



**DEPT. OF HEALTH AND HUMAN SERVICES** 

# Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

Entire December 2022 PDL with November P&T Changes Noted in Red Font that Become Effective January 20, 2023

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

### Nebraska Medicaid Preferred Drug List

### with Prior Authorization Criteria

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

#### ACNE AGENTS, TOPICAL

ACNE AGENTS, TOPICAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benzoyl peroxide (BPO) WASH, LOTION clindamycin/BPO (generic Benzaclin) PUMP clindamycin phosphate PLEDGET clindamycin phosphate SOLUTION DIFFERIN LOTION, CREAM, Rx-GEL (adapalene) DIFFERIN GEL (adapalene) OTC erythromycin GEL erythromycin SOLN 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)AL ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) AZELEX (azelaic acid) BENZACLIN PUMP (clindamycin/BPO) BENZEFOAM (benzoyl peroxide) benzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide GEL OTC benzoyl peroxide GEL Rx benzoyl peroxide GEL Rx benzoyl peroxide GEL Rx benzoyl peroxide GEL Rx benzoyl peroxide GEL CITC clindamycin FOAM, LOTION clindamycin FOAM, LOTION clindamycin phosphate (generic for Clindagel) GEL clindamycin/BPO (generic Acanya) GEL clindamycin/BPO (generic Duac) clindamycin/tretinoin (generic Veltin, Ziana) dapsone (generic Aczone) EPIDUO FORTE GEL PUMP (adapalene/BPO) erythromycin GEL, PLEDGET erythromycin-BPO (generic for Benzamycin) EVOCLIN (clindamycin) FABIOR (tazarotene foam) NEUAC (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONACE PLUS (sulfacetamide sodium) PLIXDA (adapalene) SWAB RETIN-A <sup>AL</sup> GEL, CREAM (tretinoin) sulfacetamide sulfacetamide/sulfur SUMADAN (sulfacetamide/sulfur) tazarotene CREAM,GEL <sup>NR</sup> (generic Tazorac) tazarotene FOAM (generic Fabior) TRETIN-X (tretinoin) tretinoin microspheres (generic for Retin-A Micro) AL TWYNEO (tretinoin/BPO) AL. NR CREAM	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

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
donepezil (generic Aricept) donepezil ODT (generic Aricept ODT) rivastigmine PATCH (generic for Exelon Patch)	ASE INHIBITORS  ADLARITY (donepezil) PATCH ARICEPT (donepezil) donepezil 23 (generic Aricept 23)  EXELON (rivastigmine) CAPS, PATCH galantamine (generic Razadyne) SOLN, TAB galantamine ER (generic Razadyne ER) rivastigmine CAPS (generic Exelon)	<ul> <li>Non-preferred agents will be approved for patients who have failed a 120-day trial of ONE preferred agent within this drug class within the last 6 months         OR     </li> <li>Current, stabilized therapy of the non-preferred agent within the previous 45 days</li> </ul>
NMDA RECEPT memantine (generic for Namenda)	OR ANTAGONIST  memantine ER (generic for Namenda XR) memantine SOLN (generic for Namenda)  NAMENDA (memantine)  NAMZARIC (memantine/donepezil)	<ul> <li>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)</li> </ul>

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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>	BELBUCA (buprenorphine)QL BUCCAL buprenorphine BUCCAL (generic for Belbuca)AL,QL buprenorphine PATCH (generic Butrans)QL EMBEDA (morphine sulfate/naltrexone) DURAGESIC MATRIX (fentanyI)QL fentanyI 37.5, 62.5, 87.5 mcg PATCHQL hydrocodone ER (generic for Hysingla ER)QL hydrocodone bitartrate ER (generic for Zohydro ER) hydromorphone ER (generic for Exalgo)CL HYSINGLA ER (hydrocodone ER) KADIAN (morphine ER) methadone TABLETCL methadone ORAL SYRCL MORPHABOND ER (morphine sulfate) morphine ER (generic for Avinza, Kadian) CAPS NUCYNTA ER (tapentadol)CL oxycodone ER (generic Oxycontin) oxymorphone ER (generic Opana ER) tramadol ER (generic Conzip)CL	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

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#### ANALGESICS, OPIOID SHORT-ACTINGQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acetaminophen/codeine ELIXIR, TABLET codeine TABLET hydrocodone/APAP SOLN, TABLET hydrocodone/ibuprofen hydromorphone TABLET morphine CONC SOLN, SOLN, TABLET bxycodone TABLET, SOLN bxycodone/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/aspirin/caffeine FIORINAL/CODEINE (butalbital/ ASA/codeine/caffeine) hydromorphone LIQUID, SUPPOSITORY (generic Dilaudid) IBUDONE (hydrocodone/ibuprofen) levorphanol meperidine (generic Demerol) morphine SUPPOSITORIES NALOCET (oxycodone/APAP) NUCYNTA (tapentadol) <sup>CL</sup> oxycodone/APAP SOLN oxycodone/aspirin	<ul> <li>Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the lass 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)	
		-D
BUCCAL/TRANSMUCOSAL <sup>CL</sup>		Drug-specific criteria:  • Abstral®/Actiq®/Fentora®/
	ABSTRAL (fentanyl) <sup>CL</sup>	Onsolis (fentanyl): Approved only
	fentanyl <b>TRANSMUCOSAL</b> (generic	for diagnosis of cancer AND
	Actiq) <sup>CL</sup>	current use of long-acting opiate
	FENTORA (fentanyl) <sup>CL</sup>	

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NDROGEL (testosterone) PUMP CL	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®:</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) lisinopril (generic Prinivil, Zestril) quinapril (generic Accupril) ramipril (generic Altace)	captopril (generic Capoten) EPANED (enalapril) <sup>CL</sup> ORAL SOLN enalapril (generic for Epaned) <sup>CL</sup> ORAL SOLN fosinopril (generic Monopril) moexepril (generic Univasc) perindopril (generic Aceon) QBRELIS (lisinopril) <sup>CL</sup> ORAL SOLN trandolapril (generic Mavik)  EETIC COMBINATIONS captopril/HCTZ (generic Capozide) fosinopril/HCTZ (generic Monopril HCT) 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 RECEPTOR 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 BLOC	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/	- Angiotensin Modulator/Calcium Channel Blocker Combinations:
CALCIUM CHANNEL BL	OCKER COMBINATIONS	Combination agents may be
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 Renin Inhibitors/Direct Renin Inhibitor Combinations:
DIRECT RENI	N INHIBITORS	<ul> <li>May be approved with history of</li> </ul>
	aliskiren (generic Tekturna) <sup>QL</sup>	TWO preferred ACE Inhibitors or Angiotensin Receptor Blockers
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)QL		with a diagnosis of fleart failure
ANGIOTENSIN RECEPTOR BLOCKER/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)	<ul> <li>Drug-specific criteria:         <ul> <li>ORALAIR</li> <li>Confirmed by positive skin to or in vitro testing for pollenspecific IgE antibodies for Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass Mixed Pollens.</li> <li>For use in patients 5 through 65 years of age.</li> </ul> </li> <li>PALFORZIA</li> <li>Confirmed diagnosis of pear allergy by allergist</li> <li>For use in patients ages 4 to 17; it may be continued in patients 18 years and older with documentation of previouse within the past 90 days</li> <li>Initial dose and increase titration doses should be given in a healthcare setting</li> <li>Should not be used in patient with uncontrolled asthma or concurrently on a NSAID</li> </ul>

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#### **ANTIBIOTICS, GASTROINTESTINAL**

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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> SUSP CAYSTON (aztreonam lysine) <sup>QL,CL</sup> tobramycin (generic Bethkis) tobramycin (generic Tobi) <sup>CL</sup>	<ul> <li>Diagnosis of Cystic Fibrosis is required for all agents</li> <li>ICD10 Group = E84, ICD9 = 277.00, 277.01, 277.02, 277.03, 277.09</li> </ul>
		<ul> <li>Arikayce: Requires diagnosis of refractory MAC lung disease defined as patients who did not achieve negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy</li> <li>Cayston®: Trial of tobramycin via nebulizer and demonstration of TOBI® compliance required</li> <li>Tobi Podhaler®: Requires trial of tobramycin via nebulizer or documentation why nebulized tobramycin cannot be used</li> </ul>

#### ANTIBIOTICS TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
bacitracin <b>OINT</b> bacitracin/polymyxin (generic Polysporin) mupirocin <b>OINT</b> (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/ pramoxine	CENTANY (mupirocin) gentamicin OINT, 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) metronidazole, vaginal NUVESSA (metronidazole)	CLEOCIN <b>CREAM</b> (clindamycin) METROGEL (metronidazole) VANDAZOLE (metronidazole) XACIATO (clindamycin phosphate) vaginal gel <sup>AL,NR</sup>	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

#### **ANTICOAGULANTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) noxaparin (generic Lovenox) PRADAXA (dabigatran) varfarin (generic Coumadin) CARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg CARELTO (rivaroxaban) 2.5 mg <sup>CL,QL</sup> CARELTO DOSE PACK (rivaroxaban)	BEVYXXA (betrixaban) <sup>QL</sup> dabigatran etexilate <sup>NR</sup> (generic Pradaxa) fondaparinux (generic Arixtra) FRAGMIN (dalteparin) SAVAYSA (edoxaban) <sup>QL</sup> XARELTO (rivaroxaban) <sup>CL</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:         <ul> <li>Stroke and systemic embolism (SE) risk reduction in nonvalvular atrial fibrillation (NVAF) OR</li> <li>Treatment of deep vein thrombos (DVT) and pulmonary embolism (PE) following 5-10 days of parenteral anticoagulant therapy</li> </ul> </li> <li>Xarelto 2.5mg: Use limited to reduction of risk of major cardiovascular events (cardiovascular death, myocardia infarction, and stroke) in patients with chronic coronary artery disease or peripheral artery disease</li> <li>Xarelto Suspension: Approved for patients ≤12 years of age or if there is a clinical reason why a preferred solid dosage form cannibe used.</li> </ul>

