



PDL Updated October 1, 2021 Highlights indicated change from previous posting For the most up to date list of covered drugs consult the Drug Lookup on the Nebraska Medicaid Website at <a href="https://druglookup.fhsc.com/druglookupweb/?client=nestate">https://druglookup.fhsc.com/druglookupweb/?client=nestate</a>

• **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

- 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

For a complete list of Claims Limitations visit: <a href="https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf">https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf</a>

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ACNE AGENTS, TOPICAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benzoyl peroxide (BPO) WASH, LOTION  clindamycin/BPO (generic Duac) clindamycin phosphate PLEDGET clindamycin phosphate SOLUTION DIFFERIN LOTION, CREAM, Rx-GEL (adapalene) DIFFERIN GEL (adapalene) OTC erythromycin SOLUTION erythromycin-BPO (generic for Benzamycin) PANOXYL 10% WASH (BPO) OTC RETIN-A (tretinoin) <sup>AL</sup> CREAM, GEL	adapalene (generic differin) adapalene/BPO (generic Epiduo)  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) NR benzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide GEL OTC benzoyl peroxide GEL RX benzoyl peroxide TOWELETTE OTC clindamycin FOAM, LOTION clindamycin GEL clindamycin/BPO (generic Acanya, Benzaclin) GEL clindamycin/BPO (generic Veltin, Ziana) dapsone (generic Aczone) EPIDUO FORTE GEL PUMP  (adapalene/BPO) erythromycin-BPO (generic for Benzamycin) EVOCLIN (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (sulfacetamide sodium) PLIXDA (adapalene) SWAB RETIN-A GEL, CREAMAL (tretinoin) sulfacetamide sulfacetamide/sulfur SUMADAN (sulfacetamide/sulfur) tazarotene FOAM (generic Fabior) NR TRETIN-X (tretinoin) tretinoin CREAM, GELAL (generic Avita, Retin-A) tretinoin microspheres (generic for Retin-A Micro) AL	Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class  ### Class    **The company of the patients of the patie

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PDL Updated October 1, 2021 Highlights indicated change from previous posting **ALZHEIMER'S AGENTS** 

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

deliver 20mg or 25mg)

NAMENDA (memantine)

NAMZARIC (memantine/donepezil)

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANALGESICS, OPIOID LONG-ACTING** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) <sup>QL</sup> PATCH fentanyl 25, 50, 75, 100 mcg PATCH <sup>QL</sup> morphine ER TABLET (generic MS Contin, Oramorph SR) OXYCONTIN <sup>CL</sup> (oxycodone ER) tramadol ER (generic Ultram ER) <sup>CL</sup>	ARYMO ER (morphine sulfate) DL BELBUCA (buprenorphine) DL BUCCAL (generic for Belbuca) AL, NR, QL buprenorphine PATCH (generic Butrans) DL EMBEDA (morphine sulfate/naltrexone)  DURAGESIC MATRIX (fentanyl) DL fentanyl 37.5, 62.5, 87.5 mcg PATCH DL hydrocodone ER (generic for Hysingla ER) NR, QL hydrocodone ER (generic for Zohydro ER)  hydromorphone ER (generic for Exalgo) L HYSINGLA ER (hydrocodone ER)  KADIAN (morphine ER)  methadone TABLET, ORAL SYRNR, CL MORPHABOND ER (morphine sulfate)  morphine ER (generic for Avinza, Kadian) CAPSULE  NUCYNTA ER (tapentadol) CL oxycodone ER (generic Oxycontin) oxymorphone 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

PDL Updated October 1, 2021  $\frac{\text{Highlights}}{\text{Highlights}}$  indicated change from previous posting ANALGESICS, OPIOID SHORT-ACTING QL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OR	AL	Non-preferred agents will be
acetaminophen/codeine ELIXIR,     TABLET codeine TABLET hydrocodone/APAP SOLUTION,     TABLET hydrocodone/ibuprofen hydromorphone TABLET morphine CONC SOLUTION,     SOLUTION, TABLET oxycodone TABLET, SOLUTION oxycodone/APAP Tramadol 50 TABLETAL (generic     Ultram) tramadol/APAP (generic Ultracet)	APADAZ (benzhydrocodone/APAP) <sup>CL</sup> benzhydrocodone/APAP (generic Apadaz <sup>,CL</sup> butalbital/caffeine/APAP/codeine butalbital compound w/codeine   (butalbital/ASA/caffeine/codeine) carisoprodol compound-codeine   (carisoprodol/ASA/codeine) dihydrocodeine/APAP/caffeine dihydrocodeine/APAP/caffeine 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> OXAYDO (oxycodone) <sup>CL</sup> oxycodone/APAP SOLUTION oxycodone/APAP SOLUTION oxycodone/APAP TABLET (generic Prolate) oxycodone/APAP TABLET (generic Prolate) oxycodone/Ibuprofen oxymorphone IR (generic Opana) pentazocine/naloxone PROLATE SUSP   (oxycodone/acetaminophen) <sup>NR</sup> ROXICODONE TABLET (oxycodone) tramadol 100mg TABLET(generic Ultram) <sup>AL</sup> ROXYBOND (oxycodone/APAP)	<ul> <li>approved for patients who have failed THREE preferred agents within this drug class within the last 12 months</li> <li>Note: for short acting opiate tablets and capsules there is a maximum quantity limit of #150 per 30 days.</li> <li>Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will consist of -prescriptions limited to a 7 day supply, AND -initial opiate prescription fill limited to maximum of 50 Morphine</li> </ul>

PDL Updated October 1, 2021 Highlights indicated change from previous posting ANALGESICS, OPIOID SHORT-ACTING<sup>QL</sup> (Continued)

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

# ANDROGENIC AGENTS (Topical)CL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANDROGEL (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®: Approved for Males only</li> <li>Natesto®: Approved for Males or with diagnosis of: Primary hypogonadism (congenitor acquired) OR</li></ul></li></ul>

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# **ANGIOTENSIN MODULATORS**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting ANGIOTENSIN MODULATORS (Continued)

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

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

#### ANTI-ALLERGENS, ORAL

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

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PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIBIOTICS**, **GASTROINTESTINAL** 

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

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Preferred Agents <sup>CL</sup>	Non-Preferred Agents	Prior Authorization/Class Criteria
BETHKIS (tobramycin) <sup>CL</sup> KITABIS PAK (tobramycin) <sup>CL</sup> TOBI-PODHALER (tobramycin) <sup>CL,QL</sup>	ARIKAYCE (amikacin liposomal inh) <sup>CL</sup> SUSPENSION CAYSTON (aztreonam lysine) <sup>QL,CL</sup> tobramycin (generic for Bethkis) tobramycin (generic Tobi) <sup>CL</sup>	<ul> <li>Diagnosis of Cystic Fibrosis is required for all agents         ICD10 Group = E84, ICD9 = 277.00, 277.01, 277.02, 277.03, 277.09</li> <li>Drug-specific criteria:         <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> </ul> </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 OINTMENT bacitracin/polymyxin (generic Polysporin) mupirocin OINTMENT (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/ pramoxine	CENTANY (mupirocin) gentamicin <b>OINTMENT, CREAM</b> mupirocin <b>CREAM</b> (generic Bactroban) <sup>CL</sup>	Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class within the last 12 months  Drug-specific criteria:     Mupirocin® Cream: Clinical reason the ointment cannot be used

