORI TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PYLERA (bismuth, metronidazole, tetracycline) <sup>QL</sup>	lansoprazole/amoxicillin/clarithromycin (generic Prevpac) <sup>QL</sup> OMECLAMOX-PAK (omeprazole, clarithromycin, amoxicillin) <sup>QL</sup>	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

#### HAE TREATMENTSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BERINERT (C1 esterase inhibitor, human) INTRAVENOUS HAEGARDA (C1 esterase inhibitor, human)AL,CL SUB-Q icatibant acetate (generic for FIRAZYR)AL SUB-Q	CINRYZE (C1 esterase inhibitor, human) <sup>AL,CL</sup> INTRAVENOUS FIRAZYR (icatibant acetate) <sup>AL</sup> SUB-Q ORLADEYO (berotralstat) CAP <sup>AL,QL</sup> RUCONEST (recombinant human C1 inhibitor) <sup>AL</sup> INTRAVENOUS TAKHZYRO (lanadelumab-flyo) <sup>AL,CL</sup> VIAL, SYRINGE <sup>NR</sup>	All agents require documentation of diagnosis of Type I or Type II HAE and deficient or dysfunctional C1 esterase inhibitor enzyme.     Concomitant use with ACE inhibitors, NSAIDs, or estrogencontaining products is contraindicated     Non-preferred agents will be approved for patients who have a failed trial or a contraindication to ONE preferred agent within this drug class  Drug-Specific Criteria     Cinryze, Haegarda, Orladeyo, and Takhzyro, require a history of
		two or more HAE attacks monthly, and trial and failure or contraindication to oral danazol

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#### **HEMOPHILIA TREATMENTS**

Preferred Agents	Non-Preferred Agents	Pri	or Authorization/Class Criteria
FACT	OR VIII	■ Non	-preferred agents will be
ALPHANATE HELIXATE FS HUMATE-P NOVOEIGHT NUWIQ XYNTHA KIT, SOLOFUSE	ADVATE ADYNOVATE AFSTYLA ELOCTATE ESPEROCT HEMOFIL-M JIVI <sup>AL</sup> KOATE-DVI KIT KOATE-DVI VIAL KOGENATE FS KOVALTRY OBIZUR RECOMBINATE	app faile age • Pati age to n	roved for patients who have and a trial of ONE preferred int within this drug class sents receiving a hemophilia int which moved from preferred on-preferred status on 1-21-21 be allowed to continue same
FAC	FOR IX	_	
ALPROLIX BENEFIX	ALPHANINE SD IDELVION IXINITY MONONINE PROFILNINE SD REBINYN RIXUBIS		
FACTOR VIIA AND PROTHROME	BIN COMPLEX-PLASMA DERIVED		
NOVOSEVEN RT	FEIBA NF SEVENFACT <sup>AL</sup>		
	XIII PRODUCTS		
COAGADEX CORIFACT	TRETTEN		
VON WILLEBR	AND PRODUCTS		
WILATE	VONVENDI		
BISPECIFI	C FACTORS		
HEMLIBRA			

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#### **HEPATITIS B TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
entecavir <b>TABLET</b>	adefovir dipivoxil BARACLUDE (entecavir) SOLUTION, TABLET EPIVIR HBV (lamivudine) TABLET, SOLUTION HEPSERA (adefovir dipivoxil) lamivudine hbv TABLET VEMLIDY (tenofovir alafenamide fumarate)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

#### with Prior Authorization Criteria

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#### **HEPATITIS C TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIRECT ACTIN	IG ANTI-VIRAL	Hepatitis C Treatments PA Form
sofosbuvir/velpatasvir (generic Epclusa) <sup>CL</sup> MAVYRET (glecaprevir/pibrentasvir) TABLET <sup>CL</sup> , PELLET <sup>AL,CL,NR</sup> VOSEVI (sofosbuvir/velpatasvir/ voxilaprevir) <sup>CL</sup>	HARVONI 200/45MG, TABLET (sofosbuvir/ledipasvir)CL HARVONI (ledipasvir/sofosbuvir)CL PELLET sofosbuvir/ledipasvir (generic Harvoni)CL SOVALDI (sofosbuvir)CL PELLET SOVALDI TABLET (sofosbuvir)CL VIEKIRA PAK (ombitasvir/ paritaprevir/ritonavir/dasabuvir)CL ZEPATIER (elbasvir/grazoprevir)CL	Non-preferred products require trial of preferred agents within the same group and/or will only be considered with documentation of why the preferred product within this drug class is not appropriate for patient     Patients newly eligible for Medicaid will be allowed to complete treatment with the original that treatment was initially authorized by another payor  Drug-specific criteria:  Trial with with a preferred agent not required in the following:     Harvoni:     Post liver transplant for genotype
DID	Many	1 or 4
	<b>VIRIN</b> REBETOL (ribavirin)	Vosevi: Requires documentation of non- response after previous treatment course of Direct Acting Anti-viral agent (DAA) for
	FERON	genotype 1-6 without cirrhosis or with compensated cirrhosis
PEGASYS (pegylated interferon alfa- 2a) <sup>CL</sup> PEG-INTRON (pegylated interferon alfa-2b) <sup>CL</sup>		

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#### **HISTAMINE II RECEPTOR BLOCKERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
famotidine <b>TABLET</b> (generic for Pepcid) nizatidine <b>SOLUTION</b> (generic for Axid)	cimetidine TABLET, SOLUTION <sup>CL</sup> (generic for Tagamet) famotidine SUSPENSION nizatidine CAP (generic for Axid) ranitidine CAPSULE, (generic for Zantac) ranitidine OTC, SYRUP, TABLET (generic for Zantac)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Cimetidine: Approved for viral M. contagiosum or common wart V. Vulgaris treatment</li> <li>cimetidine solution/ famotidine suspension/ranitidine syrup: Requires clinical reason why nizatidine syrup cannot be used ***famotidine suspension is authorized during shortage of nizatidine syrup.***</li> </ul>

#### with Prior Authorization Criteria

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#### HIV / AIDSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CCR5 AN	TAGONISTS	<ul> <li>Non-preferred agents will be</li> </ul>
SELZENTRY <b>SOLN</b> , <b>TAB</b> (maraviroc)	maraviroc (generic Selzentry) <sup>NR</sup>	approved for patients who have a diagnosis of HIV/AIDS and patien
FUSION I	NHIBITORS	specific documentation of why the preferred products within this drug
FUZEON <b>SUB-Q</b> (enfuvirtide) <sup>QL</sup>		class are not appropriate for patient, including, but not limited to, drug resistance or concomitan conditions not recommended with
HIV-1 ATTACH	IMENT INHIBITOR	preferred agents
	RUKOBIA ER (fostemsavir) <sup>AL,QL</sup>	<ul> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue.</li> </ul>
INTEGRASE STRAND TRA	NSFER INHIBITORS (INSTIS)	therapy
ISENTRESS (raltegravir) <sup>QL</sup> ISENTRESS HD (raltegravir)	TIVICAY PD (dolutegravir)	<ul><li>Diagnosis of HIV/AIDS required OR</li></ul>
TIVICAY (dolutegravir)		<ul> <li>Pre and Post Exposure Prophylaxis</li> </ul>
NON-NUCLEOSIDE REVERSE TRA	ANSCRIPTASE INHIBITORS (NNRTIS)	
efavirenz <b>CAPSULE, TABLET</b> (generio Sustiva) INTELENCE (etravirine) <sup>QL</sup>	EDURANT (rilpivirine) ETRAVIRINE (new generic for Intelence) <sup>NR,QL</sup>	
PIFELTRO (doravirine) <sup>QL</sup>	nevirapine IR, ER (generic Viramune/Viramune XR)	
	RESCRIPTOR (delavirdine) SUSTIVA CAPSULE, TABLET	
	(efavirenz) VIRAMUNE (nevirapine) SUSP	
NUCLEOSIDE REVERSE TRAN	SCRIPTASE INHIBITORS (NRTIs)	
abacavir <b>SOLN, TABLET</b> (generic Ziagen)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for	
EMTRIVA <b>CAPSULE</b> , <b>SOLN</b> (emtricitabine)	Emtriva) EPIVIR (lamivudine)	
amivudine <b>SOLN, TABLET</b> (generic Epivir)	RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit)	
zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir)	VIDEX (didanosine) <b>SOLN</b> ZIAGEN (abacavir)	
NUCLEOTIDE REVERSE TRAN	ISCRIPTASE INHIBITORS (NRTIs)	
tenofovir <b>TABLET</b> (generic Viread)	VIREAD (tenofovir) POWDER	
PHARMACOKIN	NETIC ENHANCER	
	TYBOST (cobicistat) <sup>QL</sup>	

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#### HIV / AIDS<sup>CL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROTEASE	INHIBITORS	
atazanavir CAPSULE (generic Reyataz) LEXIVA SUSP (fosamprenavir) ritonavir TABLET (generic Norvir)	APTIVUS CAPSULE, SOLN (tipranavir) CRIXIVAN (indinavir) fosamprenavir TAB (generic Lexiva) INVIRASE (saquinavir) LEXIVA TABLET (fosamprenavir) NORVIR POWDER, SOLN (ritonavir) NORVIR (ritonavir) TAB PREZISTA (darunavir) SUSP, TABLET REYATAZ POWDER (atazanavir) VIRACEPT (nelfinavir)	<ul> <li>Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> <li>Diagnosis of HIV/AIDS required</li> <li>OR</li> <li>Pre and Post Exposure Prophylaxis</li> </ul>

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HIV / AIDS <sup>CL</sup> (Continued)  Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
•	E INHIBITORS (PIs) or PIs plus	Non-preferred agents will be
PHARMACOKIN	NETIC ENHANCER	approved for patients who have a diagnosis of HIV/AIDS and patient
lopinavir/ritonavir <b>SOLN</b> (generic Kaletra)	KALETRA <b>SOLN</b> (lopinavir/ritonavir) KALETRA <b>TAB</b> (lopinavir/ritonavir) lopinavir/ritonavir <b>TAB</b> (generic Kaletra) <sup>NR</sup> PREZCOBIX (darunavir/cobicistat) <sup>QL</sup>	specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to drug resistance or concomitant conditions not recommended with preferred agents  Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy  Diagnosis of HIV/AIDS required OR  Pre and Post Exposure Prophylaxis
COMBINATION NUCLEOS(T)IDE RE	VERSE TRANSCRIPTASE INHIBITORS	
abacavir/lamivudine (generic Epzicom) CIMDUO (lamivudine/tenofovir) <sup>QL</sup> DESCOVY (emtricitabine/tenofovir) <sup>QL, CL</sup> lamivudine/zidovudine (generic Combivir) TRUVADA (emtricitabine/tenofovir)	abacavir/lamivudine/zidovudine (generic Trizivir)  COMBIVIR (lamivudine/zidovudine)  emtricitabine/tenofovir (generic Truvada) <sup>CL</sup> EPZICOM (abacavir sulfate/lamivudine)  TEMIXYS (lamivudine/tenofovir) <sup>QL</sup> TRIZIVIR  (abacavir/lamivudine/zidovudine)	Drug-Specific Criteria  Descovy:  • Approval will be granted for a diagnosis of HIV/AIDS  For PrEP use: Will require prior approval with a documentation of a contraindication to Truvada.