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

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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) CAPS, CAPS PACK <sup>QL</sup>	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 SOLN (generic Emetrol) prochlorperazine, oral (generic Compazine) promethazine SYRUP, TABLET (generic Phenergan) promethazine SUPPOSITORY 12.5mg, 25mg TRANSDERM-SCOP (scopolamine)	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
troche) uconazole SUSP, TAB (generic Diflucan) riseofulvin SUSP riseofulvin microsized TAB ystatin SUSP, TAB erbinafine (generic Lamisil)	BREXAFEMME (ibrexafungerp) <sup>QL</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) NOXAFIL (posaconazole) POWDERMIX <sup>AL,NR</sup> nystatin POWDER ONMEL (itraconazole) posaconazole (generic Noxafil) <sup>AL,CL</sup> TOLSURA (itraconazole) <sup>CL</sup> VIVJOA (oteseconazole) CAPS <sup>NR</sup> voriconazole (generic VFEND) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approve 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 invasive aspergillosis or invasive mucormycosis</li> <li>Flucytosine: Approved for diagnosis of:         <ul> <li>Candida: Septicemia, endocarditis, UTIs</li> <li>Cryptococcus: Meningitis, pulmonary infections</li> <li>Noxafil®: No trial for diagnosis of Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropeni hematologic malignancies, Graft vs. Host disease(GVHD), Immunosuppression secondary to hematopoietic stem cell transplant</li> <li>Noxafil® Suspension:</li></ul></li></ul>

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ANTIFUNGALS, TOPICAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole CREAM (generic Lotrimin) RX, OTC clotrimazole SOLN OTC ketoconazole CREAM, SHAMPOO (generic Nizoral) LAMISIL (terbinafine) SPRAY OTC LAMISIL AT CREAM (terbinafine) OTC miconazole CREAM, POWDER OTC nystatin terbinafine OTC (generic Lamisil AT) tolnaftate POWDER, CREAM, POWDER OTC (generic Tinactin)  ANTIFUNGAL/STEF clotrimazole/betamethasone CREAM (generic Lotrisone) nystatin/triamcinolone (generic Mycolog)	ALEVAZOL (clotrimazole) OTC ciclopirox CREAM, GEL, SUSP (generic Ciclodan, Loprox) ciclopirox NAIL LACQUERCL (generic Penlac) ciclopirox SHAMPOO (generic Loprox) clotrimazole SOLN RX (generic Lotrimin) DESENEX POWDER OTC (miconazole) econazole (generic Spectazole) ERTACZO (sertaconazole) EXELDERM (sulconazole) FUNGOID OTC JUBLIA (efinaconazole) <sup>CL</sup> ketoconazole FOAMCL (generic Extina, Ketodan) LAMISIL AT GEL, SPRAY (terbinafine) OTC LOPROX (ciclopirox) SUSP, 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 Densal HP) tavaborole SOLUTION <sup>CL</sup> (generic Kerydin) tolnaftate SPRAY, OTC	<ul> <li>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</li> <li>Extina: Requires trial and failure or contraindication to other ketoconazole forms</li> <li>Jublia and tavaborole:         Approved diagnoses include Onychomycosis of the toenails due to <i>T.rubrum OR T. Mentagrophytes</i> </li> <li>ciclopirox nail lacquer: No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine</li> </ul>
CREAM, OINT		

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
cetirizine TABLET, SOLN (Rx only) (generic Zyrtec) Ioratadine TABLET, SOLN (generic Claritin) Ievocetirizine TABLET (generic Xyzal)	cetirizine CHEWABLE (generic Zyrtec) cetirizine SOLN (OTC) desloratadine (generic Clarinex) desloratadine ODT (generic Clarinex Reditabs) fexofenadine (generic Allegra) fexofenadine 180mg (generic Allegra 180mg) <sup>QL</sup> levocetirizine (generic Xyzal) SOLN loratadine CAPS, CHEWABLE, ODT (generic 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>TAB</b> (generic for Catapres) clonidine <b>TRANSDERMAL</b> guanfacine (generic for Tenex) methyldopa	methyldopa/hydrochlorothiazide	<ul> <li>Non-preferred agents will be approved for patients who have failed a 30-day trial with ONE preferred agent within this drug class</li> </ul>

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#### **ANTIHYPERURICEMICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
allopurinol (generic Zyloprim) MITIGARE (colchicine) probenecid probenecid/colchicine (generic Col- Probenecid)	allopurinol <sup>NR</sup> 200mg colchicine <b>TAB</b> (generic Colcrys) <sup>CL</sup> colchicine <b>CAPS</b> (generic Mitigare) febuxostat (generic Uloric) <sup>CL</sup> GLOPERBA <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<sup>®</sup>: Approved without trial for familial Mediterranean fever OR pericarditis</li> <li>Gloperba: Approved for documented swallowing disorder</li> <li>Uloric<sup>®</sup>: Clinical reason why allopurinol cannot be used</li> </ul>

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#### **ANTIMIGRAINE AGENTS, OTHER**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AJOVY (fremanezumab-vfrm) CL, QL PEN, Autoinjector  AJOVY (fremanezumab-vfrm) Autoinjector 3-packCL,QL  EMGALITY 120 mg/mL (galcanezumab- gnlm) CL, QL PEN, SYRINGE  NURTEC ODT (rimegepant)AL,CL,QL  UBRELVY (ubrogepant)AL,CL, QL  TABLET	Almovig (erenumab-aooe) CL,QL CAFERGOT (ergotamine/caffeine) dihydroergotamine mesylate NASAL ELYXYB (celecoxib)AL,QL SOLN EMGALITY 100 mg (galcanezumabgnim) CL,QL SYRINGE ERGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL QULIPTA (atogepant)ALQL REYVOW (lasmiditan)AL, CL,QL TABLET TRUDHESA (dihydroergotamine mesylate)AL,QL NASAL	<ul> <li>All acute treatment agents will be approved for patients who have a failed trial or a contraindication to 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>Emgality 120mg is recommended for preventative treatment of Migraine, Emgaility 100mg is recommended for treatment of Episodic Cluster Headache</li> <li>For Prophylactic Treatment: 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> </ul>

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#### **ANTIMIGRAINE AGENTS, TRIPTANSQL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OF	RAL	Non-preferred agents will be
rizatriptan (generic Maxalt) rizatriptan ODT (generic Maxalt MLT) 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)	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
NA	SAL	
IMITREX (sumatriptan)	ONZETRA XSAIL (sumatriptan) sumatriptan (generic Imitrex Nasal) TOSYMRA (sumatriptan) zolmitriptan (generic for Zomig) ZOMIG (zolmitriptan)	
INJEC	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 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

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#### ANTIPARKINSON'S AGENTS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHO	INERGICS	Non-preferred agents will be
benztropine (generic for Cogentin)		approved for patients who have failed ONE preferred agents within
trihexyphenidyl (generic for Artane)  COMT IN	HIBITORS	this drug class
33	entacapone (generic for Comtan)	- Drug-specific criteria:
	ONGENTYS (opicapone) <sup>QL</sup> tolcapone (generic for Tasmar)	<ul> <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- - containing drug
pramipexole (generic for Mirapex) ropinirole (generic for Requip)	bromocriptine (generic Parlodel) ropinirole ER (generic Requip ER) <sup>CL</sup> NEUPRO (rotigotine) <sup>CL</sup> pramipexole ER (generic Mirapex ER) <sup>CL</sup> ropinirole ER (generic Requip XL) <sup>CL</sup>	<ul> <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	HIBITORS	For Parkinsons: Clinical reason required why preferred agent
selegiline CAPS, TABLET (generic Eldepryl)	rasagiline (generic 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
	KINSON'S DRUGS	Parkinson's disease and concurrent
amantadine CAPS, SYRUP TABLET (generic Symmetrel) carbidopa/levodopa (generic Sinemet) carbidopa/levodopa ER (generic Sinemet CR) levodopa/carbidopa/entacapone (generic Stalevo)	APOKYN (apomorphine) SUB-Q apomorphine (generic Apokyn) SUB-Q carbidopa (generic Lodosyn) carbidopa/levodopa ODT (generic Parcopa) DHIVY (carbidopa/levodopa) DUOPA (carbidopa/levodopa) GOCOVRI (amantadine) INBRIJA (levodopa) INHALER KYNMOBI (apomorphine) KYNMOBI (apomorphine) NOURIANZ (istradefylline) CL,QL NOURIANZ (istradefylline) CL,QL NOURIANZ (mantadine) 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)	<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
calcipotriene CREAM, OINT, SOLN	calcitriol (generic for Vectical) calcipotriene/betamethasone OINT	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> SUSP 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>OINT</b>	acyclovir CREAM, (generic Zovirax) DENAVIR (penciclovir) penciclovir (generic Denavir) <sup>NR</sup> 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>

#### ANYIOI VTICE

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>SOLN</b> (generic for Valium) orazepam <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> LOREEV XR (lorazepam) <sup>AL</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
atenolol (generic Tenormin) atenolol/chlorthalidone (generic Tenoretic) bisoprolol (generic Zebeta) bisoprolol/HCTZ (generic Ziac) metoprolol (generic Lopressor) metoprolol ER (generic Toprol XL) propranolol (generic Inderal) propranolol ER (generic Inderal LA)	acebutolol (generic Sectral) betaxolol (generic Kerlone) BYSTOLIC (nebivolol) HEMANGEOL (propranolol) SOLN INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) LEVATOL (penbutolol) metoprolol/HCTZ (generic Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide nebivolol (generic Bystolic) 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 Actigall) ursodiol 250mg <b>TABLET</b> (generic URSO) ursodiol 500mg <b>TABLET</b> (generic URSO FORTE)	BYLVAY (odevixibat)CAP, PELLET CHENODAL (chenodiol) CHOLBAM (cholic acid) LIVMARLI (maralixibat) SOLNAL OCALIVA (obeticholic acid) RELTONE (ursodiol 200mg,400mg) CAP	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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#### **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) fesoterodine <sup>NR</sup> (generic Toviaz) flavoxate GELNIQUE (oxybutynin) GEMTESA (vibegron) <sup>AL,QL</sup> MYRBETRIQ <b>TABLET</b> , <b>SUSP</b> <sup>AL,CL,QL</sup> (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) <sup>AL</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>Drug-specific criteria:</li> <li>Myrbetriq®: Covered without trial in contraindication to anticholinergic agents</li> <li>Myrbetriq suspension: Covered for pediatric patients ≥ 3 years old with a diagnosis of Neurogenic Detrusor Overactivity (NDO)</li> </ul>

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BISPHOSPHONATES		Non-preferred agents will be
BISPHOSI alendronate (generic Fosamax) TAB ibandronate (generic Boniva) <sup>QL</sup>	PHONATES  alendronate SOLN (generic Fosamax) <sup>QL</sup> ATELVIA DR (risedronate)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within the same group  Drug-specific criteria:

