

DEPT. OF HEALTH AND HUMAN SERVICES



Pete Ricketts, Governor

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

PDL Updated November 3, 2020 *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 <u>https://druglookup.fhsc.com/druglookupweb/?client=nestate</u>

• **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

- <u>Buprenorphine Products PA Form</u>
- <u>Buprenorphine Products Informed Consent</u>
- 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: https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf

PDL Updated November 3, 2020 Highlights indicated change from previous posting ACNE AGENTS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AZELEX (azelaic acid) benzoyl peroxide (BPO) GEL, WASH, LOTION OTC clindamycin/BPO (generic Duac) clindamycin phosphate SOLUTION DIFFERIN LOTION, CREAM, Rx-GEL (adapalene) DIFFERIN GEL (adapalene) OTC erythromycin SOLUTION PANOXYL 10% WASH (BPO) OTC tretinoin CREAM, GEL ^{AL} (generic Retin-A)	adapalene (generic differin) adapalene/BPO (generic Epiduo) <i>AKLIEF (trifarotene)^{AL}</i> ALTRENO (tretinoin) ^{AL} <i>AMZEEQ (minocycline)</i> <i>ARAZLO (tazarotene)^{AL}</i> ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) BENZACLIN PUMP (clindamycin/BPO) <i>BENZEFOAM (benzoyl peroxide)^{NR}</i> benzoyl peroxide CLEANSER , CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide GEL Rx <i>benzoyl peroxide</i> FOAM (generic Clindamycin FOAM , LOTION clindamycin FOAM , LOTION clindamycin/BPO (generic Acanya, Benzaclin) GEL <i>clindamycin/BPO</i> (generic Veltin, Ziana) dapsone (generic Aczone) EPIDUO FORTE GEL PUMP (adapalene/BPO) erythromycin GEL, PLEDGET erythromycin-BPO (generic for Benzamycin) EVOCLIN (clindamycin) FABIOR (tazarotene foam) NEUAC (clindamycin/BPO) OVACE PLUS (sulfacetamide sodium) PLIXDA (adapalene) SWAB <i>RETIN-A</i> GEL, CREAM ^{AL} (tretinoin) sulfacetamide sulfacetamide/sulfur SUMADAN (sulfacetamide/sulfur) tazarotene CREAM (generic Tazorac) TRETIN-X (tretinoin) 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

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

PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **ALZHEIMER'S AGENTS**

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

ANALGESICS, OPIOID LONG-ACTING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) ^{QL} PATCH fentanyl 25, 50, 75, 100 mcg PATCH ^{QL} morphine ER TABLET (generic MS Contin, Oramorph SR)	ARYMO ER (morphine sulfate) ^{QL} BELBUCA (buprenorphine) ^{CL} buccal buprenorphine PATCH (generic Butrans) ^{QL}	The Center for Disease Control (CDC) does not recommend long acting opioids when beginning opioid treatment.
OXYCONTIN ^{CL} (oxycodone ER)	EMBEDA (morphine sulfate/ naltrexone) DURAGESIC MATRIX (fentanyl) ^{QL} fentanyl 37.5, 62.5, 87.5 mcg PATCH ^{QL} hydrocodone bitartrate ER (generic for Zohydro ER) hydromorphone ER (generic for	 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
	Exalgo) ^{CL} HYSINGLA ER (hydrocodone ER) KADIAN (morphine ER) methadone ^{CL} MORPHABOND ER (morphine sulfate) morphine ER (generic for Avinza, Kadian) CAPSULE NUCYNTA ER (tapentadol) ^{CL} oxycodone ER (generic Oxycontin) oxymorphone ER (generic Opana ER) tramadol ER (generic Conzip, Ryzolt, Ultram ER) ^{CL}	 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 November 3, 2020 *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	APADAZ (benzhydrocodone/APAP) ^{CL} benzhydrocodone/APAP (generic Apadaz ^{,CL} 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) ^{CL} OXAYDO (oxycodone) ^{CL} oxycodone CAPSULE oxycodone (APAP SOL UTION	 Non-pretented agents win be approved for patients who have failed THREE preferred agents within this drug class within the las 12 months Note: for short acting opiate tablet: and capsules there is a maximum quantity limit of #150 per 30 days. Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will consist of -prescriptions limited to a 7 day supply, AND -initial opiate prescription fill limited to maximum of 50 Morphine Milligram Equivalents (MME) per day These limits may only be exceede with patient specific documentation of medical necessity, with examples such as, cancer diagnosis, end-of-life care, palliative care, Sickle Cell Anemia or prescriber attestation that patient is not recently opiate naive Drug-specific criteria: Apadaz: Approved only for diagnosis of acute pain, for 30 days or less Tramadol/APAP: Clinical reason why individual ingredients can't be used