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

#### with Prior Authorization Criteria

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#### HIV / AIDS<sup>CL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COMBINATION PRODU	CTS – MULTIPLE CLASSES	
ATRIPLA (tenofovir/emtricitabine/efavirenz)  BIKTARVY (bictegravir/emtricitabine/tenofovir)QL  COMPLERA (rilpivirine/emtricitabine/tenofovir)  DELSTRIGO (doravirine/lamivudine/tenofovir)QL  GENVOYA (elvitegravier/cobicistat/emtricitabine/tenofovir)QL, AL  ODEFSEY (emtricitabine/rilpivirine/tenofovir)QL  STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir)QL  SYMFI (efavirenz/lamivudine/tenofovir)QL  SYMFI LO (efavirenz/lamivudine/tenofovir)QL  TRIUMEQ (dolutegravir/abacavir/lamivudine)	DOVATO (dolutegravir/lamivudine) <sup>QL</sup> efavirenz/emtricitabine/tenofovir   (generic Atripla) <sup>CL</sup> efavirenz/lamivudine/tenofovir   (generic for Symfi) <sup>QL</sup> efavirenz/lamivudine/tenofovir   (generic for Symfi Lo) <sup>QL</sup> JULUCA (dolutegravir/rilpivirine) <sup>QL</sup> SYMTUZA (darunavir/cobicistat/   emtricitabine/tenofovir) <sup>QL</sup> TRIUMEQ PD (abacavir, dolutegravir, and lamivudine) SUSP <sup>NR</sup>	<ul> <li>Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> <li>Diagnosis of HIV/AIDS required</li> <li>OR</li> <li>Pre and Post Exposure Prophylaxis</li> </ul>

#### with Prior Authorization Criteria

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#### HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acarbose (generic for Precose)	miglitol (generic for Glyset) GLYSET (miglitol)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

#### with Prior Authorization Criteria

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#### HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GLUCAGON-LIKE PEPTIDE-1 RE	CEPTOR AGONIST (GLP-1 RA) <sup>CL</sup>	Preferred agents require metformin
BYDUREON (exenatide ER) BYDUREON <b>PEN</b> (exenatide ER) subcutaneous BYETTA (exenatide) subcutaneous TRULICITY (dulaglutide) VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) BYDUREON BCISE PEN (exenatide) <sup>QL</sup> OZEMPIC (semaglutide) RYBELSUS (semaglutide) TANZEUM (albiglutide)  A COMBINATIONS  SOLIQUA (insulin glargine/lixisenatide)	trial and diagnosis of diabetes  Non-preferred agents will be approved for patients who have:  Failed a trial of TWO preferred agents within GLP-1 RA  AND  Diagnosis of diabetes with HbA1C ≥ 7 AND  Trial of metformin, or contraindication or intolerance to metformin
	XULTOPHY (insulin degludec/liraglutide)	
AMYLIN .	ANALOG	ALL criteria must be met
	SYMLIN (pramlintide) subcutaneous	<ul> <li>Concurrent use of short-acting mealtime insulin</li> <li>Current therapy compliance</li> <li>No diagnosis of gastroparesis</li> <li>HbA1C ≤ 9% within last 90 days</li> <li>Fingerstick monitoring of glucose during initiation of therapy</li> </ul>
DIPEPTIDYL PEPTIDASE-4 (DPP-4) IN	HIBITOR <sup>QL</sup>	
GLYXAMBI (empagliflozin/linagliptin) JANUMET (sitagliptin/metformin) JANUMET XR (sitagliptin/metformin) JANUVIA (sitagliptin) JENTADUETO (linagliptin/metformin) TRADJENTA (linagliptin)	alogliptin (generic for Nesina) alogliptin/metformin (generic for Kazano) JENTADUETO XR (linagliptin/metformin) KOMBIGLYZE XR (saxagliptin/metformin) ONGLYZA (saxagliptin) alogliptin/pioglitazone (generic for Oseni) QTERN (dapagliflozin/saxagliptin) STEGLUJAN (ertugliflozin/sitagliptin) TRIJARDY XR (empagliflozin/linagliptin/metformin) <sup>AL</sup>	Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within DPP-4

#### with Prior Authorization Criteria

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#### HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

HUMALOG (insulin lispro) U-100
glargine)  LANTUS (insulin glargine) VIAL  LEVEMIR (insulin detemir) PEN, VIAL  NOVOLOG (insulin aspart)  CARTRIDGE, FLEXPEN, VIAL  NOVOLOG MIX FLEXPEN, VIAL  (insulin aspart/aspart protamine)

#### **HYPOGLYCEMICS, MEGLITINIDES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
repaglinide (generic for Prandin)	nateglinide (generic for Starlix) <sup>CL</sup> repaglinide/metformin (generic for Prandimet) <sup>CL</sup>	Non-preferred agents will be approved for patients with:     Failure of a trial of ONE preferred agent in another Hypoglycemic class OR     T2DM and inadequate glycemic control

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#### HYPOGLYCEMICS, METFORMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
metformin IR & ER (generic Glucophage/Glucophage XR)	metformin ER (generic Fortamet/Glumetza) metformin <b>SOLUTION</b> (generic Riomet) RIOMET ER (metformin ER) <sup>AL</sup>	<ul> <li>Metformin ER (generic Fortamet®)/Glumetza®: Requires clinical reason why generic Glucophage XR® cannot be used</li> <li>Metformin solution: Prior authorization not required for age &lt;7 years</li> </ul>

#### **HYPOGLYCEMICS, SGLT2**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) <sup>QL,CL</sup> INVOKAMET (canagliflozin/metformin) <sup>QL,CL</sup> INVOKANA (canagliflozin) <sup>CL</sup> JARDIANCE (empagliflozin) <sup>QL,CL</sup> SYNJARDY (empagliflozin/metformin) <sup>AL,CL,QL</sup> XIGDUO XR (dapagliflozin/metformin) <sup>QL,CL</sup>	INVOKAMET XR (canagliflozin/metformin) <sup>QL</sup> SEGLUROMET (ertugliflozin/metformin) <sup>QL</sup> STEGLATRO (ertugliflozin) <sup>QL</sup> SYNJARDY XR (empagliflozin/metformin) <sup>AL,QL</sup>	<ul> <li>Preferred agents are Approved for diagnosis of diabetes AND a trial of metformin</li> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> <li>Drug Specific Criteria:</li> <li>Farxiga and Jardiance:         <ul> <li>Approved for a diagnosis of heart failure with reduced ejection fraction (NYHA class II-IV)</li> </ul> </li> </ul>

#### HYPOGLYCEMICS, SULFONYLUREAS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
glimepiride (generic Amaryl) glipizide IR & ER (generic Glucotrol/ Glucotrol XL) glyburide (generic Diabeta/Glynase)	chlorpropamide tolazamide tolbutamide	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class
SULFONYLUREA	COMBINATIONS	
glipizide/metformin glyburide/metformin (generic Glucovance)		

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NR – Product was not reviewed - New Drug criteria will apply

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#### **HYPOGLYCEMICS, TZD**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
THIZAOLIDINEDIONES (TZDs)		<ul> <li>Non-preferred agents will be</li> </ul>
pioglitazone (generic for Actos)	AVANDIA (rosiglitazone)	approved for patients who have failed a trial of THE preferred agent
TZD COM	BINATIONS	within this drug class
	pioglitazone/glimepiride (generic for Duetact) pioglitazone/metformin (generic for Actoplus Met)	Combination products: Require clinical reason why individual ingredients cannot be used

#### **IDIOPATHIC PULMONARY FIBROSIS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OFEV (nintedanib esylate) <sup>CL</sup>	ESBRIET (pirfenidone)	<ul> <li>Non-preferred agent requires trial of preferred agent within this drug class</li> <li>FDA approved indication required – ICD-10 diagnosis code</li> </ul>

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ASENRA (benralizumab) <sup>AL</sup> <b>PEN</b>	NUCALA (mepolizumab) <sup>AL</sup>	Asthma Immunomodulator PA Form
(OLAIR (omalizumab) <b>SYR</b> <sup>AL,QL</sup>	AUTO-INJ, SYR,	<ul> <li>Non-preferred agents require a tri of a preferred agent within this dru class with the same indication</li> </ul>
		Drug Specific Criteria:
		Dupixent: is indicated for
		- Patients 6 years and older as an add- maintenance treatment in patients with moderate-to-severe asthma with an eosinophilic phenotype or with oral corticosteroid dependent asthma
		- For other indications, see
		Immunomodulators, Atopic Dermatitis
		Fasenra: is indicated for
		- Patient 12 years and older for ad on maintenance treatment of severe asthma, and with an eosinophilic phenotype
		Nucala: is indicated for
		-Patients 6 years and older for ad on maintenance treatment of severe asthma, and with an eosinophilic phenotype
		-Patients 12 years and older with hypereosinophilic syndrome (HES) for ≥6 months without identifiable non-hematologic secondary cause
		-Patients 18 years and older for add- maintenance treatment of chroni rhinosinusitis with nasal polyps (CRWSwNP) with inadequate response to nasal corticosteroids
		-Adult patients with eosinophilic granulomatosis with polyangi
		Xolair Syringe- is indicated for
		-Patients 6 years and older for moderate to severe persistent asthma with a positive skin test of in vitro reactivity to a perennial aeroallergen and symptoms that are inadequately controlled with inhaled corticosteroids
		-Patients 12 years and older with Chronic spontaneous urticaria (CSU) who remain symptomatic despite H1 antihistamine treatme
		-Patients 18 years and older with Nas Polyps with inadequate response nasal corticosteroids. As add-on maintenance treatment