## **Nebraska Medicaid Preferred Drug List**

### 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>

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROVENTIL HFA (albuterol) VENTOLIN HFA (albuterol)	albuterol HFA (generic ProAir HFA, Proventil HFA, and Ventolin HFA) levalbuterol HFA (generic Xopenex HFA) PROAIR DIGIHALER (albuterol) PROAIR RESPICLICK (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>
INHA	LERS – Long Acting	
SEREVENT (salmeterol)	STRIVERDI RESPIMAT (olodaterol)	
INHA	ALATION SOLUTION	
albuterol (2.5mg/3ml premix or 2.5mg/0.5ml) albuterol 100 mg/20 mL albuterol low dose (0.63mg/3ml & 1.25mg/3ml)	arformoterol tartrate (generic Brovana) BROVANA (arformoterol) formoterol fumarate (generic Perforomist) levalbuterol (generic for Xopenex) PERFOROMIST (formoterol)	
	ORAL	
albuterol <b>SYRUP</b>	albuterol <b>TABLET</b> albuterol ER (generic for Vospire ER) metaproterenol (formerly generic for Alupent) terbutaline (generic for Brethine)	

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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-dihydi diltiazem (generic Cardizem) verapamil (generic Calan/Isoptin) LONG-A	isradipine (generic Dynacirc) nicardipine (generic Cardene) nifedipine (generic Procardia) nimodipine (generic Nimotop) NYMALIZE (nimodipine) SOLN ropyridines  ACTING Dyridines  felodipine ER (generic Plendil) KATERZIA (amlodipine) Plevamlodipine (generic Conjupri) NR nisoldipine (generic Sular)	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
Non dibyd	NORLIQVA (amolidipine) <sup>AL,NR,QL</sup> <b>SOLN</b> copyridines	
diltiazem ER (generic Cardizem CD) verapamil ER <b>TAB</b>	CALAN SR (verapamil) diltiazem ER (generic Cardizem LA) MATZIM LA (diltiazem ER) TIAZAC (diltiazem) verapamil ER CAPS verapamil 360mg CAPS verapamil ER (generic Verelan PM)	

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#### CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

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

#### **COLONY STIMULATING FACTORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NEUPOGEN (filgrastim) <b>VIAL</b> NYVEPRIA <b>SUB-Q</b> (pegfilgrastim-apgf)	FULPHILA SUB-Q (pegfilgrastim-jmdb) • GRANIX (tbo-filgrastim) LEUKINE (sargramostim) NEULASTA SYR(pegfilgrastim) NEUPOGEN DISP SYR (filgrastim) NIVESTYM SYR,VIAL (filgrastim-aafi) RELEUKO (filgrastim-ayow) SYR,VIAL UDENYCA SUB-Q (pegfilgrastim-cbqv) ZARXIO (filgrastim-sndz) ZIEXTENZO SYR (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	FINZALA (ethinyl estradiol/norethindrone acetate) <b>CHEW</b> NR	
listed. Brand name products may be subject to Maximum Allowable Cost (MAC) pricing or require substitution with a generic equivalent	norethindrone/ethinyl estradiol FE estrophasic (generic EstropFE) <sup>NR</sup>	
Specific agents can be looked up using the Drug Look-up Tool at: <a href="https://druglookup.fhsc.com/druglookupweb/?client=nestate">https://druglookup.fhsc.com/druglookupweb/?client=nestate</a>		

## **Nebraska Medicaid Preferred Drug List**

#### 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) SPIRIVA RESPIMAT (tiotropium)	<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>
INHALATION	SOLUTION	exacerbation in last year upon initial review
albuterol/ipratropium (generic for Duoneb) ipratropium <b>SOLN</b> (generic for Atrovent)	LONHALA (glycopyrrolate inhalation soln) YUPELRI (revefenacin)	
ORAL	AGENT	
	DALIRESP (roflumilast) <sup>CL, QL</sup> roflumilast (generic Daliresp) <sup>CL,NR,QL</sup>	

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	guaifenesin/codeine <b>LIQUID</b> hydrocodone/homatropine SYRUP promethazine/codeine <b>SYRUP</b> promethazine/phenylephrine/codeine SYRUP	<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>

Entire December 2022 PDL with November P&T changes are Highlighted in Red effective January 20, 2023 CYSTIC FIBROSIS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BRONCHITOL (mannitol) <sup>AL,CL,QL</sup> KALYDECO <b>PACKET, TAB</b> (ivacaftor) <sup>QL, AL</sup> ORKAMBI (lumacaftor/ivacaftor) <b>PACKET, TAB</b> <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
COSENTYX (secukinumab) <sup>CL</sup> ENBREL (etanercept) KIT, MINI CART, PEN, SYR, VIAL <sup>QL</sup> HUMIRA (adalimumab) <sup>QL</sup> OTEZLA (apremilast) ORAL <sup>CL,QL</sup>	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIBINQO (abrocitinib) <sup>AL,QL</sup> CIMZIA (certolizumab pegol) <sup>QL</sup> ENSPRYNG (satralizumab-mwge) SUB-Q ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYR KINERET (anakinra) OLUMIANT (baricitinib) TAB <sup>CL,QL</sup> ORENCIA (abatacept) SUB-Q RINVOQ ER (upadacitinib) <sup>CL,QL</sup> SILIQ (brodalumab) SIMPONI (golimumab) SKYRIZI (risankizamab-rzaa) SYR SKYRIZI ON-BODY (risankizamab-rzaa) <sup>QL</sup> SKYRIZI PEN (risankizamab-rzaa) <sup>QL</sup> STELARA (ustekinumab) SUB-Q TALTZ (ixekizumab) <sup>AL</sup> TREMFYA (guselkumab) <sup>QL</sup> XELJANZ (tofacitinib) TAB, SOLN <sup>CL,QL</sup> XELJANZ XR (tofacitinib) TAB <sup>CL,QL</sup>	<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 TWO preferred agents 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>JAK-Inhibitors: For FDA approved indications that require a patient to have had an inadequate response to a TNF blocker, documentation of an inadequate response is required.</li> <li>Drug-specific criteria:</li> <li>Cosentyx: Requires treatment failure of Enbrel OR Humira with the same FDA-approved indications and age limits.</li> <li>Otezla: Requires a trial of Humira</li> </ul>

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#### **DIURETICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
amiloride TABLET bumetanide TABLET chlorothiazide TABLET chlorothialidone TABLET (generic Diuril) furosemide SOLN, TABLET (generic Lasix) hydrochlorothiazide CAPS, TABLET (generic Microzide) indapamide TABLET metolazone TABLET spironolactone TABLET (generic Aldactone) torsemide TABLET	CAROSPIR (spironolactone) SUSP eplerenone TABLET (generic Inspra) <sup>CL</sup> ethacrynic acid CAPS (generic Edecrin) KERENDIA (finerenone) TABLET CL,QL methyclothiazide TABLET THALITONE (chlorthalidone) TABLET triamterene (generic Dyrenium)	<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>Eplerenone: Will be approved with a failed trial or intolerance to spironolactone, a trial with two preferred agents is not required.</li> <li>Kerendia: For diagnosis of chronic kidney disease associated with Type-II diabetes in adults, trial of a preferred agent not required.</li> </ul>
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

EPINEPHRINE, SELF-INJECTED**		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
epinephrine (AUTHORIZED GENERIC for Epipen/ Epipen Jr.) AUTOINJECTOR EPIPEN (epinephrine) AUTOINJ EPIPEN JR. (epinephrine) AUTOINJ	epinephrine (generic for Adrenaclick) epinephrine (generic for Epipen/ Epipen Jr.) <b>AUTOINJECTOR</b> SYMJEPI (epinephrine) <b>PFS</b>	Non-preferred agents require clinical documentation why the preferred product within this drug class is not appropriate

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
EPOGEN (rHuEPO) RETACRIT (EPOETIN ALFA- EPBX)	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) evofloxacin <b>TABLET</b> (generic Levaquin)	BAXDELA (delafloxacin) ciprofloxacin ER ciprofloxacin SUSP (generic Cipro) levofloxacin SOLN 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, linezolic sulfamethoxazole/trimethoprim)</li> <li>Ciprofloxacin/Levofloxacin Suspensior 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
AMITIZA (lubiprostone) <sup>AL, QL</sup> LINZESS (linaclotide) <sup>QL</sup> MOVANTIK (naloxegol oxalate) <sup>QL</sup>	alosetron (generic Lotronex)  IBSRELA (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)	<ul> <li>Non-preferred agents will be approved for patients who have failed a 30-day trial of ONE preferred agent within this drug class</li> <li>Lotronex®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate</li> <li>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</li> <li>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</li> <li>Trulance®: Covered for diagnosis of either chronic idiopathic constipation or IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.)</li> <li>Viberzi®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate</li> </ul>

### **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, PEN, SYRINGE, VIAL ZEGALOGUE (dasiglucagon) <sup>AL,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>

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## **GLUCOCORTICOIDS, INHALED**

GLUCOCORTICOIDS  ASMANEX (mometasone) <sup>QL,AL</sup> FLOVENT HFA (fluticasone) PULMICORT FLEXHALER (budesonide)  ARMONAIR DIGIHALER (fluticasone) <sup>AL,QL</sup> ARMONAIR RESPICLICK (fluticasone) <sup>AL</sup> ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) <sup>AL,QL</sup> FLOVENT DISKUS (fluticasone) fluticasone HFA (generic Flovent HFA) QVAR (beclomethasone) QVAR Redihaler (beclomethasone)	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 rug-specific criteria:  budesonide respules: Covered without PA for age ≤ 8 years  OR for diagnosis of eosinophilic
FLOVENT HFA (fluticasone) PULMICORT FLEXHALER (budesonide)  ARMONAIR DIGIHALER (fluticasone) <sup>AL,QL</sup> ARMONAIR RESPICLICK (fluticasone) <sup>AL</sup> ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) <sup>AL,QL</sup> FLOVENT DISKUS (fluticasone) fluticasone HFA (generic Flovent HFA) QVAR (beclomethasone)	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  rug-specific criteria: budesonide respules: Covered without PA for age ≤ 8 years
GLUCOCORTICOID/BRONCHODILATOR COMBINATIONS  ADVAIR DISKUS (fluticasone/ salmeterol) <sup>QL</sup> ADVAIR HFA (fluticasone/salmeterol) <sup>QL</sup> DULERA (mometasone/formoterol) SYMBICORT (budesonide/ formoterol) SYMBICORT (budesonide/ formoterol)  Fluticasone/salmeterol (generic Symbicort) fluticasone/salmeterol (generic Advair Diskus) <sup>QL</sup> fluticasone/vilanterol (Breo Ellipta) TRELEGY ELLIPTA (fluticasone/ umeclidinium/vilanterol) WIXELA INHUB (generic Advair Diskus) <sup>QL</sup> INHALATION SOLUTION budesonide RESPULES (generic Pulmicort)	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.