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CLEOCIN <b>OVULES</b> (clindamycin) clindamycin <b>CREAM</b> (generic Cleocin) CLINDESSE (clindamycin) NUVESSA (metronidazole) VANDAZOLE (metronidazole)	CLEOCIN <b>CREAM</b> (clindamycin) METROGEL (metronidazole) metronidazole, vaginal	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 las 6 months

#### **ANTICOAGULANTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg <sup>CL,QL</sup> XARELTO DOSE PACK (rivaroxaban)	BEVYXXA (betrixaban) <sup>QL</sup> fondaparinux (generic Arixtra) FRAGMIN (dalteparin) SAVAYSA (edoxaban) <sup>QL</sup>	<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:         <ul> <li>Coumadin®: Clinical reason generic warfarin cannot be used</li> <li>Savaysa®: Approved diagnoses include:</li></ul></li></ul>

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# **ANTIEMETICS/ANTIVERTIGO AGENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CANNA	BINOIDS	Non-preferred agents will be
dronabinol (generic Marinol) <sup>AL</sup>	CESAMET (nabilone)	approved for patients who have failed ONE preferred agent within this drug class within the same
5HT3 RECEPT	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
NK-1 RECEPTO	R ANTAGONIST	5-HT3 antagonist     Regimens include: AC combination
EMEND (aprepitant) CAPSULE, CAPSULE PACK <sup>QL</sup>	aprepitant (generic Emend) QL,CL AKYNZEO (netupitant/palonosetron)CL VARUBI (rolapitant) <b>TABLET</b> CL	(Doxorubicin or Epirubicin with Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine, Carbplatin, Cisplatin, Clofarabine,
TRADITIONAL	ANTIEMETICS	Cyclophosphamide, Cytarabine, Dacarbazine, Dactinomycin, Daunorubicin,
DICLEGIS (doxylamine/pyridoxine) <sup>CL,QL</sup> dimenhydrinate (generic Dramamine) OTC meclizine (generic Antivert) metoclopramide (generic Reglan) phosphoric acid/dextrose/fructose SOLUTION (generic Emetrol) prochlorperazine, oral (generic Compazine) promethazine 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>Epirubicin, Etoposide, 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 ora dosage form intolerance</li> </ul>

PDL Updated October 1, 2021 Highlights indicated change from previous posting ANTIFUNGALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche) fluconazole SUSPENSION, TABLET (generic Diflucan) griseofulvin SUSPENSION griseofulvin microsized TABLET nystatin SUSPENSION, TABLET terbinafine (generic Lamisil)	BREXAFEMME (ibrexafungerp) <sup>QL,NR</sup> CRESEMBA (isavuconazonium) <sup>CL</sup> flucytosine (generic Ancobon) <sup>CL</sup> griseofulvin ultramicrosize (generic GRIS-PEG) itraconazole (generic Sporanox) <sup>CL</sup> ketoconazole (generic Nizoral) nystatin <b>POWDER</b> ONMEL (itraconazole) ORAVIG (miconazole) posaconazole (generic Noxafil) <sup>AL,CL</sup> TOLSURA (itraconazole) <sup>CL</sup> voriconazole (generic VFEND) <sup>CL</sup>	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class</li> <li>Drug-specific criteria:</li> <li>Cresemba®: Approved for diagnosis of invasive aspergillosis or invasive mucomycosis</li> <li>Flucytosine: Approved for diagnosis of: Candida: Septicemia, endocarditis, UTIs Cryptococcus: Meningitis, pulmonary infections</li> <li>Noxafil®: No trial for diagnosis of Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropenic hematologic malignancies, Graft vs. Host disease(GVHD), Immunosuppression secondary to hematopoietic stem cell transplant</li> <li>Noxafil® Suspension: Oropharyngeal/esophageal candidiasis refractory to itraconazole and/or fluconazole</li> <li>Onmel®: Requires trial and failure or contraindication to terbinafine</li> <li>Sporanox®/Itraconazole: Approved for diagnosis of Aspergillosis, Blastomycosis, Histoplasmosis, Onychomycosis due to terbinafine-resistant dermatophytes, Oropharyngeal/ esophageal candidiasis refractory to fluconazole</li> <li>Sporanox®: Requires trial and failure of generic itraconazole</li> <li>Sporanox®: Requires trial and failure of generic itraconazole</li> <li>Sporanox®: Requires trial and failure of generic itraconazole</li> <li>Vforanos Liquid: Clinical reason solid oral cannot be used</li> <li>Tolsura: Approved for diagnosis of Aspergillosis, Blastomycosis, and Histoplasmosis and requires a trial and failure of generic itraconazole</li> <li>Vfend®: No trial for diagnosis of Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Graft vs. Host disease (GVHD), Candidemia (candida krusei), Esophageal Candidiasis, Blastomycosis, S. apiospermum and Fusarium spp., Oropharyngeal/esophageal candidiasis</li> </ul>

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PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIHISTAMINES, MINIMALLY SEDATING** 

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

### **ANTIHYPERTENSIVES, SYMPATHOLYTICS**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIHYPERURICEMICS** 

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIMIGRAINE AGENTS**, **OTHER** 

AJOVY (fremanezumab-vfrm) CL, QL PEN, Autoinjector, Autoinjector 3-packNR  EMGALITY 120 mg/mL (galcanezumab- gnlm) CL, QL PEN, SYRINGE  UBRELVY (ubrogepant)^AL, CL, QL TABLET  Almovig (erenumab-aooe) CL, QL ergotamine mesylate NASAL ergotamine tartrate) MIGEROT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL NURTEC ODT (rimegepant)^AL, CL, QL TABLET  PABLET  Almovig, Ajovy and Emgality 120mg: Require ≥ 4 migraines per month for ≥ 3 months and has tried and failed a ≥ 1 month trial of two medications listed in the 2012 American Academy of Neurology/American Aeademy of Neurology/American Headache Society guidelines (examples include: antidepressants (amitriptyline, venlafaxine), Beta blockers (propraolo), atti-option), atti-option, atti-option, atti-option, atti-option, and in moving requires a trial of
Emgality 120mg or Ajovy or clinical, patient specific reason that a preferred agent cannot be used

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIMIGRAINE AGENTS**, **TRIPTANS**<sup>QL</sup>

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANTIPARASITICS**, **TOPICAL** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NATROBA (spinosad) permethrin 1% OTC (generic Nix) permethrin 5% RX (generic Elimite) pyrethrin/piperonyl butoxide (generic RID, A-200)	CROTAN (crotamiton) LOTION EURAX (crotamiton) CREAM, LOTION ivermectin (generic Sklice) LOTION NR lindane malathion (generic Ovide) SKLICE (ivermectin) 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