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#### IMMUNOMODULATORS, ATOPIC DERMATITISAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIDEL (pimecrolimus) EUCRISA (crisaborole) <sup>CL,QL</sup>	ADBRY (tralokinumab-ldrm) SUB-Q <sup>AL,NR,QL</sup> DUPIXENT (dupilumab) <sup>AL,CL</sup> DUPIXENT PEN <sup>AL</sup> OPZELURA (ruxolitinib phosphate) CREAM <sup>AL,NR,QL</sup> pimecrolimus (generic for Elidel) tacrolimus (generic for Protopic) <sup>CL</sup>	<ul> <li>Non-preferred agents require: Trial of a topical steroid AND Trial of one preferred product within this drug class</li> <li>Drug-specific criteria:</li> <li>Dupixent: Indicated for the treatment of patients aged 6 years and older with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. DUPIXENT can be used with or without topical corticosteroids.</li> <li>-as an add-on maintenance treatment of patients aged 6 years and older with moderate-to-severe asthma characterized by an eosinophilic phenotype or with oral corticosteroid dependent asthma.</li> <li>- as an add-on maintenance treatment in adult patients with inadequately controlled chronic rhinosinusitis with nasal polyposis (CRSwNP)</li> <li>Eucrisa: Requires use and failure of 1 topical steroid or Elidel.</li> </ul>

#### IMMUNOMODULATORS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
imiquimod (generic for Aldara)	ALDARA (imiquimod) imiquimod (generic for Zyclara) podofilox (generic for Condylox) VEREGEN (sinecatechins) ZYCLARA (imiquimod)	Non-preferred agents require clinical reason why preferred agent within this drug class cannot be used

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#### **IMMUNOSUPPRESSIVES, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathioprine (generic Imuran) azathioprine (generic Azasan) <sup>NR</sup> cyclosporine, modified CAPSULE (generic Neoral) mycophenolate CAPSULE, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLUTION RAPAMUNE (sirolimus) TABLET tacrolimus ZORTRESS (everolimus) AL	ASTAGRAF XL (tacrolimus) AZASAN (azathioprine) cyclosporine CAPSULE, SOFTGEL cyclosporine, modified SOLUTION   (generic Neoral) ENVARSUS XR (tacrolimus) everolimus (generic for Zortress) <sup>AL</sup> GENGRAF (cyclosporine, modified) CAPSULE, SOLUTION mycophenolate SUSPENSION   (generic Cellcept) mycophenolic acid MYFORTIC (mycophenolate sodium) PROGRAF (tacrolimus) CAPSULE, PACKET REZUROCK (belumosudil) <sup>AL,NR,QL</sup> TAB SANDIMMUNE (cyclosporine) CAPSULE, SOLUTION sirolimus SOLUTION, TABLET   (generic Rapamune) TAVNEOS (avacopan) <sup>NR,QL</sup> CAPSULE	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class  Patients established on existing therapy will be allowed to continue

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#### **INTRANASAL RHINITIS DRUGS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOLINERGICS		Non-preferred agents will be approved
ipratropium (generic for Atrovent)		for patients who have failed a 30-day trial of ONE preferred agent within this
ANTIHIS'	TAMINES	drug class
azelastine 0.1% (generic for Astelin)	azelastine 0.15% (generic for Astepro) azelastine/fluticasone (generic for Dymista) olopatadine (generic for Patanase)	<ul> <li>Drug-specific criteria:</li> <li>mometasone: Prior authorization NOT required for children ≤ 12 years</li> <li>budesonide: Approved for use in Pregnancy (Pregnancy Category</li> </ul>
CORTICO	STEROIDS	<ul> <li>B)</li> <li>Xhance: Indicated for treatment of</li> </ul>
fluticasone (generic for Flonase)	BECONASE AQ (beclomethasone) budesonide Rx (generic for Rhinocort) flunisolide (generic for Nasalide) mometasone (generic for Nasonex) OMNARIS (ciclesonide) QNASL 40 & 80 (beclomethasone) TICANASE (fluticasone) XHANCE (fluticasone) ZETONNA (ciclesonide)	nasal polyps in <u>&gt;</u> 18 years only

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#### **LEUKOTRIENE MODIFIERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
montelukast <b>TABLET/CHEWABLE</b> (generic for Singulair) <sup>AL</sup>	montelukast <b>GRANULES</b> (generic for Singulair) <sup>CL, AL</sup> zafirlukast (generic for Accolate) zileuton ER (generic for Zyflo CR) ZYFLO (zileuton)	<ul> <li>Non-preferred agents will be approved for patients who have failed a 30-day trial of THE preferred agent within this drug class</li> <li>Drug-specific criteria:         <ul> <li>montelukast granules:</li> <li>PA not required for age &lt; 2 years</li> </ul> </li> </ul>

#### LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clindamycin <b>CAPSULE</b> clindamycin palmitate <b>SOLUTION</b> linezolid <b>TABLET</b>	CLEOCIN (clindamycin ) CAPSULE CLEOCIN PALMITATE (clindamycin) linezolid SUSPENSION SIVEXTRO (tedizolid phosphate) ZYVOX (linezolid) SUSPENSION, TABLET	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

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#### LIPOTROPICS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BILE ACID SE	QUESTRANTS	<ul> <li>Non-preferred agents will be</li> </ul>
cholestyramine (generic Questran) colestipol TABLETS (generic Colestid)	colesevelam (generic Welchol)  TABLET, PACKET colestipol GRANULES (generic Colestid) QUESTRAN LIGHT (cholestyramine)	approved for patients who have failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:  Colesevelam: Trial not required for diabetes control and monotherapy with
TREATMENT OF HOMOZYGOUS FA	MILIAL HYPERCHOLESTEROLEMIA	metformin, sulfonylurea, or insulin has been inadequate
	JUXTAPID (lomitapide) <sup>CL</sup> KYNAMRO (mipomersen) <sup>CL</sup>	<ul> <li>Juxtapid®/ Kynamro®:</li> <li>Approved for diagnosis of homozygous familial hypercholesterolemia (HoFH)</li> </ul>
FIBRIC ACID	DERIVATIVES	OR
fenofibrate (generic Tricor) fenofibrate (generic Lofibra) gemfibrozil (generic Lopid)  NIA niacin ER (generic for Niaspan)	fenofibric acid (generic Fibricor/Trilipix) fenofibrate (generic Antara/Fenoglide/ Lipofen/Triglide)  CIN  NIACOR (niacin IR) NIASPAN (niacin ER)	<ul> <li>Treatment failure/maximized dosing/contraindication to ALL the following: statins, ezetimibe, niacin, fibric acid derivatives, omega-3 agents, bile acid sequestrants</li> <li>Require faxed copy of REMS PA form</li> <li>Vascepa®: Approved for TG ≥ 500</li> </ul>
OMEGA-3 F	ATTY ACIDS	-
omega-3 fatty acids (generic for Lovaza)	icosapent (generic for Vascepa) <sup>CL</sup> omega-3 OTC VASCEPA (icosapent) <sup>CL</sup>	
CHOLESTEROL ABSO	ORPTION INHIBITORS	
ezetimibe (generic for Zetia)	NEXLIZET (bempedoic acid/ ezetimibe) <sup>QL</sup>	

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#### LIPOTROPICS, OTHER (continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROPROTEIN CONVERTASE SUI		<ul> <li>Prior Authorization/Class Criteria</li> <li>Praluent®: Approved for diagnoses of:         <ul> <li>atherosclerotic cardiovascular disease (ASCVD)</li> <li>heterozygous familial hypercholesterolemia (HeFH)</li> <li>Homozygous familial hypercholesterolemia (HoFH) as an adjunct to other LDL-C lowering therapies</li> </ul> </li> <li>MND         <ul> <li>Maximized high-intensity statin WITH ezetimibe for at 3 continuous months</li> <li>Failure to reach target LDL-C levels: ASCVD - &lt; 70 mg/dL, HeFH - &lt; 100 mg/dL</li> </ul> </li> <li>Repatha®: Approved for:         <ul> <li>adult diagnoses of atherosclerotic cardiovascular disease (ASCVD)</li> <li>heterozygous familial hypercholesterolemia (HeFH)</li> <li>homozygous familial hypercholesterolemia (HoFH) in age ≥ 13</li> <li>statin-induce rhabdomyolysis</li> </ul> </li> <li>AND         <ul> <li>Maximized high-intensity statin WITH ezetimibe for 3+ continuous months</li> <li>Failure to reach target LDL-C levels: ASCVD - &lt; 70 mg/dL, HeFH - &lt; 100 mg/dL</li> <li>Concurrent use of maximally-tolerated statin must continue, except for statin-induced rhabdomyolysis or a contraindication to a statin</li> </ul> </li> </ul>