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## **GLUCOCORTICOIDS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ALKINDI (hydrocortisone)  GRANULES <sup>AL</sup> CORTEF (hydrocortisone) cortisone TAB dexamethasone INTENSOL  EMFLAZA (deflazacort) SUSP, TAB <sup>CL</sup> ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg, 32mg ORTIKOS ER (budesonide) <sup>AL,QL</sup> prednisolone sodium phosphate	<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> <li>Tarpeyo: Indicated for the treatment of primary immunoglobulin A nephropathy (IgAN)</li> </ul>

## **GROWTH HORMONES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GENOTROPIN (somatropin)	HUMATROPE (somatropin)	Growth Hormone PA Form
NUTROPIN AQ (somatropin)	OMNITROPE (somatropin)	Growth Hormone Criteria
NORDITROPIN (somatropin)	SAIZEN (somatropin)	
	SEROSTIM (somatropin)	
	SKYTROFA (lonapegsomatropin-tcgd)	
	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

BERINERT (C1 esterase inhibitor, human) INTRAVENOUS  HAEGARDA (C1 esterase inhibitor, human) AL,CL INTRAVENOUS  HAEGARDA (C1 esterase inhibitor, human) AL,CL SUB-Q  icatibant acetate (generic for FIRAZYR) L SUB-Q  CAPAL,OL RUCONEST (recombinant human C1 inhibitor) L INTRAVENOUS  TAKHZYRO (lanadelumab-flyo) AL,CL VIAL, SYRINGENR  CINRYZE (C1 esterase inhibitor, human) AL,CL INTRAVENOUS  FIRAZYR  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 with the same indication.  Drug-Specific Criteria	Preferred Agents	Non Professed Agents	Prior Authorization/Class Criteria
human) INTRAVENOUS  HAEGARDA (C1 esterase inhibitor, human) AL,CL SUB-Q icatibant acetate (generic for FIRAZYR) SUB-Q  icatibant acetate (generic for FIRAZYR) INTRAVENOUS  FIRAZYR) INTRAVENOUS  TAKHZYRO (lanadelumab-flyo) AL,CL VIAL, SYRINGENR  human) AL,CL INTRAVENOUS  (icatibant acetate) ALSUB-Q  ORLADEYO (berotralstat)  CAPAL,QL RUCONEST (recombinant human C1 inhibitor) INTRAVENOUS  TAKHZYRO (lanadelumab-flyo) AL,CL VIAL, SYRINGENR  Non-preferred agents will be approved for patients who have a failed trial or a contraindication to ONE preferred agent within this drug class with the same indication.  Drug-Specific Criteria	Preferred Agents	Non-Preferred Agents	PHOI AUTHORIZATION/Glass Griteria
human) <sup>AL,CL</sup> SUB-Q icatibant acetate (generic for FIRAZYR) <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> 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 with the same indication.  Drug-Specific Criteria			HAE Treatments PA Form
and Takhzyro, require a history of two or more HAE attacks monthly, and trial and failure or contraindication to oral danazol	human) <sup>AL,CL</sup> <b>SUB-Q</b> icatibant acetate (generic for	(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>	and deficient or dysfunctional C1 esterase inhibitor enzyme. Concomitant use with ACE inhibitors, NSAIDs, or estrogen- containing 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 with the same indication.  Drug-Specific Criteria  Cinryze, Haegarda, Orladeyo, and Takhzyro, require a history of two or more HAE attacks monthly, and trial and failure or

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### **HEMOPHILIA TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior 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	approved for patients who have failed a trial of ONE preferred agent within this drug class
FAC	FOR IX	
ALPROLIX	ALPHANINE SD	
BENEFIX	IDELVION IXINITY 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 WILLEBRA	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) SOLN, TABLET EPIVIR HBV (lamivudine) TABLET, SOLN 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>

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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) <b>TABLET<sup>CL</sup>, PELLET<sup>AL,CL,NR</sup></b> VOSEVI (sofosbuvir/velpatasvir/ voxilaprevir) <sup>CL</sup>	HARVONI 200/45MG, <b>TABLET</b> (sofosbuvir/ledipasvir) <sup>CL</sup> HARVONI (ledipasvir/sofosbuvir) <sup>CL</sup> <b>PELLET</b> sofosbuvir/ledipasvir (generic Harvoni) <sup>CL</sup> SOVALDI (sofosbuvir) <sup>CL</sup> <b>PELLET</b> SOVALDI <b>TABLET</b> (sofosbuvir) <sup>CL</sup> VIEKIRA <b>PAK</b> (ombitasvir/ paritaprevir/ritonavir/dasabuvir) <sup>CL</sup> ZEPATIER (elbasvir/grazoprevir) <sup>CL</sup>	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
DIDA	N/IDIN	1 or 4
	VIRIN	■ Vosevi: Requires documentation of non-
ribavirin 200mg CAPSULE, TABLET	REBETOL (ribavirin)	response after previous treatment course of Direct Acting Anti-viral agent (DAA) for
INTERFERON		genotype 1-6 without cirrhosis or with compensated cirrhosis
PEGASYS (pegylated interferon alfa- 2a) CL PEG-INTRON (pegylated interferon alfa-2b) CL		Somportune of Those

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## **HISTAMINE II RECEPTOR BLOCKERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
famotidine Rx, OTC TAB (generic Pepcid) famotidine SUSP	cimetidine <b>TABLET</b> , <b>SOLN</b> <sup>CL</sup> (generic Tagamet) nizatidine <b>CAPS</b> (generic Axid)	<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.</li> </ul>
		contagiosum or common wart V. Vulgaris treatment

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## HIV / AIDSCL

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

Entire December 2022 PDL with November P&T changes are Highlighted in Red effective January 20, 2023

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

Preferred Agents	Non-Preferred Agents	Pric	or Authorization/Class Criteria
	APTIVUS CAPS, SOLN (tipranavir) CRIXIVAN (indinavir) fosamprenavir TAB (generic Lexiva) INVIRASE (saquinavir) LEXIVA SUSP (fosamprenavir) LEXIVA TABLET (fosamprenavir) NORVIR POWDER, SOLN (ritonavir) NORVIR (ritonavir) TAB PREZISTA (darunavir) SUSP, TABLET REYATAZ POWDER (atazanavir) VIRACEPT (nelfinavir)		All agents require:  Diagnosis of HIV/AIDS required; OR Diagnosis of Pre and Post Exposure Prophylaxis  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  Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	KALETRA SOLN (lopinavir/ritonavir) KALETRA TAB (lopinavir/ritonavir) lopinavir/ritonavir TAB (generic Kaletra) PREZCOBIX (darunavir/cobicistat)  Coloridation of the c	<ul> <li>All agents require:         <ul> <li>Diagnosis of HIV/AIDS required; OR</li> <li>Diagnosis of Pre and Post Exposure Prophylaxis</li> </ul> </li> <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</li> </ul>
abacavir/lamivudine (generic Epzicom) CIMDUO (lamivudine/tenofovir)QL DESCOVY (emtricitabine/tenofovir)QL, CI emtricitabine/tenofovir (generic Truvada)CL lamivudine/zidovudine (generic Combivir)	abacavir/lamivudine/zidovudine (generic Trizivir) COMBIVIR (lamivudine/zidovudine) EPZICOM (abacavir sulfate/lamivudine) TEMIXYS (lamivudine/tenofovir) TRIZIVIR (abacavir/lamivudine/zidovudine) TRUVADA (emtricitabine/tenofovir)	conditions not recommended with preferred agents  Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ATRIPLA (efavirenz/emtricitabine/tenofovir) efavirenz/lamivudine/tenofovir (generic for Symfi) <sup>QL</sup> efavirenz/lamivudine/tenofovir	Prior Authorization/Class Criteria  All agents require:  Diagnosis of HIV/AIDS required; OR  Diagnosis of Pre and Post Exposure Prophylaxis  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  Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy

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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>

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## HYPOGLYCEMICS. INCRETIN MIMETICS/ENHANCERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
·	CEPTOR AGONIST (GLP-1 RA)CL	GLP-1 RA Criteria
OZEMPIC (semaglutide) TRULICITY (dulaglutide) VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) BYDUREON BCISE PEN (exenatide) <sup>QL</sup> BYDUREON (exenatide ER) BYDUREON PEN (exenatide ER) subcutaneous BYETTA (exenatide) subcutaneous MOUNJARO (tirazepatide) <sup>NR</sup> PEN RYBELSUS (semaglutide)	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR  A diagnosis of ASCVD associated with a diagnosis of Type II diabetes (no metformin trial required)  Non-preferred agents will be approved for patients who have:  Failed a trial of TWO preferred agents within GLP-1 RA
INSULIN/GLP-1 RA	A COMBINATIONS	AND Diagnosis of diabetes with HbA1C
	SOLIQUA (insulin glargine/lixisenatide) XULTOPHY (insulin degludec/liraglutide)	<ul> <li>► Diagnosis of diabetes with FIDATC</li> <li>≥ 7 AND</li> <li>■ Trial of metformin, or contraindication or intolerance to metformin</li> </ul>
AMYLIN	ANALOG	Amylin Analog Criteria
	SYMLIN (pramlintide) subcutaneous	<ul> <li>ALL criteria must be met</li> <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>Monitoring of glucose during initiation of therapy</li> </ul>
DIPEPTIDYL PEPTIDAS	E-4 (DPP-4) INHIBITOR <sup>QL</sup>	- initiation of therapy
JANUMET (sitagliptin/metformin) JANUMET XR (sitagliptin/metformin) JANUVIA (sitagliptin) JENTADUETO (linagliptin/metformin) TRADJENTA (linagliptin)	alogliptin (generic for Nesina) alogliptin/metformin (generic for Kazano) GLYXAMBI (empagliflozin/linagliptin) 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>	DPP-4 Inhibitor Criteria Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin.  Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within the DPP-4 inhibitor class