PDL Updated October 1, 2021 Highlights indicated change from previous posting

# **ANTIPARKINSON'S AGENTS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class
ANTICHOLINERGICS	<ul> <li>Non-preferred agents will be</li> </ul>	A 11 -
benztropine (generic for Cogentin) trihexyphenidyl (generic for Artane)		
COMT INHIBITORS	•	
	entacapone (generic for Comtan) • ONGENTYS (Opicapone) <sup>NR,QL</sup> tolcapone (generic for Tasmar)	
DOPAMINE AGONISTS	•	
pramipexole (generic for Mirapex) ropinirole (generic for Requip)	bromocriptine (generic for Parlodel) ropinirole ER (generic for Requip ER) <sup>CL</sup> NEUPRO (rotigotine) <sup>CL</sup> pramipexole ER (generic for Mirapex ER) <sup>CL</sup> ropinirole ER (generic for Requip XL) <sup>CL</sup> ropinirole ER (generic for Requip XL) <sup>CL</sup>	
MAO-B INHIBITORS	•	
selegiline CAPSULE, TABLET (generic for Eldepryl)	rasagiline (generic for Azilect) QL XADAGO (safinamide) ZELAPAR (selegiline)CL	

# ANTIPARKINSON'S AGENTS, ORAL

(continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class
	•	
OTHER ANTIPARKINSON'S DRUGS		

PDL Updated October 1, 2021 Highlights indicated change from previous posting

# **ANTIPSORIATICS, ORAL**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting ANTIPSORIATICS, TOPICAL

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

#### **ANTIVIRALS, ORAL**

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

#### **ANTIVIRALS, TOPICAL**

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

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **ANXIOLYTICS** 

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **BETA BLOCKERS, ORAL** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
•	acebutolol (generic Sectral) betaxolol (generic Kerlone) BYSTOLIC (nebivolol) HEMANGEOL (propranolol) SOLUTION INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) LEVATOL (penbutolol) metoprolol/HCTZ (generic	<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> </ul> </li> </ul>
	Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide nebivolol (generic Bystolic) <sup>NR</sup> pindolol (generic Viskin) propranolol/HCTZ (generic Inderide) timolol (generic Blocadren) TOPROL XL (metoprolol ER)	Sotylize®: Covered for diagnosis of life –threatening ventricular arrhythmias OR maintenance of normal sinus rhythm in highly symptomatic atrial fibrillation/flutter (AFIB/AFL)  Requires clinical reason generic sotalol cannot be used
BETA- AND ALI	PHA-BLOCKERS	
carvedilol (generic Coreg) labetalol (generic Trandate)	carvedilol ER (generic Coreg CR)	
ANTIARR	HYTHMIC	
sotalol (generic Betapace)	SOTYLIZE (sotalol)	

#### **BILE SALTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ursodiol <b>CAPSULE</b> 300mg (generic for Actigall) ursodiol 250mg <b>TABLET</b> (generic for URSO) ursodiol 500mg <b>TABLET</b> (generic for URSO FORTE)	CHENODAL (chenodiol) CHOLBAM (cholic acid) OCALIVA (obeticholic acid)	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

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

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# **BLADDER RELAXANT PREPARATIONS**

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

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#### BONE RESORPTION SUPRESSION AND RELATED DRUGS

alendronate (generic Fosamax) TABLET ibandronate (generic Boniva) ibandronate (generic Boniva)  ATELVIA DR (risedronate) BINOSTO (alendronate) etidronate disodium (generic Didronate) FOSAMAX PLUS DQL risedronate (generic Actonel)  OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS  Calcitonin-salmon NASAL raloxifene (generic Evista) teriparatide (generic Forteo)  CL,QL  TYMLOS (abaloparatide)	individual agents without prior authorization  • Atelvia DR®: Requires clinical reason
TABLET  ibandronate (generic Boniva) <sup>QL</sup> ibandronate (generic Boniva) <sup>QL</sup> ATELVIA DR (risedronate)  BINOSTO (alendronate)  etidronate disodium (generic Didrone FOSAMAX PLUS D <sup>QL</sup> risedronate (generic Actonel) <sup>QL</sup> OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS  calcitonin-salmon NASAL  aloxifene (generic Evista)  FORTEO (teriparatide) <sup>CL,QL</sup>	failed a trial of ONE preferred agent within the same group  Drug-specific criteria:  I) • Actonel® Combinations: Covered as individual agents without prior authorization  • Atelvia DR®: Requires clinical reason
	alendronate cannot be taken on an empty stomach  Binosto®: Requires clinical reason why alendronate tablets OR Fosamax® solution cannot be used  Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification  Forteo®: Covered for high risk of fracture High risk of fracture:  BMD -3 or worse  Postmenopausal women with history of non-traumatic fractures  Postmenopausal women with 2 or more clinical risk factors  Family history of non-traumatic fractures  DXA BMD T-score ≤ -2.5 at any site  Glucocorticoid use ≥ 6 months at 7.5 dose of prednisolone equivalent  Rheumatoid Arthritis  Postmenopausal women with BMD T-score ≤ -2.5 at any site with any clinical risk factors  More than 2 units of alcohol per day  Current smoker  Men with primary or hypogonadal osteoporosis  Osteoporosis associated with sustained systemic glucocorticoid

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**BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALPHA BLOCKERS		Non-preferred agents will be
alfuzosin (generic Uroxatral) doxazosin (generic Cardura) tamsulosin (generic Flomax) terazosin (generic Hytrin)	CARDURA XL (doxazosin) silodosin (generic Rapaflo)	approved for patients who have failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:
5-ALPHA-REDUCTA	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
INHALERS – Short Acting		Non-preferred agents will
PROAIR HFA (albuterol)  INHALI SEREVENT (salmeterol)	albuterol HFA (generic for ProAir HFA, Proventil HFA, Ventolin HFA) levalbuterol HFA (generic for Xopenex HFA) PROAIR DIGIHALER (albuterol) PROAIR RESPICLICK (albuterol) PROVENTIL HFA (albuterol)  ERS - Long Acting  ARCAPTA NEOHALER (indacaterol)	be approved for patients who have failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:  • Xopenex®: Covered for cardiac diagnoses or side effect of tachycardia with albuterol product
INHAL	STRIVERDI RESPIMAT (olodaterol)  ATION 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) <sup>NR</sup> BROVANA (arformoterol) formoterol fumarate (generic Performist) <sup>NR</sup> 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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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SHORT-ACTING Dihydropyridines		Non-preferred agents will be approved for patients who have
diltiazem (generic Cardizem) verapamil (generic Calan/Isoptin) LONG-	isradipine (generic Dynacirc) nicardipine (generic Cardene) nifedipine (generic Procardia) nimodipine (generic Nimotop) NYMALIZE (nimodipine) SOLUTION ropyridines  ACTING byridines felodipine ER (generic Plendil) KATERZIA (amlodipine) nisoldipine (generic Sular)	<ul> <li>failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:         <ul> <li>Nifedipine: May be approved without trial for diagnosis of Preterm Labor or Pregnancy Induced Hypertension (PIH)</li> <li>Nimodipine: Covered without trial for diagnosis of subarachnoid hemorrhage</li> <li>Katerzia: May be approved with documented swallowing difficulty</li> </ul> </li> </ul>
Non-dihyd	ropyridines	
diltiazem ER (generic Cardizem CD) verapamil ER <b>TABLET</b>	CALAN SR (verapamil) diltiazem ER (generic Cardizem LA) MATZIM LA (diltiazem ER) TIAZAC (diltiazem) verapamil ER CAPSULE verapamil 360mg CAPSULE verapamil ER (generic Verelan PM)	