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#### LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
STATINS		Non-preferred agents will be
atorvastatin (generic Lipitor) <sup>QL</sup> lovastatin (generic Mevacor) pravastatin (generic Pravachol) rosuvastatin (generic Crestor) simvastatin (generic Zocor)	ALTOPREV (lovastatin ER) <sup>CL</sup> EZALLOR SPRINKLE (rosuvastatin) <sup>QL</sup> fluvastatin IR/ER (generic Lescol/ Lescol XL) LIVALO (pitavastatin) ZYPITAMAG (pitavastatin)	<ul> <li>approved for patients who have failed a trial of TWO preferred agent within this drug class, within the last 12 months</li> <li>Drug-specific criteria:         <ul> <li>Altoprev®: One of the TWO trials must be IR lovastatin</li> </ul> </li> <li>Combination products: Require clinical</li> </ul>
STATIN CO	BINATIONS	reason why individual ingredients cannot be used
	atorvastatin/amlodipine (generic Caduet) simvastatin/ezetimibe (generic Vytorin)	<ul> <li>fluvastatin ER: Requires trial of TWO preferred agents AND trial of IR fluvastatin OR clinical reason IR cannot be used</li> <li>simvastatin/ezetimibe: Approved for 3-month continuous trial of ONE standard dose statin</li> </ul>

#### **MACROLIDES AND KETOLIDES, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MACROLIDES		Require clinical reason why
· · · · · · · · · · · · · · · · · · ·	clarithromycin ER (generic Biaxin XL) E.E.S. SUSPENSION (erythromycin ethylsuccinate) E.E.S. TABLET (erythromycin ethylsuccinate) ERY-TAB (erythromycin) ERYPED SUSPENSION (erythromycin) ERYTHROCIN (erythromycin) erythromycin base TABLET, CAPSULE	preferred products within this drug class cannot be used AND ≥ 3-day trial on a preferred product

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#### **METHOTREXATE**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
methotrexate PF VIAL, TABLET, VIAL	OTREXUP (methotrexate) SUB-Q RASUVO (methotrexate) SUB-Q REDITREX	<ul> <li>Non-preferred agents will be approved for FDA-approved indications</li> <li>Drug-specific criteria:</li> <li>Xatmep<sup>TM</sup>:Indicated for pediatric patients only</li> </ul>

#### **MOVEMENT DISORDERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) <sup>CL</sup> INGREZZA (valbenazine) <sup>AL,CLQL</sup> CAP	INGREZZA (valbenazine) <sup>CL</sup> <b>INITIATION PACK</b> XENAZINE (tetrabenazine) <sup>CL</sup>	All drugs require an FDA approved indication – ICD-10 diagnosis code required.
tetrabenazine (generic for Xenazine) <sup>CL</sup>		Non-preferred agents require trial of Austedo
		<ul> <li>Austedo: Diagnosis of Tardive Dyskinesia or chorea associated with Huntington's Disease; Requires a Step through tetrabenazine with the diagnosis of chorea associated with Huntington's Disease</li> <li>Ingrezza: Diagnosis of Tardive Dyskinesia in adults</li> <li>tetrabenazine: Diagnosis of chorea with Huntington's Disease</li> </ul>

#### with Prior Authorization Criteria

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#### **MULTIPLE SCLEROSIS DRUGS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AVONEX (interferon beta-1a) <sup>QL</sup> BETASERON (interferon beta-1b) <sup>QL</sup> COPAXONE 20mg (glatiramer) <sup>QL</sup> KESIMPTA (Ofatumumab) <sup>CL,QL</sup> TECFIDERA (dimethyl fumarate)	AUBAGIO (teriflunomide)  BAFIERTAM (monomethyl fumarate) <sup>QL</sup> dalfampridine (generic Ampyra) <sup>QL</sup> dimethyl fumarate (generic for Tecfidera)  EXTAVIA (interferon beta-1b) <sup>QL</sup> GILENYA (fingolimod) <sup>QL</sup> glatiramer (generic Copaxone) <sup>QL</sup> MAVENCLAD (cladribine) MAYZENT (siponimod) <sup>QL</sup> PLEGRIDY (peginterferon beta-1a) <sup>QL</sup> PONVORY (ponesimod) <sup>NR</sup> REBIF (interferon beta-1a) <sup>QL</sup> VUMERITY (diroximel) <sup>QL</sup> ZEPOSIA (ozanimod) <sup>AL,QL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:         <ul> <li>Ampyra®: Approved for diagnosis of gait disorder associated with MS AND EDSS score ≤ 7</li> <li>Plegridy: Approved for diagnosis of relapsing MS</li> <li>Kesimpta: Approved for patients who have failed a trial of a preferred injectable agent within this class</li> </ul> </li> </ul>

#### **NITROFURAN DERIVATIVES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
nitrofurantoin macrocrystals CAPSULE (generic for Macrodantin) nitrofurantoin monohydrate- macrocrystals CAPSULE (generic for Macrobid)	nitrofurantoin <b>SUSPENSION</b> (generic for Furadantin)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

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#### **NSAIDs, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
·	diclofenac potassium (generic for Cataflam, Zipsor) diclofenac SR (generic for Voltaren-XR) diflunisal (generic for Dolobid) etodolac & SR (generic for Lodine/XL) fenoprofen (generic for Nalfon) flurbiprofen (generic for Ansaid) ibuprofen OTC (generic for Advil, Motrin) CAPSULE	Prior Authorization/Class Criteria  Non-preferred agents within COX-1 SELECTIVE group will be approved for patients who have failed no less than 30-day trial of TWO preferred agents within this drug class  Drug-specific criteria:  Arthrotec®: Requires clinical reason why individual ingredients cannot be used  Duexis®/Vimovo®: Requires clinical reason why individual agents cannot be used
nabumetone (generic for Relafen) naproxen Rx, OTC (generic for Naprosyn) naproxen enteric coated sulindac (generic for Clinoril)	indomethacin ER (generic for Indocin) INDOCIN RECTAL, SUSPENSION ketoprofen & ER (generic for Orudis) meclofenamate (generic for Meclomen) mefenamic acid (generic for Ponstel) meloxicam CAP   (generic Vivlodex) <sup>CL, QL</sup> naproxen CR (generic for Naprelan) naproxen SUSPENSION (generic for Naprosyn) naproxen sodium (generic for Anaprox) naproxen-esomeprazole (generic for Vimovo) oxaprozin (generic for Daypro) piroxicam (generic for Feldene) RELAFEN DS (nabumetone) tolmetin (generic for Tolectin) Ketorolac Nasal QL (generic for Sprix)	<ul> <li>meclofenamate: Approvable without trial of preferred agents for menorrhagia</li> </ul>

#### with Prior Authorization Criteria

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#### **NSAIDs, ORAL (Continued)**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COX-I SELECT	IVE (continued)	
	ALL BRAND NAME NSAIDs including:  CAMBIA (diclofenac oral solution)  DUEXIS (ibuprofen/famotidine) <sup>CL</sup> ibuprofen/famotidine (generic Duexis) <sup>CL</sup> SPRIX (ketorolac nasal spray)  NASAL <sup>QL, CL</sup> TIVORBEX (indomethacin)  VIVLODEX (meloxicam submicronized)  ZIPSOR (diclofenac)  ZORVOLEX (diclofenac)	<ul> <li>Drug-specific criteria:</li> <li>Sprix®: Approved for patients unable to tolerate, swallow OR absorb oral NSAIDs OR contraindication OR trial of TWO preferred oral NSAIDs</li> <li>Tivorbex®: Requires clinical reason why indomethacin capsules cannot be used</li> <li>Zorvolex®: Requires trial of oral diclofenac OR clinical reason why diclofenac potassium/sodium cannot be used</li> </ul>
NSAID/GI PROTECTA	ANT COMBINATIONS	
	diclofenac/misoprostol (generic for Arthrotec)	
COX-II SE	ELECTIVE	
celecoxib (generic for Celebrex)		

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#### **NSAIDs, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
diclofenac sodium GEL (OTC only)	diclofenac (generic for Pennsaid Solution) <sup>CL</sup> FLECTOR <b>PATCH</b> (diclofenac) <sup>CL</sup> LICART <b>PATCH</b> (diclofenac) <sup>CL</sup> PENNSAID <b>PACKET</b> , <b>PUMP</b> (diclofenac) <sup>CL</sup> VOLTAREN <b>GEL</b> (diclofenac) <sup>CL</sup>	Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class  Drug Specific Criteria  Flector®/Licart: Approved for diagnosis of acute pain due to sprain/strain/contusion AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form  Pennsaid®: Approved for osteoarthritis of the knees AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form  Pennsaid® Pump: Requires clinical reason why 1.5% solution cannot be used  Voltaren®: Approved for diagnosis of osteoarthritis AND trial of oral diclofenac OR clinical resaon patient cannot use oral dosage form

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NOTE: Other oral oncology agents not listed here may also be available. See <a href="https://nebraska.fhsc.com/default.asp">https://nebraska.fhsc.com/default.asp</a> for coverage information and prior authorization status for products not listed.