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## HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HUMALOG (insulin lispro) U-100 CARTRIDGE, PEN, VIAL  HUMALOG JR. (insulin lispro) U-100 KWIKPEN  HUMALOG MIX VIAL (insulin lispro/lispro protamine)  HUMALOG MIX KWIKPEN (insulin lispro/lispro protamine)  HUMULIN (insulin) VIAL  HUMULIN 70/30 VIAL  HUMULIN U-500 VIAL  HUMULIN OTC PEN  HUMULIN 70/30 OTC PEN  insulin aspart (generic for Novolog) insulin aspart/insulin aspart protamine PEN, VIAL (generic for Humalog) PEN, VIAL, JR KWIKPEN  insulin lispro/lispro protamine KWIKPEN (Humalog Mix Kwikpen)  LANTUS SOLOSTAR PEN (insulin glargine)  LANTUS (insulin glargine) VIAL  LEVEMIR (insulin detemir) PEN, VIAL  NOVOLIN (insulin) PEN  NOVOLOG (insulin aspart) CARTRIDGE, FLEXPEN, VIAL  (insulin aspart/aspart protamine)	ADMELOG (insulin lispro) PEN, VIAL AFREZZA (regular insulin) INHALATION APIDRA (insulin glulisine) BASAGLAR (insulin glargine, rec) PEN, TEMPO PEN FIASP (insulin aspart) CARTRIDGE, PEN, VIAL HUMALOG U-100 TEMPO PEN HUMALOG (insulin lispro) U-200 KWIKPEN insulin degludec (generic Tresiba) Insulin glargine PEN, VIAL 200U/mL PEN insulin Glargine-YFGN PEN, VIAL (generic for Semglee-YFGN) LYUMJEV KWIKPEN, TEMPO PEN, VIAL (insulin lispro-aabc) NOVOLIN (insulin) NOVOLIN 70/30 VIAL(insulin) NOVOLOG MIX (insulin aspart/aspart protamine) VIAL TOUJEO SOLOSTAR (insulin glargine) SEMGLEE (insulin glargine) PEN, VIAL SEMGLEE YFGN (insulin glargine) PEN, VIAL TRESIBA (insulin degludec)	<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>Afrezza®: Approved for T1DM on long-acting insulin with no current history of smoking or chronic lung disease</li> <li>Humulin® R U-500 Kwikpen:</li></ul></li></ul>

## **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>	<ul> <li>Non-preferred agents will be approved for patients with:         Failure of a trial of ONE preferred agent in another Hypoglycemic class OR     </li> <li>T2DM and inadequate glycemic control</li> </ul>

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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>SOLN</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>	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR  A diagnosis of ASCVD or Heart Failure, or Chronic Kidney Disease associated with a diagnosis of Type II diabetes (no metformin trial required)  Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class  Drug Specific Criteria:  Farxiga: May be approved for a diagnosis of heart failure with reduced ejection fraction (NYHA class II-IV) without a diagnosis of diabetes  May be approved for a diagnosis of chronic kidney disease at risk of progression without a diagnosis of diabetes  Jardiance: May be approved for a diagnosis of diabetes  Jardiance: May be approved for a diagnosis of diabetes

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## 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	<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>
SULFONYLUREA	COMBINATIONS	
glipizide/metformin glyburide/metformin (generic Glucovance)		

## HYPOGLYCEMICS, TZD

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
THIZAOLIDINEDIONES (TZDs)		Non-preferred agents will be
pioglitazone (generic for Actos)	AVANDIA (rosiglitazone)	approved for patients who have failed a trial of THE preferred agen
TZD COMBINATIONS		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) <sup>QL</sup> pirfenidone (generic for Esbriet) <sup>QL</sup>	<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
FASENRA (benralizumab) AL PEN XOLAIR (omalizumab) SYRAL,QL	NUCALA (mepolizumab) <sup>AL</sup> AUTO-INJ, SYR	<ul> <li>All agents require prior authorization AND an FDA-approved diagnosis for approval</li> <li>Non-preferred agents require a trial of a preferred agent within this drug class with the same indication</li> <li>For asthma indications: All agents must be prescribed by or in consultation with an allergist, immunologist, or pulmonologist</li> <li>Agents listed may have other FDA approved indications, and will be subject to prior authorization</li> <li>Drug Specific Criteria:</li> <li>Dupixent: (for other indications, please see the Immunomodulators, Atopic Dermatitis therapeutic class) – For Eosinophilic Asthma or Corticosteroid Dependent Asthma: Patients must be ages 6 and older. Documentation of moderate to severe asthma with either eosinophils &gt;/= 150 + 1 exacerbation OR oral corticosteroid dependency AND prior drug therapy of med-high or max-tolerated inhaled corticosteroid + controller OR max-tolerated inhaled corticosteroid / long acting beta agonist combo</li> </ul>

# **Nebraska Medicaid Preferred Drug List**

## with Prior Authorization Criteria

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DUPIXENT (dupilumab) PEN,SYR*L.C. ELIDEL (pimecrolimus)  SUB-Q <sup>AL_OL</sup> OPZELURA (cuxolitinib phosphate)  CREAM*L.C. G.  PROTOPIC (tacrolimus)  Drug-specific criteria:  Dupixent:  1. Atopic Dermatitis: Trial or failure of a topical corticosteroid AND a topical cacineum inhibitor.  2. Essinghilis, power field.  2. Essinghilis, power field.  3. Nasal Potyps: Documentation of treatment failure of a reposited of the special corticosteroid of treatment failure of a reposited of the special corticosteroid of treatment failure of a reposited of the special corticosteroid of treatment failure of a reposited corticosteroid of treatment failure of the special corticosteroid of treatment failure of a reposited corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of the special corticosteroid of treatment failure of a reposite of the special corticosteroid of the

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## **IMMUNOMODULATORS, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
imiquimod (generic for Aldara)	HYFTOR (sirolimus) <sup>AL,NR</sup> <b>GEL</b> 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

## **IMMUNOSUPPRESSIVES, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathioprine (generic Imuran) azathioprine (generic Azasan) <sup>NR</sup> cyclosporine, modified CAPS (generic Neoral) everolimus (generic for Zortress) <sup>AL</sup> mycophenolate CAPS, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLN RAPAMUNE (sirolimus) TABLET tacrolimus	ASTAGRAF XL (tacrolimus) AZASAN (azathioprine) cyclosporine CAP, SOFTGEL cyclosporine, modified SOLN (generic Neoral) ENVARSUS XR (tacrolimus) GENGRAF (cyclosporine, modified) CAP, SOLN mycophenolate SUSP (generic Cellcept) mycophenolic acid MYFORTIC (mycophenolate sodium) PROGRAF (tacrolimus) CAPS, PACKET REZUROCK (belumosudil) <sup>AL,QL</sup> TAB SANDIMMUNE (cyclosporine) CAPS, SOLN sirolimus SOLN, TABLET (generic Rapamune) TAVNEOS (avacopan) <sup>QL</sup> CAPS ZORTRESS (everolimus) AL	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

# **Nebraska Medicaid Preferred Drug List**

## with Prior Authorization Criteria

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## **INTRANASAL RHINITIS DRUGS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
		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 B)</li> <li>Xhance: Indicated for treatment of</li> </ul>
CORTICO	STEROIDS	nasal polyps in <u>&gt;</u> 18 years only
fluticasone <b>Rx</b> (generic for Flonase Rx)	BECONASE AQ (beclomethasone) budesonide Rx (generic Rhinocort) flunisolide (generic Nasalide) fluticasone OTC (generic Flonase OTC) mometasone (generic Nasonex) OMNARIS (ciclesonide) QNASL 40 & 80 (beclomethasone) XHANCE (fluticasone) ZETONNA (ciclesonide)	

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### **LEUKOTRIENE MODIFIERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
montelukast <b>TAB/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>CAPS</b> clindamycin palmitate <b>SOLN</b> linezolid <b>TAB</b>	CLEOCIN (clindamycin ) CAPS CLEOCIN PALMITATE (clindamycin) linezolid SUSP SIVEXTRO (tedizolid phosphate) ZYVOX (linezolid) SUSP, TAB	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 cholestyramine (generic Questran) colestipol TABLETS (generic Colestid)	colestipol GRANULES (generic Colestid) QUESTRANTS  colestid) QUESTRAN LIGHT (cholestyramine)	<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>Colesevelam: Trial not required for</li> </ul>
TREATMENT OF HOMOZYGOUS FA	MILIAL HYPERCHOLESTEROLEMIA  JUXTAPID (lomitapide) <sup>CL</sup> KYNAMRO (mipomersen) <sup>CL</sup>	diabetes control and monotherapy with metformin, sulfonylurea, or insulin has been inadequate Juxtapid®/ Kynamro®:  Approved for diagnosis of homozygous familial hypercholesterolemia (HoFH)
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)	OR  o Treatment failure/maximized dosing/contraindication to ALL the following: statins, ezetimibe, niacin, fibric acid derivatives, omega-3 agents, bile acid sequestrants  o Require faxed copy of REMS PA form  Vascepa®: Approved for TG ≥ 500
omega-3 fatty acids (generic for Lovaza)	icosapent (generic for Vascepa) <sup>CL</sup> omega-3 OTC VASCEPA (icosapent) <sup>CL</sup> ORPTION INHIBITORS  NEXLIZET (bempedoic acid/ezetimibe) <sup>QL</sup>	

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LIPOTROPICS, OTHER (continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BTILISIN/KEXIN TYPE 9 (PCSK9) BITORS  PRALUENT (alorocumab) <sup>CL</sup> REPATHA (evolocumab) <sup>CL</sup>	<ul> <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>AND         <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
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>Non-preferred agents will be 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:</li> <li>Altoprev®: One of the TWO trials must be IR lovastatin</li> </ul>
STATIN COI	MBINATIONS	<ul> <li>Combination products: Require clinical reason why individual ingredients cannot be</li> </ul>
	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
MACRO	OLIDES	Require clinical reason why
azithromycin (generic Zithromax) clarithromycin <b>TABLET</b> , <b>SUSP</b> (generic Biaxin) E.E.S. <b>SUSP</b> (erythromycin ethylsuccinate)	clarithromycin ER (generic Biaxin XL) E.E.S. <b>TABLET</b> (erythromycin ethylsuccinate) ERY-TAB (erythromycin) erythromycin ethylsuccinate <b>SUSP</b> ERYPED <b>SUSP</b> (erythromycin) ERYTHROCIN (erythromycin) erythromycin base <b>TABLET</b> , <b>CAPS</b>	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	Non-preferred agents require a trial of the preferred agent AND will be approved for an FDA-approved indication  Drug-specific criteria:  Xatmep <sup>TM</sup> :Indicated for pediatric patients only

## **MOVEMENT DISORDERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) <sup>CL</sup> INGREZZA (valbenazine) <sup>AL,CLQL</sup> <b>CAPS</b>	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 a trial and failure of a preferred agent with the same indication or a clinical reason why a preferred agent in this class cannot be used.
		<ul> <li>Drug-specific criteria:</li> <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>

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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> dimethyl fumarate (generic for Tecfidera) KESIMPTA (Ofatumumab) <sup>CL,QL</sup>	AUBAGIO (teriflunomide) BAFIERTAM (monomethyl fumarate) <sup>QL</sup> dalfampridine (generic Ampyra) <sup>QL</sup> EXTAVIA (interferon beta-1b) <sup>QL</sup> fingolimod (generic Gilenya) <sup>NR,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) REBIF (interferon beta-1a) <sup>QL</sup> TASCENSO ODT (fingolimod) TABLET <sup>AL,NR</sup> TECFIDERA (dimethyl fumarate) VUMERITY (diroximel) <sup>QL</sup> ZEPOSIA (ozanimod) <sup>AL,CL,QL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <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> <li>Zeposia: Approved for a diagnosis of relapsing forms of multiple sclerosis (MS) with trial of ONE preferred agent OR a diagnosis of moderately to severely active ulcerative colitis and treatment failure of Humira.</li> </ul>