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

# **COLONY STIMULATING FACTORS**

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

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# **CONTRACEPTIVES, ORAL**

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

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# COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANORO ELLIPTA (umeclidinium/vilanterol) ATROVENT HFA (ipratropium) COMBIVENT RESPIMAT (albuterol/ipratropium) SPIRIVA (tiotropium) STIOLTO RESPIMAT (tiotropium/olodaterol)	BEVESPI AEROSPHERE (glycopyrolate/formoterol)  DUAKLIR PRESSAIR (aclidinium br and formoterol fum) INCRUSE ELIPTA (umeclidnium) SEEBRI NEOHALER (glycopyrolate) SPIRIVA RESPIMAT (tiotropium) TUDORZA PRESSAIR (aclidinium br) UTIBRON NEOHALER (indacaterol/glycopyrolate)	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.  Drug-specific criteria:  Daliresp®:  Covered for diagnosis of severe COPD associated with chronic bronchitis  Requires trial of a bronchodilator Requires documentation of one exacerbation in last year upon
albuterol/ipratropium (generic for Duoneb) ipratropium <b>SOLUTION</b> (generic for Atrovent)	N SOLUTION  LONHALA (glycopyrrolate inhalation soln)  YUPELRI (revefenacin)  AGENT  DALIRESP (roflumilast) <sup>CL, QL</sup>	initial review

#### **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 pseudoephedrine/codeine/ guaifenesin (generic for Lortuss EX, Tusnel C, Virtussin DAC)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE dextromethorphan product</li> <li>All codeine or hydrocodone containing cough and cold combinations are limited to &gt; 18 years of age</li> </ul>

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# CYSTIC FIBROSIS, ORAL

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ENBREL (etanercept) KIT, MINI CART, PENQL HUMIRA (adalimumab)QL ENBREL (etanercept) VIALQL TEZLA (apremilast) ORAL CL,QL	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIMZIA (certolizumab pegol) <sup>QL</sup> COSENTYX (secukinumab) ENSPRYNG (satralizumab-mwge) SUB-Q ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYRINGE KINERET (anakinra) OLUMIANT (baricitinib) ORAL <sup>CL,QL</sup> ORENCIA (abatacept) SUB-Q RINVOQ ER (upadacitinib, <sup>CL,QL</sup> SILIQ (brodalumab) SIMPONI (golimumab) SKYRIZI (risankizamab-rzaa) SKYRIZI PEN (risankizamab-rzaa) SKYRIZI PEN (risankizamab) SUB-Q TALTZ (ixekizumab) <sup>AL</sup> TREMFYA (guselkumab) <sup>QL</sup> XELJANZ (tofacitinib) ORAL, SOLN <sup>CL,QL</sup> XELJANZ XR (tofacitinib) ORAL	<ul> <li>Preferred agents will be approved with FDA-approved indication – ICD-10 diagnosis code is required.</li> <li>Non-preferred agents will be approved for FDA-approved indications in patients who have failed a trial of ONE preferred agent within this drug class, or upon diagnosis for non-preferred agent with FDA-approved indication if no preferred agent has FDA approval for diagnosis.</li> <li>Drug-specific criteria:         <ul> <li>Otezla: Requires a trial of Humira</li> <li>Olumiant: Requires documentation of inadequate response or intolerance to methotrexate and an inadequate response to one or more TNF antagonist therapies.</li> <li>Rinvoq: Requires documentation of inadequate response or intolerance to methotrexate</li> <li>Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to methotrexate. Diagnosis of Juvenile Idiopathic Arthritis for ages 2 years old and older does not require documentation of treatment failure with methotrexate. Diagnosis of moderate to severe ulcerative colitis (UC) requires documentation of treatment failure with a Tumor Necrosis Factor blocker agent; does not require documentation of treatment failure with methotrexate.</li> </ul> </li> </ul>

PDL Updated October 1, 2021 Highlights indicated change from previous posting **DIURETICS** 

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

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

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

# EPINEPHRINE, SELF-INJECTEDQL

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

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#### **ERYTHROPOIESIS STIMULATING PROTEINS**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting

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

## **FLUOROQUINOLONES, ORAL**

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AMITIZA (lubiprostone) <sup>AL, QL</sup> LINZESS (linaclotide) <sup>QL</sup> MOVANTIK (naloxegol oxalate) <sup>QL</sup>	alosetron (generic Lotronex)  Iubiprostone (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>Drug-specific criteria:</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 <b>SUSP</b> (generic Proglycem) GLUCAGON EMERGENCY (glucagon) <sup>QL</sup> <b>INJ KIT</b> (Fresenius) GVOKE (glucagon) <sup>AL,QL</sup> <b>PEN</b> , <b>SYRINGE</b>	<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>

PDL Updated October 1, 2021 Highlights indicated change from previous posting **GLUCOCORTICOIDS, INHALED** 

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

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### **GLUCOCORTICOIDS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ALKINDI (hydrocortisone)  GRANULES <sup>AL/NR</sup> CORTEF (hydrocortisone) cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSPENSION, TABLET <sup>CL</sup> ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg, 32mg ORTIKOS ER (budesonide) <sup>AL,QL</sup> PEDIAPRED (prednisolone sodium phosphate) prednisolone sodium phosphate (generic for Millipred/Veripred) prednisolone sodium phosphate ODT prednisone SOLUTION prednisone INTENSOL RAYOS DR (prednisone) TABLET	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agents within this drug class within the last 6 months</li> <li>Drug-specific criteria:</li> <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>

#### **GROWTH HORMONES**

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

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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>	<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>

#### HAE TREATMENTSCL

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

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#### **HEMOPHILIA TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
F/	ACTOR VIII	<ul> <li>Non-preferred agents will be</li> </ul>
ALPHANATE HELIXATE FS HUMATE-P	ADVATE ADYNOVATE AFSTYLA	approved for patients who have failed a trial of ONE preferred agent within this drug class
NOVOEIGHT NUWIQ XYNTHA KIT, SOLOFUSE	ELOCTATE ESPEROCT HEMOFIL-M JIVI <sup>AL</sup> KOATE-DVI KIT KOATE-DVI VIAL KOGENATE FS KOVALTRY OBIZUR RECOMBINATE	<ul> <li>Patients receiving a hemophilia agent which moved from preferred to non-preferred status on 1-21-21 will be allowed to continue same therapy</li> </ul>
F	ACTOR IX	
BENEFIX	ALPHANINE SD ALPROLIX IDELVION IXINITY MONONINE PROFILNINE SD REBINYN RIXUBIS	
	OMBIN COMPLEX-PLASMA DERIVED	
NOVOSEVEN RT	FEIBA NF SEVENFACT <sup>AL,NR</sup>	
FACTOR X	AND XIII PRODUCTS	
COAGADEX CORIFACT	TRETTEN	
VON WILLE	BRAND PRODUCTS	
WILATE	VONVENDI	
	CIFIC FACTORS	
HEMLIBRA		