#### **ONCOLOGY AGENTS, ORAL, BREAST**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
IBRANCE (palbociclib)	NHIBITOR  KISQALI (ribociclib)  KISQALI FEMARA CO-PACK  VERZENIO (abemaciclib)	Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
cyclophosphamide XELODA (capecitabine)	CHERAPY  capecitabine (generic for Xeloda) <sup>CL</sup>	<ul> <li>Drug-specific critera</li> <li>anastrozole: May be approved for malignant neoplasm of male breast (male breast cancer)</li> </ul>
anastrozole (generic for Arimidex) exemestane (generic for Aromasin) letrozole (generic for Femara) tamoxifen citrate (generic for Nolvadex)	BLOCKADE  SOLTAMOX SOLN (tamoxifen) <sup>CL</sup> toremifene (generic for Fareston) <sup>CL</sup>	<ul> <li>capecitabine: Requires trial of Xeloda or clinical reason Xeloda cannot be used</li> <li>Fareston®: Require clinical reason why tamoxifen cannot be used</li> <li>letrozole: Approved for diagnosis of breast cancer with day supply greater than 12 – NOT approved</li> </ul>
OTI	NERLYNX (neratinib) PIQRAY (alpelisib) lapatinib (generic Tykerb) <sup>CL</sup> TALZENNA (talazoparib tosylate) QL TUKYSA(tucatinib) <sup>QL</sup>	for short term use  Soltamox: May be approved with documented swallowing difficulty

#### with Prior Authorization Criteria

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#### **ONCOLOGY AGENTS, ORAL, HEMATOLOGIC**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
mercaptopurine	PURIXAN (mercaptopurine) <sup>AL</sup>	<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation</li> </ul>
	AML	submitted supporting off-label use from current treatment guidelines
IMBRUVICA (ibrutinib) LEUKERAN (chlorambucil) VENCLEXTA (venetoclax)	DAURISMO (glasdegib maleate) <sup>QL</sup> IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) <sup>QL</sup> XOSPATA (gilteritinib) <sup>QL</sup> CLL  COPIKTRA (duvelisib) <sup>QL</sup> ZYDELIG (idelalisib)	<ul> <li>Drug-specific critera</li> <li>Hydrea®: Requires clinical reasor why generic cannot be used</li> <li>Melphalan: Requires trial of Alkeran or clinical reason Alkeran cannot be used</li> <li>Purixan: Prior authorization not required for age ≤12 or for documented swallowing disorder</li> <li>Tabloid: Prior authorization not required for age &lt;19</li> </ul>
	CML	Tasigna: Patients receiving
hydroxyurea (generic for Hydrea) imatinib (generic for Gleevec) MYLERAN (busulfan) SPRYCEL (dasatinib)	BOSULIF (bosutinib) GLEEVEC (imatinib) HYDREA (hydroxyurea) ICLUSIG (ponatinib) SCEMBLIX (asciminib) <sup>NR</sup> TASIGNA (nilotinib) <sup>CL</sup>	<ul> <li>Tasigna, which changed from preferred to non-preferred on 1-17-19 will be allowed to continue therapy</li> <li>Xpovio: Indicated for relapsed or refractory multiple myeloma. Requires concomitant therapy with</li> </ul>
	MPN	dexamethasone
JAKAFI (ruxolitinib)		
MY	ELOMA	
ALKERAN (melphalan) REVLIMID <sup>QL</sup> (lenalidomide)	FARYDAK (panobinostat)  lenalidomide <sup>NR,QL</sup> (generic for Revlimid)  melphalan (generic for Alkeran)  NINLARO (ixazomib)  POMALYST (pomalidomide)  THALOMID (thalidomide)  XPOVIO (selinexor) CL	
0	THER	
MATULANE (procarbazine) TABLOID (thioguanine) tretinoin (generic for Vesanoid) <sup>AL</sup>	BRUKINSA (zanubrutinib <sup>QL</sup> CALQUENCE (acalabrutinib) <sup>QL</sup> INREBIC (fedratinib dihydrochloride) <sup>QL</sup> INQOVI (decitabine/cedazuridine) UKONIQ (umbralisb) <sup>NR</sup> VONJO (pacritinib) <sup>NR,QL</sup> ZOLINZA (vorinostat)	

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#### **ONCOLOGY AGENTS, ORAL, LUNG**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALECENSA (alectinib)	ALK  ALUNBRIG (brigatinib) <sup>QL</sup> LORBRENA (lorlatinib) <sup>QL</sup> ZYKADIA (ceritinib) CAPSULE, TABLET	<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines</li> <li>Drug-Specific Criteria</li> <li>Iressa/ Xalkori: Patients receiving Iressa or Xalkori prior to 1/21/21 (which changed from preferred to non-preferred) will be allowed to continue current treatment</li> </ul>
ALK /	ROS1 / NTRK	
	ROZLYTREK (entrectinib) <sup>AL,QL</sup> XALKORI (crizotinib)	
	EGFR	
TAGRISSO (osimertinib)	erlotinib (generic for Tarceva) EXKIVITY (mobocertinib) <sup>NR,QL</sup> GILOTRIF (afatinib) IRESSA (gefitinib) TARCEVA (erlotinib) VIZIMPRO (dacomitinib) <sup>QL</sup>	
(	OTHER	
	GAVRETO (pralsetinib) <sup>QL</sup> HYCAMTIN (topotecan) LUMAKRAS (sotrasib) <sup>QL</sup> RETEVMO (selpercatinib) <sup>AL</sup> TABRECTA (capmatinib) <sup>QL</sup> TEPMETKO (tepotinib) <sup>QL</sup>	

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#### **ONCOLOGY AGENTS, ORAL, OTHER**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CAPRELSA (vandetanib) GLEOSTINE (lomustine) LYNPARZA (olaparib) temozolomide (generic for Temodar) ZEJULA (niraparib)	AYVAKIT (avapritinib) <sup>AL,NR,QL</sup> BALVERSA (erdafitinib) COMETRIQ (cabozantinib) HEXALEN (altretamine) KOSELUGO (selumetinib) <sup>AL</sup> LONSURF (trifluridine/tipiracil) PEMAZYRE (pemigatinib) <sup>QL</sup> RUBRACA (rucaparib) STIVARGA (regorafenib) TAZVERIK (tazemetostat) <sup>AL</sup> TURALIO (pexidartinib) <sup>QL</sup> TRUSELTIQ (infigratinib) CAPSULE VITRAKVI (larotrectinib) CAPSULE, SOLUTION <sup>QL</sup>	Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines

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#### **ONCOLOGY AGENTS, ORAL, PROSTATE**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
abiraterone (generic for Zytiga) <sup>AL,QL</sup> bicalutamide (generic for Casodex) flutamide XTANDI (enzalutamide) <sup>AL,QL</sup> ZYTIGA (abiraterone) <sup>AL,QL</sup>	EMCYT (estramustine) ERLEADA (apalutamide) <sup>QL</sup> nilutamide (generic for Nilandron) NUBEQA (darolutamide) <sup>QL</sup> YONSA (abiraterone acetonide, submicronized)	Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines

#### **ONCOLOGY AGENTS, ORAL, RENAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INLYTA (axitinib) LENVIMA (lenvatinib) SUTENT (sunitinib) VOTRIENT (pazopanib	AFINITOR DISPERZ (everolimus) <sup>CL</sup> CABOMETYX (cabozantinib) everolimus (generic for Afinitor) everolimus <b>SUSP</b> (generic for Afinitor Disperz) <sup>NR</sup> FOTIVDA (tivozanib) <sup>NR</sup> NEXAVAR (sorafenib) sunitinib malate (generic for Sutent) WELIREG (belzutifan) <sup>NR,QL</sup>	<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines</li> <li>Drug-specific critera</li> <li>Afinitor: Patients receiving Afinitor, which changed from preferred to non-preferred on 1-17-19 will be allowed to continue therapy</li> </ul>

#### **ONCOLOGY AGENTS, ORAL, SKIN**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BASAL CELL		<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent,</li> </ul>
ERIVEDGE (vismodegib)	ODOMZO (sonidegib) <sup>CL</sup>	but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
BRAF MUTATION		
MEKINIST (trametinib)	BRAFTOVI (encorafenib)	Drug-specific critera
TAFINLAR (dabrafenib)	COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)	<ul> <li>Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17- 19 will be allowed to continue therapy</li> </ul>

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

NR – Product was not reviewed - New Drug criteria will apply

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#### with Prior Authorization Criteria

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#### **OPHTHALMICS, ALLERGIC CONJUNCTIVITIS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALREX (loteprednol 0.2%) cromolyn (generic for Opticrom) ketotifen OTC (generic for Zaditor) olopatadine 0.1% (generic for Patanol) olopatadine 0.2% (generic for Pataday once daily, Pataday twice daily)	ALOCRIL (nedocromil) ALOMIDE (lodoxamide) azelastine (generic for Optivar) BEPREVE (bepotastine besilate) bepotastine besilate (generic for Bepreve) EMADINE (emedastine) epinastine (generic for Elestat) LASTACAFT (alcaftadine) LASTACAFT (alcaftadine) PATADAY XS (olopatadine 0.7%) PATADAY OTC (olopatadine 0.2%) PAZEO (olopatadine 0.7%) ZERVIATE (certirizine) AL	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul>

## with Prior Authorization Criteria

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### **OPHTHALMICS, ANTIBIOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FLUOROQU	JINOLONES	<ul> <li>Non-preferred agents will be</li> </ul>
ciprofloxacin <b>SOLUTION</b> (generic for Ciloxan) ofloxacin (generic for Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic for Zymaxid) levofloxacin MOXEZA (moxifloxacin) moxifloxacin (generic for Vigamox) moxifloxacin (generic for Moxeza) VIGAMOX (moxifloxacin)	<ul> <li>approved for patients who have failed a one-month trial of TWO preferred agent within this drug class</li> <li>Azasite®: Approval only requires trial of erythromycin</li> <li>Drug-specific criteria:</li> <li>Natacyn®: Approved for documented fungal infection</li> </ul>
MACRO	DLIDES	_
erythromycin	AZASITE (azithromycin) <sup>CL</sup>	
AMINOGL	YCOSIDES	
gentamicin <b>SOLUTION</b> tobramycin (generic for Tobrex drops)		
	ALMIC AGENTS	
bacitracin/polymyxin B (generic Polysporin) polymyxin B/trimethoprim (generic for Polytrim)	bacitracin NATACYN (natamycin) <sup>CL</sup> neomycin/bacitracin/polymyxin B OINTMENT neomycin/polymyxin B/gramicidin NEOSPORIN (neomycin/polymyxin B/gramcidin) sulfacetamide SOLUTION (generic for Bleph-10) sulfacetamide OINTMENT	

# with Prior Authorization Criteria

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### **OPHTHALMICS, ANTIBIOTIC-STEROID COMBINATIONS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
neomycin/polymyxin/dexamethasone (generic for Maxitrol) sulfacetamide/prednisolone TOBRADEX SUSPENSION, OINTMENT (tobramycin and dexamethasone)	BLEPHAMIDE (prednisolone and sulfacetamide) BLEPHAMIDE S.O.P. neomycin/polymyxin/HC neomycin/bacitracin/poly/HC PRED-G SUSPENSION, OINTMENT (prednisolone/gentamicin) tobramycin/dexamethasone SUSPENSION (generic for Tobradex) TOBRADEX S.T. (tobramycin and dexamethasone) ZYLET (loteprednol, tobramycin)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul>