## **NITROFURAN DERIVATIVES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
nitrofurantoin macrocrystals CAPSULE (generic Macrodantin) nitrofurantoin monohydrate- macrocrystals CAPS (generic Macrobid)	nitrofurantoin <b>SUSPENSION</b> (genericFuradantin)	<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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#### **Preferred Agents Non-Preferred Agents** Prior Authorization/Class Criteria Non-preferred agents within COX-**COX-I SELECTIVE** 1 SELECTIVE group will be diclofenac sodium (generic for diclofenac potassium (generic approved for patients who have Voltaren) Cataflam, Zipsor) failed no less than 30-day trial of TWO preferred agents within this ibuprofen OTC, Rx (generic for Advil, diclofenac SR (generic Voltaren-XR) Motrin) CHEW, DROPS, SUSP. drug class diflunisal (generic Dolobid) All combination agents require a TAB etodolac & SR (generic Lodine/XL) clinical reason why individual ibuprofen OTC (generic for Advil, fenoprofen (generic Nalfon) agents can't be used separately Motrin) CAPS flurbiprofen (generic Ansaid) indomethacin CAPS (generic for ibuprofen/famotidine (generic Drug-specific criteria: Indocin) Duexis)CL Arthrotec®: Requires clinical ketorolac (generic for Toradol) reason why individual ingredients indomethacin ER (generic Indocin) cannot be used meloxicam **TAB** (generic for Mobic) ketoprofen & ER (generic Orudis) meclofenamate: Approvable nabumetone (generic for Relafen) meclofenamate (generic Meclomen) without trial of preferred agents for naproxen Rx, OTC (generic for menorrhagia mefenamic acid (generic Ponstel) Naprosyn) Sprix®: Approved for patients meloxicam CAP naproxen enteric coated unable to tolerate, swallow OR (generic Vivlodex)CL, QL absorb oral NSAIDs OR sulindac (generic for Clinoril) naproxen CR (generic Naprelan) contraindication OR trial of TWO preferred oral NSAIDs naproxen **SUSP** (generic Naprosyn) naproxen sodium (generic Anaprox) naproxen-esomeprazole (generic Vimovo) oxaprozin (generic Daypro) piroxicam (generic Feldene) tolmetin (generic Tolectin) ketorolac NASALQL (generic Sprix)

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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:	-
	DUEXIS (ibuprofen/famotidine) <sup>CL</sup> INDOCIN <b>RECTAL</b> , <b>SUSP</b> NALFON (flurbiprofen) RELAFEN DS (nabumetone) ZIPSOR (diclofenac)	
NSAID/GI PROTECTA	ZORVOLEX (diclofenac, submicronized)  ANT COMBINATIONS	-
133, 115, 51, 1 10 12 01	diclofenac/misoprostol (generic for Arthrotec)	
COX-II SE	LECTIVE	
celecoxib (generic for Celebrex)		

### **NSAIDs, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
diclofenac sodium <b>GEL (OTC only)</b> diclofenac (generic Pennsaid Solution)	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 AND a clinical reason why 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
CDK 4/6 INHIBITOR		Non-preferred agents DO NOT
	IBRANCE (palbociclib) KISQALI (ribociclib) KISQALI FEMARA CO-PACK VERZENIO (abemaciclib)	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  Patients undergoing treatment at the time of any preferred etatus.
CHEMO	THERAPY	the time of any preferred status change will be allowed to continue
capecitabine (generic for Xeloda) cyclophosphamide	XELODA (capecitabine)	<ul> <li>therapy</li> <li>Drug-specific critera</li> <li>anastrozole: May be approved for malignant neoplasm of male breast</li> </ul>
HORMONE	BLOCKADE	<ul> <li>(male breast cancer)</li> <li>Fareston®: Require clinical reason</li> </ul>
anastrozole (generic for Arimidex) exemestane (generic for Aromasin) letrozole (generic for Femara) tamoxifen citrate (generic for Nolvadex)	SOLTAMOX <b>SOLN</b> (tamoxifen) <sup>CL</sup> toremifene (generic for Fareston) <sup>CL</sup>	why tamoxifen cannot be used  letrozole: Approved for diagnosis of breast cancer with day supply greater than 12 – NOT approved for short term use  Soltamox: May be approved with documented swallowing difficulty
OTHER		
	NERLYNX (neratinib) PIQRAY (alpelisib) lapatinib (generic Tykerb) <sup>CL</sup> TALZENNA (talazoparib tosylate) <sup>QL</sup> TUKYSA(tucatinib) <sup>QL</sup>	

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

Non-Preferred Agents	Prior Authorization/Class Criteria
ALL	<ul> <li>Non-preferred agents DO NOT</li> </ul>
PURIXAN (mercaptopurine) <sup>AL</sup>	require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use
AML	from current treatment guidelines
DAURISMO (glasdegib maleate)QL IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib)QL XOSPATA (gilteritinib) QL  CLL  COPIKTRA (duvelisib) QL IMBRUVICA (ibrutinib) VENCLEXTA (venetoclax) ZYDELIG (idelalisib)  CML  BOSULIF (bosutinib) GLEEVEC (imatinib) HYDREA (hydroxyurea) ICLUSIG (ponatinib) SCEMBLIX (asciminib) SPRYCEL (dasatinib)	
, ,	
	_
JAKAFI (ruxolitinib)	
ELOMA	-
FARYDAK (panobinostat)  lenalidomide <sup>QL</sup> (generic Revlimid)  melphalan (generic for Alkeran)  NINLARO (ixazomib)  POMALYST (pomalidomide)  THALOMID (thalidomide)  XPOVIO (selinexor) CL	
BRUKINSA (zanubrutinib <sup>QL</sup> CALQUENCE (acalabrutinib) <sup>QL</sup> INREBIC (fedratinib dihydrochloride) <sup>QL</sup> INQOVI (decitabine/cedazuridine) VONJO (pacritinib) <sup>QL</sup> ZOLINZA (vorinostat)	
	AML  DAURISMO (glasdegib maleate) <sup>QL</sup> IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) <sup>QL</sup> XOSPATA (gilteritinib) <sup>QL</sup> COPIKTRA (duvelisib) <sup>QL</sup> IMBRUVICA (ibrutinib) VENCLEXTA (venetoclax) ZYDELIG (idelalisib)  CML  BOSULIF (bosutinib) GLEEVEC (imatinib) HYDREA (hydroxyurea) ICLUSIG (ponatinib) SCEMBLIX (asciminib) SCEMBLIX (asciminib) SPRYCEL (dasatinib) TASIGNA (nilotinib) <sup>CL</sup> MPN  JAKAFI (ruxolitinib)  FARYDAK (panobinostat) lenalidomide <sup>QL</sup> (generic Revlimid) melphalan (generic for Alkeran) NINLARO (ixazomib) POMALYST (pomalidomide) THALOMID (thalidomide) XPOVIO (selinexor) <sup>CL</sup> OTHER  BRUKINSA (zanubrutinib <sup>QL</sup> INREBIC (fedratinib dihydrochloride) INQOVI (decitabine/cedazuridine) VONJO (pacritinib) <sup>QL</sup>

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CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL – Age Limit

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ALK  ALECENSA (alectinib)  ALUNBRIG (brigatinib) <sup>QL</sup> LORBRENA (lorlatinib) <sup>QL</sup> ZYKADIA (ceritinib) CAPS, TAB	<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>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>
ALK / F	ROS1 / NTRK	
	ROZLYTREK (entrectinib) <sup>AL,QL</sup> XALKORI (crizotinib)	_
	EGFR	
	erlotinib (generic for Tarceva) EXKIVITY (mobocertinib) <sup>QL</sup> GILOTRIF (afatinib) IRESSA (gefitinib) TARCEVA (erlotinib) TAGRISSO (osimertinib) 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
temozolomide (generic for Temodar)	AYVAKIT (avapritinib) <sup>AL,QL</sup> BALVERSA (erdafitinib) CAPRELSA (vandetanib) COMETRIQ (cabozantinib) HEXALEN (altretamine) KOSELUGO (selumetinib) <sup>AL</sup> LONSURF (trifluridine/tipiracil) LYNPARZA (olaparib) LYTGOBI (futibatinib) <sup>NR</sup> PEMAZYRE (pemigatinib) <sup>QL</sup> RUBRACA (rucaparib) STIVARGA (regorafenib) TAZVERIK (tazemetostat) <sup>AL</sup> TURALIO (pexidartinib) <sup>QL</sup> TRUSELTIQ (infigratinib) CAPS VITRAKVI (larotrectinib) CAPS, SOLN ZEJULA (niraparib)	<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>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>

## ONCOLOGY AGENTS, ORAL, PROSTATE

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
abiraterone (generic Zytiga) <sup>AL,QL</sup> bicalutamide (generic Casodex) lutamide	EMCYT (estramustine) ERLEADA (apalutamide) <sup>QL</sup> nilutamide (generic Nilandron) NUBEQA (darolutamide) <sup>QL</sup> ORGOVYX (relugolix) XTANDI (enzalutamide) <sup>AL,QL</sup> ZYTIGA (abiraterone) <sup>AL,QL</sup> YONSA (abiraterone acetonide, submicronized)	<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>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
UTENT (sunitinib)	AFINITOR DISPERZ (everolimus) <sup>CL</sup> CABOMETYX (cabozantinib) everolimus (generic Afinitor) everolimus SUSP (generic Afinitor Disperz) FOTIVDA (tivozanib) INLYTA (axitinib) LENVIMA (lenvatinib) NEXAVAR (sorafenib) sorafenib (generic Nexavar) sunitinib malate (generic Sutent) VOTRIENT (pazopanib) WELIREG (belzutifan) <sup>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>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ERIVEDGE (vismodegib)	CELL ODOMZO (sonidegib) <sup>CL</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> </ul>
BRAF M	UTATION	<ul> <li>Patients undergoing treatment at</li> </ul>
MEKINIST (trametinib) TAFINLAR (dabrafenib)	BRAFTOVI (encorafenib) COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)	the time of any preferred status change will be allowed to continue therapy