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#### **HEPATITIS B TREATMENTS**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **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) <sup>CL</sup> VOSEVI (sofosbuvir/velpatasvir/ voxilaprevir) <sup>CL</sup>	HARVONI 200/45MG, TABLET,	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
RIBA	VIRIN	1 or 4  Vosevi: Requires documentation of non-
	REBETOL (ribavirin)	response after previous treatment course of Direct Acting Anti-viral agent (DAA) for
INTER	FERON	genotype 1-6 without cirrhosis or with compensated cirrhosis
PEGASYS (pegylated interferon alfa- 2a) <sup>CL</sup> PEG-INTRON (pegylated interferon alfa-2b) <sup>CL</sup>		Compensated crimosis

PDL Updated October 1, 2021 Highlights indicated change from previous posting

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

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting HIV / AIDSCL

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

PDL Updated October 1, 2021  $\frac{\text{Highlights}}{\text{HIV}}$  indicated change from previous posting  $\frac{\text{HIV}}{\text{AIDS}^{\text{CL}}}$  (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atazanavir CAPSULE (generic Reyataz) LEXIVA SUSP (fosamprenavir) ritonavir TABLET (generic Norvir)	INHIBITORS	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

PDL Updated October 1, 2021 Highlights indicated change from previous posting HIV / AIDSCL (Continued)

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

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting HIV / AIDS<sup>CL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COMBINATION PRODU	CTS – MULTIPLE CLASSES	
ATRIPLA (tenofovir/emtricitabine/efavirenz)  BIKTARVY (bictegravir/emtricitabine/tenofovir)QL  COMPLERA (rilpivirine/emtricitabine/tenofovir)  DELSTRIGO (doravirine/lamivudine/tenofovir)QL  GENVOYA (elvitegravier/cobicistat/emtricitabine/tenofovir)QL, AL  ODEFSEY (emtricitabine/rilpivirine/tenofovir)QL  STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir)QL  SYMFI (efavirenz/lamivudine/tenofovir)QL  SYMFI LO (efavirenz/lamivudine/tenofovir)QL  TRIUMEQ (dolutegravir/abacavir/lamivudine)	SYMTUZA (darunavir/cobicistat/ emtricitabine/tenofovir) <sup>QL</sup>	class are not appropriate for

PDL Updated October 1, 2021 Highlights indicated change from previous posting HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting

# HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting 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 R U-500 KWIKPENCL HUMULIN 70/30 OTC PEN INSULIN ASPART (generic for Novolog) INSULIN ASPART (generic for Novolog) INSULIN ASPART (generic for Humalog) PEN, VIAL (generic for Humalog) PEN, VIAL, JR KWIKPEN INSULIN IISPRO (Humalog Mix Kwikpen) LANTUS SOLOSTAR PEN (INSULIN glargine) LANTUS (INSULIN glargine) VIAL LEVEMIR (INSULIN detemir) PEN, VIAL 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 FIASP (insulin aspart) CARTRIDGE, PEN, VIAL HUMALOG (insulin lispro) U-200 KWIKPEN LYUMJEV KWIKPEN, VIAL(insulin lispro-aabc) NOVOLIN (insulin) NOVOLIN 70/30 VIAL(insulin) TOUJEO SOLOSTAR (insulin glargine) SEMGLEE (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) repaglinide/metformin (generic for Prandimet)	Non-preferred agents will be approved for patients with:     Failure of a trial of ONE preferred agent in another Hypoglycemic class OR     T2DM and inadequate glycemic control

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#### HYPOGLYCEMICS, METFORMINS

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

#### **HYPOGLYCEMICS, SGLT2**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) <sup>QL,CL</sup> INVOKAMET (canagliflozin/metformin) <sup>QL, CL</sup> INVOKANA (canagliflozin) <sup>CL</sup> JARDIANCE (empagliflozin) <sup>QL, CL</sup> SYNJARDY (empagliflozin/metformin) <sup>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>QL</sup>	<ul> <li>Preferred agents are Approved for diagnosis of diabetes AND a trial of metformin</li> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> </ul>

#### HYPOGLYCEMICS, SULFONYLUREAS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
glimepiride (generic Amaryl) glipizide IR & ER (generic Glucotrol/ Glucotrol XL) glyburide (generic Diabeta/Glynase)	chlorpropamide tolazamide tolbutamide	<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>
SULFONYLURE	COMBINATIONS	
glipizide/metformin glyburide/metformin (generic Glucovance)		

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
THIZAOLIDINE	THIZAOLIDINEDIONES (TZDs)	
pioglitazone (generic for Actos)	AVANDIA (rosiglitazone)	approved for patients who have failed a trial of THE preferred agent
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)	<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>

#### IMMUNOMODULATORS, ASTHMACL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ASENRA (benralizumab) <sup>AL</sup> <b>PEN</b>	NUCALA (mepolizumab) <sup>AL</sup> AUTO-INJ, SYR,  XOLAIR (omalizumab) SYR <sup>AL,NR,QL</sup>	Drug Specific Criteria:  Dupixent: See criteria listed under Immunomodulator, Atopic Dermatitis class  Fasenra: is indicated for patient 12 years and older for add on maintenance treatment of severe asthma, and with an eosinophilic phenotype  Nucala: is indicated for  -Patients 6 years and older for add on maintenance treatment of severe asthma, and with an eosinophilic phenotype  -Patients 12 years and older with hypereosinophilic syndrome (HES) for ≥6 months without identifiable non-hematologic secondary cause  -Adult patients with eosinophilic granulomatosis with polyangiit

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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## IMMUNOMODULATORS, ATOPIC DERMATITISAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIDEL (pimecrolimus) EUCRISA (crisaborole) <sup>CL,QL</sup>	DUPIXENT (dupilumab) <sup>AL,CL</sup> DUPIXENT <b>PEN<sup>AL</sup></b> Opzelura (ruxolitinib phosphate) CREAM <sup>AL,NR,QL</sup> pimecrolimus (generic for Elidel) tacrolimus (generic for Protopic) <sup>CL</sup>	<ul> <li>Non-preferred agents require: Trial of a topical steroid AND Trial of one preferred product within this drug class</li> <li>Drug-specific criteria:</li> <li>Dupixent: Indicated for moderate to severe atopic dermatitis, must have trial of Eucrisa; For moderate to severe asthma, must have eosinophilic phenotype or oral corticosteroid dependent asthma uncontrolled with maintenance controller medication; For adults with chronic rhinosinusitis with nasal polyposis, must document inadequate control on current treatment regimen and be used as add-on maintenance treatment with intranasal steroid</li> <li>Eucrisa: Requires use and failure of 1 topical steroid or Elidel.</li> </ul>