### with Prior Authorization Criteria

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### **OPHTHALMICS, ANTI-INFLAMMATORIES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CORTICO	STEROIDS	Non-preferred agents will be
fluorometholone 0.1% (generic for FML) <b>OINTMENT</b> LOTEMAX <b>SOLUTION</b> (loteprednol 0.5%) MAXIDEX (dexamethasone) PRED MILD (prednisolone 0.12%)	dexamethasone (generic for Maxidex) difluprednate (generic Durezol) <sup>NR</sup> DUREZOL (difluprednate) FLAREX (fluorometholone) FML (fluorometholone 0.1% SOLUT.) FML FORTE (fluorometholone 0.25%) FML S.O.P. (fluorometholone 0.1%) INVELTYS (loteprednol etabonate) LOTEMAX OINTMENT, GEL (loteprednol) loteprednol GEL (generic for Lotemax Gel) loteprednol 0.5% SOLUTION (generic for Lotemax SOLUTION) prednisolone acetate 1% (gen. for Omnipred, Pred Forte) prednisolone sodium phosphate	<ul> <li>approved for patients who have failed a trial of TWO preferred agents within this drug class</li> <li>NSAID class: Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent</li> </ul>
NS	AID	-
diclofenac (generic for Voltaren) ketorolac 0.5% (generic for Acular)	ACUVAIL (ketorolac 0.45%) BROMSITE (bromfenac) bromfenac 0.09% (generic for Bromday) flurbiprofen (generic for Ocufen) ILEVRO (nepafenac 0.3%) ketorolac LS 0.4% (generic for Acular LS) NEVANAC (nepafenac) PROLENSA (bromfenac 0.07%)	

### OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
RESTASIS (cyclosporine) RESTASIS MULTIDOSE (cyclosporine) XIIDRA (lifitegrast)	CEQUA (cyclosporine) QL maravi EYSUVIS (loteprednol etabonate)QL TYRVAYA (varenicline tartrate)NR, QL	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  –

AL – Age Limit

## with Prior Authorization Criteria

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### **OPHTHALMICS, GLAUCOMA**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIOTICS		<ul> <li>Non-preferred agents will be</li> </ul>
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide)  VUITY (pilocarpine) <sup>NR</sup>	approved for patients who have failed a trial of ONE preferred agent within this drug class
SYMPATHO	MIMETICS	
Alphagan P (brimonidine 0.15%) brimonidine 0.2% (generic for Alphagan)	Alphagan P (brimonidine 0.1%) apraclonidine (generic for lopidine) brimonidine P 0.15%	
BETA BLO	OCKERS	
levobunolol (generic for Betagan) timolol (generic for Timoptic)	betaxolol (generic for Betoptic) BETIMOL (timolol) BETOPTIC S (betaxolol) carteolol (generic for Ocupress) timolol (generic for Istalol) timolol (generic for Timoptic Ocudose) TIMOPTIC OCUDOSE TIMOPTIC XE (timolol gel forming solution)	
CARBONIC ANHYDR	RASE INHIBITORS	
. ,	AZOPT (brinzolamide) brinzolamide (generic for Azopt)	
PROSTAGLAND	IN ANALOGS	
latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATIO	ON DRUGS	
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	brimonidine/timolol (generic Combigan) <sup>NR</sup> dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine)	
ОТН	IER	
RHOPRESSA (netarsudil) <sup>CL</sup> ROCKLATAN (netarsudil and latanoprost) <sup>CL</sup>		Drug-specific criteria:  Rhopressa and Rocklatan: Electronically approved for patients who have a trial of ONE generic agent, within ophthalmics - glaucoma within 60 days

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#### **OPIOID DEPENDENCE TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
buprenorphine SL buprenorphine/naloxone TAB (SL) SUBOXONE FILM (buprenorphine/naloxone)	buprenorphine/naloxone <b>FILM</b> LUCEMYRA (lofexidine) <sup>CL,QL</sup> ZUBSOLV (buprenorphine/naloxone)	Buprenorphine PA Form Buprenorphine Informed Consent  Non-Preferred buprenorphine and buprenorphine /naloxone agents:  Diagnosis of Opioid Use Disorder, NOT approved for pain management Verification of "X" DEA license number of prescriber No concomitant opioids Failed trial of preferred drug or patient-specific documentation of why preferred product not appropiriate for patient  Drug-specific criteria: Lucemyra: Approved for FDA approved indication and dosing per label. Trial of preferred product not required.

#### **OPIOID-REVERSAL TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naloxone SYRINGE, VIAL naltrexone TABLET NARCAN (naloxone) SPRAY	KLOXXADO (naloxone) <sup>NR</sup> <b>NASAL</b> naloxone <b>SPRAY</b> (generic for Narcan) <sup>NR</sup> ZIMHI (naloxone) <sup>AL,NR</sup> <b>SYRINGE</b>	<ul> <li>Non-preferred agents will be approved with documentation of why preferred products within this drug class are not appropriate for the patient</li> </ul>

#### **OTIC ANTI-INFECTIVES & ANESTHETICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acetic acid (generic for Vosol)	acetic acid/hydrocortisone (generic for Vosol HC)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of the preferred agent within this drug class</li> </ul>

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#### **OTIC ANTIBIOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CIPRODEX (ciprofloxacin/dexamethasone) neomycin/polymyxin/hydrocortisone (generic for Cortisporin) ofloxacin (generic for Floxin)	CIPRO HC (ciprofloxacin/ hydrocortisone) ciprofloxacin ciprofloxacin/dexamethasone (generic for CIPRODEX) COLY-MYCIN S(neomycin/ hydrocortisone/colistin) OTOVEL (ciprofloxacin/fluocinolone)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

# PAH (PUI MONARY ARTERIAL HYPERTENSION AGENTS) ORAL AND INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ambrisentan (generic Letairis) sildenafil (generic Revatio) <sup>CL</sup> TABLET tadalafil (generic for Adcirca) <sup>CL</sup> TRACLEER (bosentan) TABLET TYVASO (treprostinil) INHALATION VENTAVIS (iloprost) INHALATION	ADEMPAS (riociguat) <sup>CL</sup> ADCIRCA (tadalafil) <sup>CL</sup> bosentan (generic Tracleer) <b>TABLET</b> LETAIRIS (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) REVATIO (sildenafil) <sup>CL</sup> <b>SUSP</b> , <b>TABLET</b> sildenafil (generic Revatio) <sup>CL</sup> <b>SUSPENSION</b> TRACLEER (bosentan) <b>TABLETS FOR SUSPENSION</b> UPTRAVI (selexipag)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> <li>Drug-specific criteria:         <ul> <li>Adcirca®/Revatio®: Approved for diagnosis of Pulmonary Arterial Hypertension (PAH)</li> <li>Adempas®:</li></ul></li></ul>

#### **PANCREATIC ENZYMES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CREON ZENPEP (pancrelipase)	PANCREAZE (pancrelipase) PERTZYE (pancrelipase) VIOKACE (pancrelipase)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul>

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

NR – Product was not reviewed - New Drug criteria will apply

## with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

#### PEDIATRIC VITAMIN PREPARATIONS

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#### **PENICILLINS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
amoxicillin CAPSULE, CHEWABLE TABLET, SUSP, TABLET ampicillin CAPSULE dicloxacillin penicillin VK		<ul> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class</li> </ul>

#### **PHOSPHATE BINDERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcium acetate <b>TABLET</b> , <b>CAPSULE</b> CALPHRON OTC (calcium acetate) RENVELA (sevelamer carbonate)	AURYXIA (ferric citrate) ELIPHOS (calcium acetate) lanthanum (generic FOSRENOL) PHOSLO (calcium acetate) PHOSLYRA (calcium acetate) RENAGEL (sevelamer HCI) sevelamer HCI (generic Renagel) sevelamer carbonate (generic Renvela) VELPHORO (sucroferric oxyhydroxide)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> </ul>

#### **PLATELET AGGREGATION INHIBITORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AGGRENOX (dipyridamole/aspirin) aspirin BRILINTA (ticagrelor) clopidogrel (generic Plavix) dipyridamole (generic Persantine) prasugrel (generic Effient)	aspirin/dipyridamole (generic Aggrenox) ticlopidine (generic Ticlid) YOSPRALA (aspirin/omeprazole) ZONTIVITY (vorapaxar) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class OR documented clopidogrel resistance</li> <li>Drug-specific criteria:</li> <li>Zontivity®: Approved for reduction of thrombotic cardiovascular events in history of MI or with peripheral artery disease (PAD) Use with aspirin and/or clopidogrel</li> </ul>

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#### **PRENATAL VITAMINS**

Additional covered agents can be looked up using the Drug Look-up Tool at: https://druglookup.fhsc.com/druglookupweb/?client=nestate

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
c-nate dha SOFTGEL complete natal dha (pnv2/iron b-g suc-p/fa/omega-3) classic prenatal TABLET (prenatal vit/fe fum/fa) COMPLETENATE CHEWABLE pnv with ca, #72/iron/fa prenatal vitamin TABLET (pnv#124/iron/fa) prenatal no.137/iron/fa OTC pretab 29mg-1 TABLET (pnv#78/iron/fa) PUREFE PLUS PUREFE OB PLUS TRINATAL RX 1 virt-nate dha SOFTGEL (pnv 11-iron fum-fa-om3) zatean-pn plus SOFTGEL (pnv/ca no.68/iron/fa1/dha)	DERMACINRX PRENATRIX CAPLET (prenatal vit no. 170/fe/fa) DERMACINRX PRENATRYL CAPLET (prenatal vit no.170/fe/fa) DERMACINRX PRETRATE CAPLET (prenatal vit no. 170/fe/fa) folivane-ob CAPSULE (pnv#15/iron fum & ps cmp/fa) niva-plus TABLET (pnv with ca,no.74/iron/fa) pnv-dha SOFTGEL (pnv combo#47/iron/fa #1/dha) taron-c dha CAPSULE (pnv#16/iron fum &ps/fa/om-3) virt-c dha SOFTGEL (pnv dha SOFTGEL (pnv combo#47/iron/fa #1/dha) WESTGEL DHA (prenatal 93/iron/folate 9/dha) zatean-pn dha CAPSULE (pnv #47/iron/fa #1/dha)	Non-preferred agents will be approved for patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class