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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 OTC (Pataday once daily)	ALOCRIL (nedocromil) ALOMIDE (lodoxamide) azelastine (generic for Optivar) BEPREVE (bepotastine besilate) bepotastine besilate (generic Bepreve) EMADINE (emedastine) epinastine (generic for Elestat) LASTACAFT (alcaftadine) OTC olopatadine 0.2% (generic Pataday once daily, Pataday OTC twice daily) PATADAY XS (olopatadine 0.7%) PATADAY OTC (olopatadine 0.2%) ZERVIATE (certirizine) <sup>AL</sup>	<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>

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## **OPHTHALMICS, ANTIBIOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FLUOROQUINOLONES		Non-preferred agents will be
ciprofloxacin <b>SOLN</b> (generic Ciloxan) ofloxacin (generic Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic Zymaxid) levofloxacin MOXEZA (moxifloxacin) moxifloxacin (generic Vigamox) moxifloxacin (generic Moxeza) VIGAMOX (moxifloxacin)	approved for patients who have failed a one-month trial of TWO preferred agent within this drug class  Azasite®: Approval only requires trial of erythromycin
MACROLIDES		
erythromycin	AZASITE (azithromycin) <sup>CL</sup>	
AMINOGLYCOSIDES		
gentamicin <b>OINT</b> gentamicin <b>SOLN</b> tobramycin (generic Tobrex drops)	TOBREX <b>OINT</b> (tobramycin)	
OTHER OPHTHALMIC AGENTS		
bacitracin/polymyxin B (generic Polysporin) polymyxin B/trimethoprim (generic Polytrim)	bacitracin neomycin/bacitracin/polymyxin B OINT neomycin/polymyxin B/gramicidin NEOSPORIN (neomycin/polymyxin B/gramcidin) sulfacetamide SOLN (generic Bleph-10) sulfacetamide OINT	

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
neomycin/polymyxin/dexamethasone (generic Maxitrol) sulfacetamide/prednisolone TOBRADEX SUSP, OINT (tobramycin and dexamethasone)	BLEPHAMIDE (prednisolone and sulfacetamide) BLEPHAMIDE S.O.P. neomycin/polymyxin/HC neomycin/bacitracin/poly/HC PRED-G SUSP, OINT (prednisolone/gentamicin) tobramycin/dexamethasone SUSP (generic 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>

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## **OPHTHALMICS, ANTI-INFLAMMATORIES**

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

#### OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
RESTASIS (cyclosporine) RESTASIS <b>MULTIDOSE</b> (cyclosporine) XIIDRA (lifitegrast)	CEQUA (cyclosporine) QL EYSUVIS (loteprednol etabonate)QL TYRVAYA (varenicline tartrate)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>

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## **OPHTHALMICS, GLAUCOMA**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIO	TICS	Non-preferred agents will be
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide)  VUITY (pilocarpine)	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 ANHYDI	RASE INHIBITORS	
,,	AZOPT (brinzolamide) brinzolamide (generic for Azopt)	
PROSTAGLAND	DIN 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) dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine)	
ОТІ	HER	
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 agents require a treatment failure of a preferred drug or patient-specific documentation of why a preferred product is not appropriate for the 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 TAB NARCAN (naloxone) SPRAY	KLOXXADO (naloxone) <b>NASAL</b> naloxone <b>SPRAY</b> (generic for Narcan) ZIMHI (naloxone) <sup>AL</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
CIPRO HC (ciprofloxacin/ hydrocortisone) CIPRODEX (ciprofloxacin/dexamethasone) neomycin/polymyxin/hydrocortisone (generic for Cortisporin) ofloxacin (generic for Floxin)	ciprofloxacin ciprofloxacin/dexamethasone (generic for CIPRODEX) COLY-MYCIN S(neomycin/ hydrocortisone/colistin) OTOVEL (ciprofloxacin/fluocinolone)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

## PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ambrisentan (generic Letairis) REVATIO (sildenafil) <sup>CL</sup> SUSP, TAB tadalafil (generic for Adcirca) <sup>CL</sup> TRACLEER (bosentan) TAB TYVASO (treprostinil) INHALATION VENTAVIS (iloprost) INHALATION	ADEMPAS (riociguat) <sup>CL</sup> ADCIRCA (tadalafil) <sup>CL</sup> bosentan (generic Tracleer) <b>TAB</b> LETAIRIS (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) sildenafil (generic Revatio) <sup>CL</sup> <b>SUSP</b> , <b>TAB</b> TADLIQ (tadalafil) <sup>NR</sup> <b>SUSP</b> TRACLEER (bosentan) <b>TAB FOR SUSPENSION</b> TYVASO DPI (treprostini) <sup>NR</sup> INHALATION POWDER 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 PANCREAZE (pancrelipase) ZENPEP (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>

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#### PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD CHEW + IRON (MULTIVITAMIN WITH IRON) CHEW  CHILDREN'S CHEWABLES (PEDI MULTIVIT NO.31/IRON/FOLIC, PEDI MULTIVIT NO.25/FOLIC ACID, PEDI MULTIVIT NO.23/FOLIC	FLORIVA (PEDI MULTIVIT NO.85/FLUORIDE) CHEW  FLORIVA PLUS (PEDI MULTIVIT NO.161/FLUORIDE DROP  MULTI-VIT-FLOR (PEDI MULTIVIT	<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>Drug specific criteria:</li> <li>DEKAs Plus: Approved for diagnosis of Cystic Fibrosis and does not require a trial of a</li> </ul>
ACID)  MULTIVIT-FLUOR (PEDI MULTIVIT NO.17 W-FLUORIDE, PEDI MULTIVIT NO.16 W- FLUORIDE) CHEW	NO.205/FLUORIDE) <b>CHEW</b> POLY-VI-FLOR (PEDI MULTIVIT NO.33/FLUORIDE) <b>CHEW</b> POLY-VI-FLOR (PEDI MULTIVIT NO.37 W-FLUORIDE) <b>DROPS</b>	preferred agent
MULTIVIT-FLUOR (PEDI MULTIVIT NO.2 W-FLUORIDE) <b>DROP</b> MULTIVIT-IRON-FLUOR (PEDI MULTIVIT 45/FLUORIDE/IRON)	POLY-VI-FLOR /0.25mg IRON (PEDI MULTIVIT 37/FLUORIDE/IRON) POLY-VI-FLOR /0.5mg IRON (PEDI MULTIVIT 33/FLUORIDE/IRON)	
PED MVIT A,C,D3 NO.21/FLUORIDE	POLY-VI-SOL (PEDIATRIC MULTIVITAMIN NO.192) <b>DROP</b>	
POLY-VI-SOL WITH IRON (PEDI MV NO.189/FERROUS SULFATE) DROPS	QUFLORA (PEDI MULTIVIT NO.157/FLUORIDE) <b>GUMMIES</b>	
TRI-VI-SOL (VIT A PALMITATE/VIT C/VIT D3) <b>DROP</b> S	QUFLORA FE (PED MULTIVIT 142/IRON/FLUORIDE) <b>CHEW</b> QUFLORA FE (PED MULTIVIT	
TRI-VITE-FLUORIDE (PED MVIT A,C,D3 NO.21/FLUORIDE)	QUFLORA FE (PED MOLTIVIT 151/IRON/FLUORIDE) <b>DROP</b> QUFLORA PED (PEDI MULTIVIT NO.63 W-FLUORIDE) <b>CHEW</b>	
	QUFLORA PED (PEDI MULTIVIT 84 WITH FLUORIDE, PEDI MULTIVIT NO.83 W-FLUORIDE) <b>DROP</b>	
	TRI-VI-FLOR (PED MVIT A,C,D3 NO.38/FLUORIDE) <b>DROPS</b>	

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#### **PENICILLINS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
amoxicillin CAPS, CHEWABLE TAB, SUSP, TAB ampicillin CAPS 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>TAB</b> CALPHRON OTC (calcium acetate) RENVELA (sevelamer carbonate)	AURYXIA (ferric citrate) calcium acetate CAPS 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
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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## **Nebraska Medicaid Preferred Drug List**

#### with Prior Authorization Criteria

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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
COMPLETENATE CHEW TABLET EXPECTA PRENATAL OTC FE C/FA FE C/FI FC C/FA FE C/VIT C/VIT B12/FA OTC MARNATAL-F O-CAL FA PNV2/IRON B-G SUC-P/FA/OMEGA-3 PNV 11-IRON FUM-FOLIC ACID-OM3 PNV NO.118/IRON FUMARATE/FA CHEW TABLET PNV NO.15/IRON FUM & PS CMP/FA PNV W-CA NO.40/IRON FUM/FA CMB NO.1 PNV WITH CA NO.68/IRON/FA NO.1/DHA PNV WITH CA,NO.72/IRON/FA PNV#16/IRON FUM & PS/FA/OMEGA-3 PNV2/IRON B-G SUC-P/FA/OMEGA-3 PRENATAL VIT #76/IRON,CARB/FA PRENATAL VIT/FE FUMARATE/FA OTC PUREFE OB PLUS PUREFE PLUS STUART ONE OTC TRINATAL RX 1 VITAFOL TAB CHEW VITAFOL ULTRA VP-PNV-DHA	CITRANATAL B-CALM COMPLETENATE CHEW TABLET DERMACINRX PRENATRIX OTC DERMACINRX PRETRATE TABLET OTC ENBRACE HR MULTI-MAC OTC NESTABS NESTABS ABC NESTABS ONE OB COMPLETE ONE OB COMPLETE ONE OB COMPLETE PREMIER OB COMPLETE TABLET OB COMPLETE WITH DHA PNV COMBO#47/IRON/FA #1/DHA PNV WITH CA,NO.72/IRON/FA PNV WITH CA,NO.74/IRON/FA OTC PNV119/IRON FUMARATE/FA/DSS PRENATAL H DHA OTC PRENATE AM PRENATE CHEWABLE TABLET PRENATE CHEWABLE TABLET PRENATE ENHANCE PRENATE ESSENTIAL PRENATE ESSENTIAL PRENATE FIXIE PRENATE FIXIE PRENATE STAR PRIMACARE SELECT-OB + DHA SELECT-OB + DHA SELECT-OB TAB CHEW TENDERA-OB OTC TRICARE TRINATAL RX 1 TRISTART DHA VITAFOL NANO VITAFOL-OB VITAFOL-OB VITAFOL-OB VITAFOL-ONE WESTGEL DHA	<ul> <li>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</li> </ul>

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AL – Age Limit

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## PROGESTERONE (hydroxyprogesterone caproate)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	hydroxyprogesterone caproate     (generic Makena)  MAKENA AUTO INJECTOR     (hydroxyprogesterone caproate)  MAKENA (hydroxyprogesterone caproate) SDV	<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> </ul> </li> <li>Maximum of 30 days per dispensing</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) esomeprazole magnesium (generic Nexium) RXQL esomeprazole magnesium (generic Nexium) OTCQL esomeprazole strontium lansoprazole (generic Prevacid)QL 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 omeprazole Rx AND pantoprazole OR Protonix SUSP.</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 compounded suspension.</li> <li>Patients ≥ 5 years of age- Only approve non-preferred for Gl diagnosis if:</li></ul></li></ul>