### **IMMUNOMODULATORS, TOPICAL**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting **IMMUNOSUPPRESSIVES, ORAL** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathiaprine (generic Imuran) cyclosporine, modified CAPSULE (generic Neoral) mycophenolate CAPSULE, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLUTION RAPAMUNE (sirolimus) TABLET tacrolimus ZORTRESS (everolimus) AL	ASTAGRAF XL (tacrolimus) AZASAN (azathioprine) cyclosporine CAPSULE, SOFTGEL cyclosporine, modified SOLUTION   (generic Neoral) ENVARSUS XR (tacrolimus) GENGRAF (cyclosporine, modified)   CAPSULE, SOLUTION mycophenolate SUSPENSION   (generic Cellcept) mycophenolic acid MYFORTIC (mycophenolate sodium) PROGRAF (tacrolimus) CAPSULE,   PACKET SANDIMMUNE (cyclosporine)   CAPSULE, SOLUTION sirolimus SOLUTION, TABLET (generic Rapamune) everolimus (generic for Zortress) 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

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#### **INTRANASAL RHINITIS DRUGS**

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

#### **LEUKOTRIENE MODIFIERS**

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

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#### LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

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

#### LIPOTROPICS, OTHER

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

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PROPROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 9 (PCSK9) INHIBITORS  PRALUENT (alorocumab) <sup>CL</sup> REPATHA (evolocumab) <sup>CL</sup> ** atherosclerotic cardiovascular disease (ASCVD)  ** heterozygous familial hypercholesterolemia (HoFH) as an adjunct to other LDL-C lowering therapies  ** AND  ** Maximized high-intensity statin WITH ezetimibe for at 3 continuous months  ** Failure to reach target LDL-C levels: ASCVD < 70 mg/dL, HeFH < 100 mg/dL  ** Repatha**: Approved for:  ** adult diagnoses of atherosclerotic cardiovascular disease (ASCVD)  ** heterozygous familial hypercholesterolemia (HoFH) in age ≥ 13  ** statin-induce rhabdomyolysis AND  ** Maximized high-intensity statin WITH ezetimibe for 3+ continuous months  ** Failure to reach target LDL-C levels: ASCVD < 70 mg/dL, HeFH < 100 mg/dL	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
● Concurrent use of maximally-tolerated statin must continue	PROPROTEIN CONVERTASE SUI	BTILISIN/KEXIN TYPE 9 (PCSK9) BITORS PRALUENT (alorocumab) <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</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:         <ul> <li>Altoprev®: One of the TWO trials must be IR lovastatin</li> </ul> </li> <li>Combination products: Require clinical</li> </ul>
STATIN COMBINATIONS		reason why individual ingredients cannot be
	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
MACR	OLIDES	Require clinical reason why
· · · · · · · · · · · · · · · · · · ·	clarithromycin ER (generic Biaxin XL) E.E.S. SUSPENSION (erythromycin ethylsuccinate) E.E.S. TABLET (erythromycin ethylsuccinate) ERY-TAB (erythromycin) ERYPED SUSPENSION (erythromycin) ERYTHROCIN (erythromycin) erythromycin base TABLET, CAPSULE	preferred products within this drug class cannot be used AND ≥ 3-day trial on a preferred product

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#### **METHOTREXATE**

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

#### **MOVEMENT DISORDERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) <sup>CL</sup> tetrabenazine (generic for Xenazine) <sup>CL</sup>	INGREZZA (valbenazine) <sup>CL</sup> <b>CAP</b> , INITIATION PACK XENAZINE (tetrabenazine) <sup>CL</sup>	Non-preferred agent requires trial of Austedo  All drugs require an FDA approved indication – ICD-10 diagnosis code required.  Drug-specific criteria:  • 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  • Ingrezza: Diagnosis of Tardive Dyskinesia in adults and trial of Austedo  • tetrabenazine:Diagnosis of chorea with Huntington's Disease

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#### **MULTIPLE SCLEROSIS DRUGS**

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

#### **NITROFURAN DERIVATIVES**

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

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# NSAIDs, ORAL

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

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

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

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#### **NSAIDs, TOPICAL**

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

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#### **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 <b>CO-PACK</b> 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
CHEMO	THERAPY	- - Drug-specific critera
cyclophosphamide XELODA (capecitabine)	capecitabine (generic for Xeloda) <sup>CL</sup>	<ul> <li>anastrozole: May be approved for malignant neoplasm of male breast (male breast cancer)</li> </ul>
HORMONE	HORMONE BLOCKADE	
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>	<ul> <li>Xeloda or clinical reason Xeloda cannot be used</li> <li>Fareston®: Require clinical reason why tamoxifen cannot be used</li> <li>letrozole: Approved for diagnosis of breast cancer with day supply greater than 12 – NOT approved</li> </ul>
ОТ	HER	for short term use
	NERLYNX (neratinib) PIQRAY (alpelisib) lapatinib (generic Tykerb) <sup>CL,NR</sup> TALZENNA (talazoparib tosylate) QL TUKYSA(tucatinib) <sup>QL</sup>	<ul> <li>Soltamox: May be approved with documented swallowing difficulty</li> </ul>

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

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

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#### ONCOLOGY AGENTS, ORAL, LUNG

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

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

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
abiraterone (generic for Zytiga) <sup>CL</sup> bicalutamide (generic for Casodex) flutamide XTANDI (enzalutamide) <sup>AL,QL</sup>	EMCYT (estramustine) ERLEADA (apalutamide) <sup>QL</sup> nilutamide (generic for Nilandron) NUBEQA (darolutamide) <sup>QL</sup> YONSA (abiraterone acetonide, submicronized) ZYTIGA (abiraterone) <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> <li>Drug Specific Critieris</li> <li>Zytiga: Patients receiving Zytiga prior to 1/21/21 (which changed from preferred to non-preferred) will be allowed to continue current treatment</li> </ul>

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

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BASAL CELL		<ul> <li>Non-preferred agents DO NOT</li> </ul>
ERIVEDGE (vismodegib)	ODOMZO (sonidegib) <sup>CL</sup>	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
BRAF MUTATION		

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MEKINIST (trametinib) TAFINLAR (dabrafenib) BRAFTOVI (encorafenib) COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)

Drug-specific critera

 Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17-19 will be allowed to continue therapy