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### PROGESTERONE (hydroxyprogesterone caproate)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MAKENA <b>AUTO INJECTOR</b> (hydroxyprogesterone caproate)	hydroxyprogesterone caproate (generic Makena) MAKENA (hydroxyprogesterone caproate) <b>SDV</b>	<ul> <li>When filled as outpatient prescription, use limited to:         <ul> <li>Singleton pregnancy AND</li> <li>Previous Pre-term delivery AND</li> <li>No more than 20 doses (administered between 16 -36 weeks gestation)</li> <li>Maximum of 30 days per dispensing</li> </ul> </li> </ul>

#### PROTON PUMP INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
omeprazole (generic Prilosec) <b>RX</b> pantoprazole (generic Protonix) <sup>QL</sup> PROTONIX <b>SUSP</b> (pantoprazole)	DEXILANT (dexlansoprazole) dexlansoprazole (generic Dexilant) <sup>NR</sup> esomeprazole magnesium (generic Nexium) RX, OTC <sup>NR, QL</sup> esomeprazole strontium lansoprazole (generic Prevacid) <sup>QL</sup> NEXIUM SUSPENSION (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) pantoprazole GRANULES QL rabeprazole (generic Aciphex)	<ul> <li>Non-preferred agents will be approved for patients who have failed an 8-week trial of BOTH preferred agents within this drug class</li> <li>Pediatric Patients:         <ul> <li>Patients ≤ 4 years of age – No PA required for Prevacid 30mg or omeprazole 20mg capsules (used to compound suspensions).</li> </ul> </li> <li>Drug-specific criteria:         <ul> <li>Prilosec®OTC/Omeprazole OTC EXCLUDED from coverage Acceptable as trial instead of Omeprazole 20mg</li> <li>Prevacid Solutab: may be approved after trial of compounde suspension.</li> <li>Patients ≥ 5 years if age- Only approve non-preferred for Gl diagnosis if:</li></ul></li></ul>

# with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

#### **SEDATIVE HYPNOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
temazepam 15mg, 30mg (generic for Restoril)	estazolam (generic for ProSom) flurazepam (generic for Dalmane) temazepam (generic for Restoril) 7.5mg, 22.5mg triazolam (generic for Halcion)  IERS  BELSOMRA (suvorexant) <sup>AL,QL</sup> DAYVIGO (lemborexant) <sup>ALQL</sup> doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopiclone (generic for Lunesta) HETLIOZ (tasimelteon) SUSP AL,QL ramelteon (generic for Rozerem) zolpidem ER (generic for Ambien CR) zolpidem SL (generic for Intermezzo)	■ Lunesta®/ Rozerem®/zolpidem ER: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used ■ Edluar®: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used and Requires documentation of swallowing disorder ■ flurazepam/triazolam: Requires trial of preferred benzodiazepine ■ Hetlioz®: Requires trial with generic zolpidem within last 12 months AND clinical reason why zaleplon AND preferred benzodiazepine cannot be used ■ Silenor®: Must meet ONE of the following:
		2350 4054

PDL Updated May 2, 2022 Highlights indicated change from previous posting SICKLE CELL ANEMIA TREATMENT<sup>AL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DROXIA (hydroxyurea)	ENDARI (L-glutamine) <sup>CL</sup> OXBRYTA (voxelotor) <sup>CL</sup> SIKLOS (hydroxyurea)	<ul> <li>Endari: Patient must have documented two or more hospital admissions per year due to sickle cell crisis despite maximum hydroxyurea dosage.</li> <li>Oxbryta: Not inidcated for sickle cell crisis. Patient must have had at least one sickle cell-related vaso-occlusive event within the past 12 months; AND baseline hemoglobin is 5.5 g/dL ≤ 10.5 g/dL; AND patient is not receiving concomitant, prophylactic blood tranfusion therapy</li> <li>Siklos: Approved for use in patients ages 2 to 17 years old</li> </ul>

#### **SINUS NODE INHIBITORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORLANOR <b>SOLUTION</b> , <b>TABLET</b> (ivabradine)	<ul> <li>Diagnosis of Chronic Heart Failure with left ventricular ejection fraction less than or equal to 35%, AND</li> <li>Sinus rhythm with resting heart rate greater than or equal to 70 beats per minute, AND</li> <li>On maximally tolerated doses of beta-blockers OR have a contraindication to beta-blocker use</li> </ul>

## with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

#### **SKELETAL MUSCLE RELAXANTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
baclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) methocarbamol (generic Robaxin) tizanidine TABLET (generic Zanaflex)	baclofen (generic for Ozobax) <sup>NR,QL</sup> SOLN carisoprodol (generic Soma) <sup>CL,QL</sup> carisoprodol compound cyclobenzaprine ER (generic     Amrix) <sup>CL</sup> dantrolene (generic Dantrium) FEXMID (cyclobenzaprine ER) FLEQSUVY (baclofen) <sup>NR</sup> SUSP LORZONE (chlorzoxazone) <sup>CL</sup> metaxalone (generic Skelaxin) NORGESIC FORTE     (orphenadrine/ASA/caffeine) orphenadrine ER PARAFON FORTE (chlorzoxazone) tizanidine CAPSULE ZANAFLEX (tizanidine) CAPSULE,     TABLET	<ul> <li>Non-preferred agents will be approved for patients who have failed a 1-week trial of TWO preferred agents within this drug class</li> <li>Drug-specific criteria:         <ul> <li>cyclobenzaprine ER:</li> <li>Requires clinical reason why IR cannot be used</li> <li>Approved only for acute muscle spasms</li> <li>NOT approved for chronic use</li> </ul> </li> <li>carisoprodol:         <ul> <li>Approved for Acute, musculoskeletal pain - NOT for chronic pain</li> <li>Use is limited to no more than 30 days</li> <li>Additional authorizations will not be granted for at least 6 months following the last dayy of previous course of therapy</li> </ul> </li> <li>Dantrolene: Trial NOT required for treatment of spasticity from spinal cord injury</li> <li>Lorzone®: Requires clinical reason why chlorzoxazone cannot be used</li> <li>Soma® 250mg: Requires clinical reason why 350mg generic strength cannot be used</li> <li>Zanaflex® Capsules: Requires clinical reason generic cannot be used</li> </ul>

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#### STEROIDS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
LOW POTENCY		Low Potency Non-preferred agents
hydrocortisone OTC & RX CREAM, LOTION, OINTMENT (Rx only) hydrocortisone/aloe OINTMENT SCALPICIN OTC (hydrocortisone)	ALA-CORT (hydrocortisone) CREAM ALA-SCALP HP (hydrocortisone) alclometasone dipropionate (generic for Aclovate) CAPEX SHAMPOO (fluocinolone) DESONATE (desonide) GEL desonide LOTION (generic for Desowen) desonide CREAM, OINTMENT   (generic for former products Desowen, Tridesilon) fluocinolone 0.01% OIL (generic for DERMA-SMOOTHE-FS) hydrocortisone/aloe CREAM hydrocortisone OTC OINTMENT MICORT-HC (hydrocortisone)	will be approved for patients who have failed a trial of ONE preferred agent within this drug class
MEDIIM	POTENCY •	Medium Potency Non-preferred
fluticasone propionate CREAM,    OINTMENT (generic for Cutivate) mometasone furoate CREAM,    OINTMENT, SOLUTION (generic for Elocon)	betamethasone valerate (generic for Luxiq) clocortolone (generic for Cloderm) fluocinolone acetonide (generic for Synalar) flurandrenolide (generic for Cordran) fluticasone propionate LOTION   (generic for Cutivate) hydrocortisone butyrate (generic for Locoid) hydrocortisone butyrate/emoll (generic for Locoid Lipocream) hydrocortisone valerate (generic for Westcort) PANDEL (hydrocortisone probutate 0.1%) prednicarbate (generic for Dermatop)	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class

## with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

## **STEROIDS, TOPICAL (Continued)**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HIGH POTENCY		High Potency Non-preferred
triamcinolone acetonide <b>OINTMENT</b> , <b>CREAM</b>	amcinonide CREAM, LOTION, OINTMENT	agents will be approved for patients who have failed a trial of TWO preferred agents within this
triamcinolone <b>LOTION</b>	betamethasone dipropionate betamethasone / propylene glycol	drug class
	betamethasone valerate	
	desoximetasone	
	diflorasone diacetate fluocinonide <b>SOLUTION</b>	
	fluocinonide CREAM, GEL, OINTMENT	
	fluocinonide emollient	
	halcinonide <b>CREAM</b> (generic for Halog)	
	HALOG (halcinonide) CREAM, OINT, SOLN	
	KENALOG AEROSOL (triamcinolone)	
	SERNIVO (betamethasone	
	dipropionate) triamcinolone <b>SPRAY</b> (generic for	
	Kenalog spray)	
	TRIANEX <b>OINTMENT</b> (triamcinolone)	
	VANOS (fluocinonide)	
VERY HIG	H POTENCY	vory riigiri otorioy rion protonou
clobetasol emollient (generic for	APEXICON-E (diflorasone)	agents will be approved for patients who have failed a trial of
Temovate-E)	BRYHALI (halobetasol prop) <b>LOTION</b>	TWO preferred agents within this
clobetasol propionate CREAM, GEL, OINTMENT, SOLUTION	clobetasol SHAMPOO, LOTION	drug class
halobetasol propionate (generic for	clobetasol propionate FOAM, SPRAY	
Ultravate)	CLOBEX (clobetasol)	
	halobetasol propionate <b>FOAM</b> (generic for Lexette) AL,QL	
	IMPEKLO (clobetasol) LOTION <sup>AL</sup>	
	LEXETTE(halobetasol propionate) AL,QL	
	OLUX-E /OLUX/OLUX-E CP (clobetasol)	