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#### **SEDATIVE HYPNOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BENZODI.	AZEPINES	Benzodiazepines 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)	<ul> <li>Non-preferred agents require a trial of the preferred benzodiazepine agent</li> <li>temazepam 7.5/22.5mg: Requires clinical reason why 15mg/30mg cannot be used</li> <li>Others Criteria</li> <li>Non-preferred agents require a trial</li> </ul>
ОТН	IERS	<ul> <li>of TWO preferred agents in the</li> </ul>
zaleplon (generic for Sonata) zolpidem (generic for Ambien)	BELSOMRA (suvorexant) <sup>AL,QL</sup> DAYVIGO (lemborexant) <sup>AL,QL</sup> doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopiclone (generic for Lunesta) HETLIOZ (tasimelteon) <sup>CL</sup> HETLIOZ LQ (tasimelteon) SUSP AL,QL QUVIVIQ (daridorexant) <sup>QL</sup> ramelteon (generic for Rozerem) zolpidem ER (generic for Ambien CR) zolpidem SL (generic for Intermezzo)	<ul> <li>of TWO preferred agents in the OTHERS sub-category</li> <li>Silenor: Must meet ONE of the following:         <ul> <li>Contraindication to all of the preferred oral sedative hypnotics agents in the OTHERS sub-category</li> <li>Medical necessity for doxepin dose &lt; 10mg</li> <li>Age greater than 65 years old or hepatic impairment (3mg dose will be approved if this criteria is met)</li> </ul> </li> <li>zolpidem/zolpidem ER: Maximum daily dose for females: zolpidem 5mg; zolpidem ER 6.25mg</li> <li>zolpidem SL: Requires clinical reason why half of zolpidem tablet cannot be used or documented swallowing disorder</li> </ul>

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#### SICKLE CELL ANEMIA TREATMENTAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DROXIA (hydroxyurea) ENDARI (Lglutamine) <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: May be approved for use in patients ages 2 to 17 years old without a trial of Droxia</li> </ul>

#### SINUS NODE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORLANOR <b>SOLN, TAB</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>

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#### **SKELETAL MUSCLE RELAXANTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
baclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) <sup>QL</sup> methocarbamol (generic Robaxin) tizanidine <b>TAB</b> (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) SUSP LORZONE (chlorzoxazone) <sup>CL</sup> LYVISPAH (baclofen) <sup>NR,QL</sup> GRANULES metaxalone (generic Skelaxin) NORGESIC FORTE     (orphenadrine/ASA/caffeine) orphenadrine ER PARAFON FORTE (chlorzoxazone) tizanidine CAPS ZANAFLEX (tizanidine) CAPS, TAB	<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		<ul> <li>Low Potency Non-preferred agents will be approved for patients who</li> </ul>
hydrocortisone OTC & RX CREAM, LOTION, OINT (Rx only) hydrocortisone/aloe OINT	alclometasone dipropionate (generic for Aclovate)  CAPEX SHAMPOO (fluocinolone)  DESONATE (desonide) GEL  desonide LOTION (generic for Desowen)  desonide CREAM, OINT (generic Desowen, Tridesilon)  fluocinolone 0.01% OIL (generic DERMA-SMOOTHE-FS)  hydrocortisone/aloe CREAM  hydrocortisone OTC OINT  TEXACORT (hydrocortisone)	have failed a trial of ONE preferred agent within this drug class
MEDIUM	POTENCY	Medium Potency Non-preferred
fluticasone propionate CREAM, OINT (generic Cutivate) mometasone furoate CREAM, OINT, SOLN (generic Elocon)	betamethasone valerate (generic Luxiq) clocortolone (generic Cloderm) fluocinolone acetonide (generic Synalar) flurandrenolide (generic Cordran) fluticasone propionate LOTION   (generic Cutivate) hydrocortisone butyrate (generic Locoid) hydrocortisone butyrate/emoll (generic Locoid Lipocream) hydrocortisone valerate (generic Westcort) PANDEL (hydrocortisone probutate 0.1%) prednicarbate (generic Dermatop)	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class

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**STEROIDS, TOPICAL (Continued)** 

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

## **Nebraska Medicaid Preferred Drug List**

## with Prior Authorization Criteria

Entire December 2022 PDL with November P&T changes are Highlighted in Red effective January 20, 2023 STIMULANTS AND RELATED AGENTS<sup>AL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Ampheta  ADDERALL XR (amphetamine salt combo)  amphetamine salt combination IR	Non-Preferred Agents  MULANTS  mine type  ADZENYS XR (amphetamine) amphetamine ER (generic Adzenys ER) SUSP  amphetamine salt combination ER (generic for Adderall XR) amphetamine sulfate (generic for Evekeo) dextroamphetamine (generic for Dexedrine) dextroamphetamine SOLN (generic Procentra) dextroamphetamine ER (generic for Dexedrine ER)	<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>Procentra®: May be approved with documentation of swallowing disorder</li> <li>Zenzedi®: Requires clinical reason</li> </ul>

## with Prior Authorization Criteria

Entire December 2022 PDL with November P&T changes are Highlighted in Red effective January 20, 2023 STIMULANTS AND RELATED ADHD DRUGS (Continued)<sup>AL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Methylphe CONCERTA (methylphenidate ER) <sup>QL</sup> 18mg, 27mg, 36mg, 54mg dexmethylphenidate (generic Focalin IR) dexmethylphenidate XR (generic Focalin XR) METHYLIN SOLN (methylphenidate) methylphenidate (generic Ritalin) methylphenidate SOLN (generic Methylin) QUILLICHEW ER CHEWTAB (methylphenidate)	Non-Preferred Agents  Penidate type  ADHANSIA XR (methylphenidate) QL APTENSIO XR (methylphenidate) AZSTARYS (serdexmethylphenidate and dexmethylphenidate)QL 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 108mg per day for ages &lt; 18</li> <li>Maximum accumulated dose of 72mg per day for ages &gt; 19</li> </ul>
	methylphenidate ER 18mg, 27mg, 36mg, 54mg (generic Concerta) <sup>QL</sup> methylphenidate ER <b>CAP</b> (generic	

## **Nebraska Medicaid Preferred Drug List**

## with Prior Authorization Criteria

Entire December 2022 PDL with November P&T changes are Highlighted in Red effective January 20, 2023 STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Non-Preferred Agents	Prior Authorization/Class Criteria
Preferred Agents Non-Preferred Agents  MISCELLANEOUS	
clonidine ER (generic for Kapvay) <sup>QL</sup> STRATTERA (atomoxetine)	—clonidine IR are available without prior authorization
	Drug-specific criteria:  armodafinil and Sunosi: Require trial of modafinil
	armodafinil and modafinil: approved only for:
modafanil (generic for Provigil) <sup>CL</sup> SUNOSI (solriamfetol) <sup>CL,QL</sup> WAKIX (pitolisant) <sup>CL,QL</sup>	<ul> <li>Sleep Apnea with documentation/confirmation via sleep study and documentation that C-PAP has been maxed</li> <li>Narcolepsy with documentation of diagnosis via sleep study</li> <li>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</li> <li>Sunosi approved only for:         <ul> <li>Sleep Apnea with documentation/confirmation via sleep study and documentation that C-PAP has been maxed</li> <li>Narcolepsy with documentation of diagnosis via sleep study</li> </ul> </li> <li>Wakix: approved only for excessive daytime sleepiness in adults with narcolepsy with documentation of narcolepsy diagnosis via sleep study</li> </ul>
	clonidine ER (generic for Kapvay) <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>

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#### **TETRACYCLINES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR (generic Vibramycin) doxycycline monohydrate 50MG, 100MG CAPS doxycycline monohydrate SUSP, TAB (generic Vibramycin) minocycline HCI CAPS, TAB (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 <b>CAP</b> (generic Adoxa/Monodox/ Oracea) minocycline HCI ER (generic Solodyn) NUZYRA (omadacycline) tetracycline VIBRAMYCIN <b>SUSP</b> (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:</li> <li>Demeclocycline: Approved for diagnosis of SIADH</li> <li>doxycycline suspension: May be approved with documented swallowing difficulty</li> </ul>

#### THROMBOPOIESIS STIMULATING PROTEINSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROMACTA (eltrombopag) <b>TAB</b> <sup>cL</sup>	DOPTELET (avatrombopag) MULPLETA (lusutrombopag) PROMACTA (eltrombopag) SUSP 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>

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#### **THYROID HORMONES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
levothyroxine <b>TAB</b> (generic Synthroid) liothyronine <b>TAB</b> (generic Cytomel) thyroid, pork <b>TAB</b> UNITHROID (levothyroxine)	EUTHYROX (levothyroxine) LEVO-T (levothyroxine) levothyroxine CAPS (generic Tirosint) THYROLAR TAB (liotrix) THYQUIDITY (levothyroxine) SOLN TIROSINT CAPS (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
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 ER (generic Pentasa) <sup>NR</sup> mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	<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>Asacol HD®/Delzicol DR®/Pentasa®: Requires clinical reason why preferred mesalamine products cannot be used</li> <li>Giazo®: Requires clinical reason why generic balsalazide cannot be used</li> </ul>
REC	TAL	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
MYFEMBREE (relugolix/ estradiol/ norethindrone acetate) <sup>AL, CL,QL</sup> ORIAHNN (elagolix/ estradiol/ norethindrone) <sup>AL,CL</sup> ORILISSA (elagolix sodium) <sup>QL,CL</sup>		Myfembree, Orilissa, and     Oriahnn: Requires an FDA     approved indication, must     follow FDA dosing guidelines,     and have had a trial and failure     of an NSAID and oral     contraceptive

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## **VASODILATORS, CORONARY**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate TAB isosorbide dinitrate ER, SA TAB (generic Dilatrate-SR/Isordil) isosorbide mono IR/SR TAB nitroglycerin SUBLINGUAL, TRANSDERMAL nitroglycerin ER TAB	BIDIL (isosorbide dinitrate/ hydralazine) <sup>CL</sup> GONITRO (nitroglycerin) isosorbide dinitrate <b>TAB</b> (Oceanside Pharm MFR only) isosorbide dinitrate/hydralazine (Bidil) <sup>CL,NR</sup> NITRO-BID OINT (nitroglycerin) NITRO-DUR (nitroglycerin) nitroglycerin <b>TRANSLINGUAL</b> (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>