#### **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) PAZEO (olopatadine 0.7%)	ALOCRIL (nedocromil) ALOMIDE (lodoxamide) azelastine (generic for Optivar) BEPREVE (bepotastine besilate) bepotastine besilate (generic for Bepreve) <sup>NR</sup> EMADINE (emedastine) epinastine (generic for Elestat) LASTACAFT (alcaftadine) olopatadine 0.1% (generic for Patanol) olopatadine 0.2% (generic for Pataday) PATADAY (olopatadine 0.7%) <sup>NR</sup> 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>SOLUTION</b> (generic for Ciloxan) ofloxacin (generic for Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic for Zymaxid) levofloxacin MOXEZA (moxifloxacin) moxifloxacin (generic for Vigamox) moxifloxacin (generic for Moxeza) VIGAMOX (moxifloxacin)	<ul> <li>approved for patients who have failed a one-month trial of TWO preferred agent within this drug class</li> <li>Azasite®: Approval only requires trial of erythromycin</li> <li>Drug-specific criteria:</li> <li>Natacyn®: Approved for documented fungal infection</li> </ul>
MACROLIDES		
erythromycin	AZASITE (azithromycin) <sup>CL</sup>	
AMINOGLY	YCOSIDES	
gentamicin <b>OINTMENT</b> gentamicin <b>SOLUTION</b> tobramycin (generic for Tobrex drops)	TOBREX <b>OINTMENT</b> (tobramycin)	
OTHER OPHTH	ALMIC AGENTS	_
bacitracin/polymyxin B (generic Polysporin) polymyxin B/trimethoprim (generic for Polytrim)	bacitracin NATACYN (natamycin) <sup>CL</sup> neomycin/bacitracin/polymyxin B OINTMENT neomycin/polymyxin B/gramicidin NEOSPORIN (neomycin/polymyxin B/gramcidin) sulfacetamide SOLUTION (generic for Bleph-10) sulfacetamide OINTMENT	

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

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

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

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

## OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
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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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RESTASIS (cyclosporine)
RESTASIS MULTIDOSE
(cyclosporine)
XIIDRA (lifitegrast)

CEQUA (cyclosporine) QL EYSUVIS (loteprednol etabonate)NR,QL

Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

#### OPHTHALMICS, GLAUCOMA

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIOTICS		Non-preferred agents will be
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide)	approved for patients who have failed a trial of ONE preferred agent within this drug class
SYMPATHO	MIMETICS	
brimonidine 0.2% (generic for Alphagan)	Alphagan P (brimonidine 0.1%) Alphagan P (brimonidine 0.15%) apraclonidine (generic for lopidine)	
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) <sup>NR</sup> TIMOPTIC OCUDOSE TIMOPTIC XE (timolol gel forming solution)	
CARBONIC ANHYDI	·	•
dorzolamide (generic for Trusopt)	AZOPT (brinzolamide) brinzolamide (generic for Azopt) <sup>NR</sup>	•
PROSTAGLAND	OIN ANALOGS	•
latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATION	ON DRUGS	
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine	

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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ОТН	IER	•
RHOPRESSA (netarsudil) <sup>CL</sup> ROCKLATAN (netarsudil and latanoprost) <sup>CL</sup>		Drug-specific criteria:  Rhopressa and Rocklatan: Electronically approved for patients who have a trial of ONE generic agent, within ophthalmics- glaucoma within 60 days

#### **OPIOID DEPENDENCE TREATMENTS**

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naloxone SYRINGE, VIAL naltrexone TABLET NARCAN (naloxone) SPRAY	KLOXXADO (naloxone) <sup>NR</sup> <b>NASAL</b>	<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
i ioioiioa rigonio	11011 1 101011047 (901110	1 1101 / tatilo il Lation of Good Office Id

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. AL – Age Limit

CL – Prior Authorization / Class Criteria apply QL – Quantity/Duration Limit

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•		_	•	-	
acetic acid (generic for Vosol)	acetic acid/hydrocortisone (generic for Vosol HC)	•	Non-preferr approved fo failed a trial within this o	or patients of the pr	s who have referred agent

#### **OTIC ANTIBIOTICS**

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ambrisentan (generic Letairis) sildenafil TABLET (generic Revatio) <sup>CL</sup> tadalafil (generic for Adcirca) <sup>CL</sup> TRACLEER TABLET (bosentan) TYVASO INHALATION (treprostinil) VENTAVIS INHALATION (iloprost)	ADEMPAS (riociguat) <sup>CL</sup> ADCIRCA (tadalafil) <sup>CL</sup> bosentan <b>TABLET</b> (generic Tracleer) LETAIRIS (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) sildenafil <b>SUSPENSION</b> (generic Revatio) <sup>CL</sup> TRACLEER <b>TABLETS FOR</b> SUSPENSION (bosentan) 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>

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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## **PANCREATIC ENZYMES**

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

#### PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
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CHILD LITTLE ANIMALS VITAMINS
CHEW OTC (pedi multivit 91/iron
fum) <b>CHEW</b>

child multivitamins chew otc (pedi multivit 19/folic acid) CHEW

CHILDREN'S CHEW MULTIVIT-IRON OTC (pedi multivit 91/iron fum) **CHEW** 

children's chewables otc (pedi multivit 23/folic acid) CHEW

children's vitamins with iron otc (pedi multivit/iron)

fluoride/vitamins A,C,AND D (ped multivit A,C,D3, 21/fluoride) **DROPS** 

infant-toddler multivit drop OTC (pediatric multivit no. 165 drops)

no.164/ferrous sulfate drops)

infant-toddler tri-vit drop (vit a palmitate/vit c/vit d3 drops)

multivitamins with fluoride (pedi multivit 2/fluoride) DROPS

multivits with iron and fluoride (pedi multivit 45/fluoride/iron) DROPS

MVC-FLUORIDE (pedi multivit 12/fluoride) CHEW TAB

ped mvi A,C,D3,No 21/fluoride DROPS pedi mvi no. 16 with fluoride CHEW pedi mvi 17 with fluoride CHEW

POLY-VI-SOL OTC (pedi multivit 81) **DROPS** 

POLY-VI-SOL WITH IRON (pedi multivit 80/ferrous sulfate) DROPS

TRI-VI-SOL OTC (vit A palmitate/vit C/Vit D3) DROPS

tri-vite-fluoride 0.25 mg/ml, and 0.5 mg/ml

VITALETS OTC (pedi multivit 36/iron) **CHEW** 

DEKAs PLUS (ped multivitamin no.128/vitamin K)NR

ESCAVITE (pedi multivit 47/iron/fluoride)

ESCAVITE D (pedi multivit 78/iron/fluoride) CHEW

ESCAVITE LQ (pedi multivit 86/iron/fluoride)

FLORIVA (pedi multivit 85/fluoride) **CHEW** 

FLORIVA PLUS OTC and Rx (pedi multivit 130/fluoride) DROPS

multivit A, B, D, E, K, ZN (pediatric multivit 153/D3/K)

POLY-VI-FLOR (pedi multivit 33/fluoride) CHEW

POLY-VI-FLOR (pedi multivit 37/fluoride) DROPS

infant-toddler multivit-iron OTC (pedi mv POLY-VI-FLOR w/IRON (pedi multivit 33/fluoride/iron) CHEW

> POLY-VI-FLOR w/IRON (pedi multivit 37/fluoride/iron) DROPS

> QUFLORA OTC and Rx (pedi multivit 84/fluoride)

QUFLORA FE (pedi multivit 142/iron/fluoride)

TRI-VI-FLORO (ped multivit A, C, D3, 38/fluoride)

Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class

Drug specific criteria:

**DEKAs Plus**: Approved for diagnosis of Cystic Fibrosis

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#### **PENICILLINS**

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

#### **PHOSPHATE BINDERS**

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

#### **PLATELET AGGREGATION INHIBITORS**

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

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#### **PRENATAL VITAMINS**

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
c-nate dha SOFTGEL complete natal dha (pnv2/iron b-g suc-p/fa/omega-3) calcium-pnv 28-1-250mg SOFTGEL classic prenatal TABLET (prenatal vit/fe fum/fa) COMPLETENATE CHEWABLE CONCEPT DHA CAPSULE CONCEPT DHA CAPSULE elite-ob CAPLET (fe c/fa) MARNATAL-F CAPSULE PRENATA TAB CHEW pnv with ca, #72/iron/fa pnv-ob+dha combo pack (pnv22/iron cbn&gluc/fa/dss/dha) pnv-vp-u CAPSULE prenaissance CAPSULE (pnv80/iron fum/fa/dss/dha) prenaissance plus SOFTGEL (pnv69/iron/fa/dss/dha) prenatal vitamin TABLET (pnv#124/iron/fa) prenatal no.137/iron/fa OTC pretab 29mg-1 TABLET (pnv#78/iron/fa) PUREFE PLUS PUREFE OB PLUS TARON-PREX PRENATAL TRINATAL RX 1 triveen-duo dha combo pack		<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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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MAKENA <b>AUTO INJECTOR</b> (hydroxyprogesterone caproate)	hydroxyprogesterone caproate (generic Makena) MAKENA (hydroxyprogesterone caproate) <b>SDV</b>	<ul> <li>When filled as outpatient prescription, use limited to:         <ul> <li>Singleton pregnancy AND</li> <li>Previous Pre-term delivery AND</li> </ul> </li> <li>No more than 20 doses (administered between 16 -36 weeks gestation)</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) RX pantoprazole (generic Protonix) <sup>QL</sup> PROTONIX SUSP (pantoprazole)	DEXILANT (dexlansoprazole) esomeprazole magnesium (generic Nexium) esomeprazole strontium lansoprazole (generic Prevacid) <sup>QL</sup> NEXIUM SUSPENSION (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) pantoprazole GRANULES QL rabeprazole (generic Aciphex)	<ul> <li>Non-preferred agents will be approved for patients who have failed an 8-week trial of BOTH preferred agents within this drug class</li> <li>Pediatric Patients:         Patients ≤ 4 years of age − No PA required for Prevacid 30mg or omeprazole 20mg capsules (used to compound suspensions).     </li> <li>Drug-specific criteria:         <ul> <li>Prilosec®OTC/Omeprazole OTC EXCLUDED from coverage Acceptable as trial instead of Omeprazole 20mg</li> <li>Prevacid Solutab: may be approved after trial of compounde suspension.</li> <li>Patients ≥ 5 years if age- Only approve non-preferred for GI diagnosis if:</li></ul></li></ul>

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#### **SEDATIVE HYPNOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BENZODI.	AZEPINES	■ Lunesta®/ Rozerem®/zolpidem
temazepam 15mg, 30mg (generic for Restoril)		

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

#### **SINUS NODE INHIBITORS**

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

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#### **SKELETAL MUSCLE RELAXANTS**

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

PDL Updated October 1, 2021 Highlights indicated change from previous posting STEROIDS, TOPICAL

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

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### STEROIDS, TOPICAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HIGH POTENCY		High Potency Non-preferred
triamcinolone acetonide OINTMENT, CREAM triamcinolone LOTION	amcinonide CREAM, LOTION, OINTMENT  betamethasone dipropionate betamethasone / propylene glycol betamethasone valerate desoximetasone diflorasone diacetate fluocinonide SOLUTION fluocinonide CREAM, GEL, OINTMENT 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 OINTMENT (triamcinolone) VANOS (fluocinonide)	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class
VERY HIG	H POTENCY	Very High Potency Non-preferred
clobetasol emollient (generic for Temovate-E) clobetasol propionate CREAM, GEL, OINTMENT, SOLUTION halobetasol propionate (generic for Ultravate)	APEXICON-E (diflorasone) BRYHALI (halobetasol prop) LOTION clobetasol SHAMPOO, LOTION clobetasol propionate FOAM, SPRAY CLOBEX (clobetasol) halobetasol propionate FOAM (generic for Lexette) AL,QL IMPEKLO (clobetasol) LOTIONAL,NR 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

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## STIMULANTS AND RELATED AGENTS<sup>AL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CNS STIMULANTS		<ul> <li>Non-preferred agents will be approved for patients who have</li> </ul>
Ampheta	mine type	failed a trial of ONE preferred
ADDERALL XR (amphetamine salt combo) amphetamine salt combination IR VYVANSE (lisdexamfetamine) CAPSULE, CHEWABLE	ADZENYS XR (amphetamine) amphetamine ER (generic for Adzenys ER) SUSPENSION amphetamine salt combination ER	agent within this drug class  Drug-specific criteria:  Procentra®: May be approved with documentation of swallowing disorder  Zenzedi®: Requires clinical reason generic dextroamphetamine IR cannot be used

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## STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Methylph	enidate type	<ul> <li>Non-preferred agents will be approved for patients who have</li> </ul>
CONCERTA (methylphenidate ER) <sup>QL</sup> 18mg, 27mg, 36mg, 54mg dexmethylphenidate (generic for Focalin IR) FOCALIN XR (dexmethylphenidate) METHYLIN SOLUTION (methylphenidate) methylphenidate (generic for Ritalin) methylphenidate SOLUTION (generic for Methylin) methylphenidate ER (generic for Ritalin SR) QUILLICHEW ER CHEWTAB (methylphenidate)	ADHANSIA XR (methylphenidate) QL APTENSIO XR (methylphenidate) COTEMPLA XR-ODT	failed a trial of TWO preferred agents within this drug class  Maximum accumulated dose of 108mg per day for ages < 18  Maximum accumulated dose of 72mg per day for ages > 19  Drug-specific criteria:  Daytrana®: May be approved in history of substance use disorder by parent, caregiver, or patient.

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## STIMULANTS AND RELATED ADHD DRUGS (Continued)<sup>AL</sup>

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

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#### **TETRACYCLINES**

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

## THROMBOPOIESIS STIMULATING PROTEINSCL

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

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#### **THYROID HORMONES**

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

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#### **ULCERATIVE COLITIS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OR	AL	Non-preferred agents will be
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine) LIALDA (mesalamine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) GIAZO (balsalazide) mesalamine ER (generic Apriso) mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	approved for patients who have failed a trial of ONE preferred agent within this drug class  Drug-specific criteria:  Asacol HD®/Delzicol DR®/Pentasa®: Requires clinical reason why preferred mesalamine products cannot be used  Giazo®: Requires clinical reason why generic balsalazide cannot be used
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
ORIAHNN (elagolix/ estradiol/ norethindrone) <sup>AL,CL</sup> ORILISSA (elagolix sodium) <sup>QL,CL</sup>	MYFEMBREE (relugolix/ estradiol/ norethindrone acetate) <sup>AL, NR, QL</sup>	Orilissa/Oriahnn: Requires an FDA approved indication, must follow FDA dosing guidelines, and have had a trial and failure of an NSAID and oral contraceptive

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## **VASODILATORS, CORONARY**

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