## with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting STIMULANTS AND RELATED AGENTS<sup>AL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CNS STIMULANTS		Non-preferred agents will be
Amphetamine type		approved for patients who have failed a trial of ONE preferred
ADDERALL XR (amphetamine salt combo) amphetamine salt combination IR VYVANSE (lisdexamfetamine) <sup>QL</sup> CAPSULE, CHEWABLE	ADZENYS XR (amphetamine) amphetamine ER (generic for Adzenys ER) SUSPENSION amphetamine salt combination ER   (generic for Adderall XR) amphetamine sulfate (generic for Evekeo) AZSTARYS (serdexmethylphenidate and dexmethylphenidate) dextroamphetamine (generic for Dexedrine) dextroamphetamine SOLUTION   (generic for Procentra) dextroamphetamine ER (generic for Dexedrine ER) DYANAVEL XR (amphetamine) EVEKEO ODT (amphetamine sulfate) MYDAYIS (amphetamine salt combo) QL methamphetamine (generic for Desoxyn) ZENZEDI (dextroamphetamine)	agent within this drug class  Drug-specific criteria:  Procentra®: May be approved with documentation of swallowing disorder  Zenzedi®: Requires clinical reason generic dextroamphetamine IR cannot be used

## with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

# STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Methylph CONCERTA (methylphenidate ER) <sup>QL</sup> 18mg, 27mg, 36mg, 54mg	ADHANSIA XR (methylphenidate) QL APTENSIO XR (methylphenidate) COTEMPLA XR-ODT	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> <li>Maximum accumulated dose of</li> </ul>
dexmethylphenidate (generic for Focalin IR) FOCALIN XR (dexmethylphenidate) METHYLIN <b>SOLUTION</b> (methylphenidate) methylphenidate (generic for Ritalin)	(methylphenidate) <sup>QL</sup> DAYTRANA <b>PATCH</b> (methylphenidate) <sup>QL</sup> dexmethylphenidate XR (generic for Focalin XR) FOCALIN IR (dexmethylphenidate)	<ul> <li>108mg per day for ages &lt; 18</li> <li>Maximum accumulated dose of 72mg per day for ages &gt; 19</li> <li>Drug-specific criteria:</li> <li>Daytrana®: May be approved in history of substance use disorder</li> </ul>
methylphenidate <b>SOLUTION</b> (generic for Methylin)  QUILLICHEW ER <b>CHEWTAB</b> (methylphenidate)	JORNAY PM (methylphenidate) QL methylphenidate 50/50 (generic for Ritalin LA) methylphenidate 30/70 (generic for Metadate CD) methylphenidate ER 18mg, 27mg,	by parent, caregiver, or patient.  May be approved with documentation of difficulty swallowing
	methylphenidate ER Tonig, 27mg, 36mg, 54mg (generic Concerta) <sup>QL</sup> methylphenidate ER <b>CAP</b> (generic for Aptensio XR) <sup>QL</sup> Methylphenidate ER (generic for Metadate ER)	
	methylphenidate ER 72mg (generic for RELEXXII) <sup>QL</sup> methylphenidate ER (generic for Ritalin SR)	
	QUILLIVANT XR <b>SUSP</b> (methylphenidate) RITALIN (methylphenidate)	

### with Prior Authorization Criteria

PDL Updated May 2, 2022 Highlights indicated change from previous posting

# STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Non-Preferred Agents	Prior Authorization/Class Criteria
ANEOUS	Note: generic guanfacine IR and —clonidine IR are available without
clonidine ER (generic for Kapvay) <sup>QL</sup> QELBREE (viloxazine) <sup>QL</sup> STRATTERA (atomoxetine)	prior authorization
	Drug-specific criteria:  armodafinil and Sunosi: Require trial of modafinil
	armodafinil and modafinil:
modafanil (generic for Provigil) <sup>CL</sup> SUNOSI (solriamfetol) <sup>CL,QL</sup> WAKIX (pitolisant) <sup>CL,QL</sup>	approved only for:  Sleep Apnea with documentation/confirmation via sleep study and documentation that C-PAP has been maxed  Narcolepsy with documentation of diagnosis via sleep study  Shift Work Sleep Disorder (only approvable for 6 months) with work schedule verified and documented. Shift work is defined as working the all night shift  Sunosi approved only for:  Sleep Apnea with documentation/confirmation via sleep study and documentation that C-PAP has been maxed  Narcolepsy with documentation of diagnosis via sleep study  Wakix: approved only for excessive daytime sleepiness in adults with narcolepsy with documentation of narcolepsy diagnosis via sleep study
	clonidine ER (generic for Kapvay) <sup>QL</sup> QELBREE (viloxazine) <sup>QL</sup> STRATTERA (atomoxetine)  EPTICS  armodafinil (generic for Nuvigil) <sup>CL</sup> modafanil (generic for Provigil) <sup>CL</sup> SUNOSI (solriamfetol) <sup>CL,QL</sup>

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.  $^{CL}$  – Prior Authorization / Class Criteria apply  $^{QL}$  – Quantity/Duration Limit  $^{AL}$  – Age Limit

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#### **TETRACYCLINES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR (generic Vibramycin) doxycycline monohydrate 50MG, 100MG CAPSULE doxycycline monohydrate SUSP, TABLET (generic Vibramycin) minocycline HCI CAPSULE, TABLET (generic Dynacin/ Minocin/Myrac)	demeclocycline (generic Declomycin) <sup>CL</sup> DORYX MPC DR (doxycycline pelletized) doxycycline hyclate DR (generic Doryx) doxycycline monohydrate 40MG, 75MG and 150MG CAPSULES (generic for Adoxa/Monodox/ Oracea) minocycline HCI ER (generic Solodyn) NUZYRA (omadacycline) tetracycline VIBRAMYCIN SUSP (doxycycline) XIMINO (minocycline ER) <sup>QL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of TWO preferred agents within this drug class</li> <li>Drug-specific criteria:         <ul> <li>Demeclocycline: Approved for diagnosis of SIADH</li> <li>Doryx®/doxycycline hyclate DR/Dynacin®/Oracea®/Solodyn®: Requires clinical reason why generic doxycycline, minocycline or tetracycline cannot be used</li> <li>doxycycline suspension: May be approved with documented swallowing difficulty</li> </ul> </li> </ul>

#### THROMBOPOIESIS STIMULATING PROTEINSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROMACTA (eltrombopag) <b>TABLET</b> <sup>CL</sup>	DOPTELET (avatrombopag) MULPLETA (lusutrombopag) PROMACTA (eltrombopag) <b>SUSP</b> TAVALISSE (fostamatinib)	<ul> <li>All agents will be approved with FDA-approved indication, ICD-10 code is required.</li> <li>Non-preferred agents require a trial of a preferred agent with the same indication or a contraindication.</li> <li>Drug-Specific Criteria</li> <li>Doptelet/Mulpleta: Approved for one course of therapy for a scheduled procedure with a risk of bleeding for treatment of thrombocytopenia in adult patients with chronic liver disease</li> </ul>

# PDL Updated May 2, 2022 Highlights indicated change from previous posting

#### **THYROID HORMONES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
levothyroxine <b>TABLET</b> (generic Synthroid) liothyronine <b>TABLET</b> (generic Cytomel) thyroid, pork <b>TABLET</b> UNITHROID (levothyroxine)	EUTHYROX (levothyroxine) LEVO-T (levothyroxine) levothyroxine CAPSULE (generic for Tirosint) THYROLAR TABLET (liotrix) THYQUIDITY (levothyroxine) SOLN TIROSINT CAPSULE (levothyroxine) TIROSINT-SOL LIQUID (levothyroxine) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Tirosint-Sol: May be approved with documented swallowing difficulty</li> </ul>

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#### **ULCERATIVE COLITIS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORAL		Non-preferred agents will be
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine) LIALDA (mesalamine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) GIAZO (balsalazide) mesalamine ER (generic Apriso) mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	approved for patients who have failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:  Asacol HD®/Delzicol DR®/Pentasa®: Requires clinical reason why preferred mesalamine products cannot be used  Giazo®: Requires clinical reason why generic balsalazide cannot be used
RECTAL		NOT covered in females
CANASA (mesalamine) ROWASA (mesalamine)	mesalamine <b>ENEMA</b> (generic Rowasa) mesalamine <b>SUPPOSITORY</b> (generic Canasa) UCERIS (budesonide)	

#### **UTERINE DISORDER TREATMENT**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORIAHNN (elagolix/ estradiol/ norethindrone) <sup>AL,CL</sup> ORILISSA (elagolix sodium) <sup>QL,CL</sup>	MYFEMBREE (relugolix/ estradiol/ norethindrone acetate) <sup>AL, NR, QL</sup>	Orilissa/Oriahnn: Requires an FDA approved indication, must follow FDA dosing guidelines, and have had a trial and failure of an NSAID and oral contraceptive     Total duration of treatment is max of 24 months

PDL Updated May 2, 2022 Highlights indicated change from previous posting

## **VASODILATORS, CORONARY**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate TABLET isosorbide dinitrate ER, SA TABLET (generic Dilatrate-SR/Isordil) isosorbide mono IR/SR TABLET nitroglycerin SUBLINGUAL, TRANSDERMAL nitroglycerin ER TABLET	BIDIL (isosorbide dinitrate/ hydralazine) <sup>CL</sup> GONITRO (nitroglycerin) isosorbide dinitrate TABLET (Oceanside Pharm MFR only) isosorbide dinitrate/hydralazine (Bidil) <sup>CL,NR</sup> NITRO-BID OINTMENT (nitroglycerin) NITRO-DUR (nitroglycerin) nitroglycerin TRANSLINGUAL (generic Nitrolingual) VERQUVO (vericiguat) <sup>AL,CL,QL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>BiDil: Approved for the treatment of heart failure as an adjunct therapy to standard therapy in self-identified black patients</li> <li>Verquvo: Approved for use in patients following a recent hospitalization for HF within the past 6 months OR need for outpatient IV diuretics, in adults with symptomatic chronic HF and EF less than 45%</li> </ul>