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PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **ANALGESICS**, **OPIOID SHORT-ACTING**^{QL} (Continued)

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NASAL		
	butorphanol SPRAY ^{QL} LAZANDA (fentanyl citrate)	
BUCCAL/TRANSMUCOSAL ^{CL}		Drug-specific criteria: Abstral [®] /Actiq [®] /Fentora [®] /
	ABSTRAL (fentanyl) ^{CL} fentanyl TRANSMUCOSAL (generic Actiq) ^{CL} FENTORA (fentanyl) ^{CL}	Onsolis (fentanyl): Approved only for diagnosis of cancer AND current use of long-acting opiate
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ANDROGENIC AGENTS (Topical)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
testosterone PUMP (generic Androgel) ^{CL}	ANDRODERM (testosterone) ^{CL} NATESTO (testosterone) ^{CL} testosterone PACKET (generic Androgel) ^{CL} <i>testosterone GEL, PACKET, PUMP</i> <i>(generic Vogelxo)</i> testosterone (generic Axiron) testosterone (generic Fortesta) testosterone (generic Testim)	 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 In addition, non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 6 months Drug-specific criteria: Androderm®/Androgel®: Approved for Males only Natesto®: Approved for Males only with diagnosis of: Primary hypogonadism (congenital or acquired) OR Hypogonadotropic hypogonadism (congenital or acquired)

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PDL Updated November 3, 2020 Highlights indicated change from previous posting ANGIOTENSIN MODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ACE INHIBITORS		Non-preferred agents will be
benazepril (generic Lotensin) enalapril (generic Vasotec) fosinopril (generic Monopril) lisinopril (generic Prinivil, Zestril) quinapril (generic Accupril) ramipril (generic Altace)	captopril (generic Capoten) EPANED (enalapril) ^{CL} ORAL SOLUTION moexepril (generic Univasc) perindopril (generic Aceon) QBRELIS (lisinopril) ^{CL} ORAL SOLUTION trandolapril (generic Mavik)	 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
benazepril/HCTZ (generic Lotensin HCT) enalapril/HCTZ (generic Vaseretic) fosinopril/HCTZ (generic Monopril HCT) lisinopril/HCTZ (generic Prinzide, Zestoretic) quinapril/HCTZ (generic Accuretic)	captopril/HCTZ (generic Capozide) moexipril/HCTZ (generic Uniretic)	tablet is not appropriate
ANGIOTENSIN REG	CEPTOR BLOCKERS	_
irbesartan (generic Avapro) Iosartan (generic Cozaar) valsartan (generic Diovan)	candesartan (generic Atacand) EDARBI (azilsartan) eprosartan (generic Teveten) olmesartan (generic Benicar) telmisartan (generic Micardis)	

^{NR} – Product was not reviewed - New Drug criteria will apply Page **7** of **79**

PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **ANGIOTENSIN MODULATORS (Continued)**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANGIOTENSIN RECEPTOR BLOCKER/DIURETIC COMBINATIONS		Non-preferred agents will be
irbesartan/HCTZ (generic Avalide) Iosartan/HCTZ (generic Hyzaar) valsartan/HCTZ (generic Diovan-HCT)	candesartan/HCTZ (generic Atacand- HCT) EDARBYCLOR (azilsartan/ chlorthalidone) olmesartan/HCTZ (generic Benicar- HCT) telmisartan/HCTZ (generic Micardis- HCT)	 approved for patients who have failed TWO preferred agents within this drug class within the last 12 months Non-preferred combination products may be covered as individual prescriptions without prior authorization
	MODULATOR/	- Angiotensin Modulator/Calcium Channel Blocker Combinations:
	OCKER COMBINATIONS	_ Combination agents may be
amlodipine/benazepril (generic Lotrel) amlodipine/valsartan (generic Exforge)	amlodipine/olmesartan (generic Azor) amlodipine/olmesartan/HCTZ (generic Tribenzor) amlodipine/telmisartan (generic Twynsta) <i>amlodipine/valsartan/HCTZ (generic Exforge HCT)</i> PRESTALIA (perindopril/amlodipine) trandolapril/verapamil (generic Tarka)	approved if there has been a trial and failure of preferred agent
		 Direct Renin Inhibitors/Direct Renin Inhibitor Combinations:
DIRECT RENI	N INHIBITORS	- May be approved witha history of
	aliskiren (generic Tekturna) ^{QL}	TWO preferred ACE Inhibitors or Angiotensin Receptor Blockers
DIRECT RENIN INHIBITOR COMBINATIONS		within the last 12 months
	TEKTURNA/HCT (aliskiren/HCTZ)	
NEPRILYSIN INHIBI	TOR COMBINATION	
ENTRESTO (sacubitril/valsartan) ^{QL}		

ANGIOTENSIN RECEPTOR BLOCKER/BETA-BLOCKER COMBINATIONS

BYVALSON (nevibolol/valsartan)

ANTHELMINTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALBENZA (albendazole) BILTRICIDE (praziquantel) ivermectin (generic for Stromectol)	EMVERM (mebendazole) ^{CL} praziquantel (generic for Biltricide) STROMECTOL (ivermectin)	 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 Drug-specific criteria: Emverm: Approval will be considered for indications not covered by preferred agents

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 ^{NR,AL} (peanut allergen powder-dnfp)	 Class Criteria: Approved for immunotherapy for the treatment of grass pollen-induced allergic rhinitis with or without conjunctivitis. Patient has had treatment failure with or contraindication to: antihistamines AND montelukast Clinical reason as to why allergy shots cannot be used. Drug-specific criteria: ORALAIR Confirmed by positive skin test or in vitro testing for pollen- specific 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.

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ANTIBIOTICS, GASTROINTESTINAL

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting ANTIBIOTICS, INHALED

Preferred Agents ^{CL}	Non-Preferred Agents	Prior Authorization/Class Criteria
BETHKIS (tobramycin) ^{CL} KITABIS PAK (tobramycin) ^{CL} TOBI-PODHALER (tobramycin) ^{CL,QL}	ARIKAYCE (amikacin liposomal inh) ^{CL} SUSPENSION CAYSTON (aztreonam lysine) ^{QL,CL} <i>tobramycin (generic for Bethkis)^{NR}</i> tobramycin (generic Tobi) ^{CL}	 Diagnosis of Cystic Fibrosis is required for all agents ICD10 Group = E84, ICD9 = 277.00, 277.01, 277.02, 277.03, 277.09 Drug-specific criteria: 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 Cayston[®]: Trial of tobramycin via nebulizer and demonstration of TOBI[®] compliance required Tobi Podhaler[®]: Requires trial of tobramycin via nebulizer or documentation why nebulized tobramycin cannot be used

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 OINTMENT, CREAM mupirocin CREAM (generic Bactroban) ^{CL}	 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

PDL Updated November 3, 2020 Highlights indicated change from previous posting ANTIBIOTICS, VAGINAL

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

ANTICOAGULANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg ^{CL,QL} XARELTO DOSE PACK (rivaroxaban)	BEVYXXA (betrixaban) ^{QL} fondaparinux (generic Arixtra) FRAGMIN (dalteparin) SAVAYSA (edoxaban) ^{QL}	 Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 12 months Drug-specific criteria: Coumadin[®]: Clinical reason generic warfarin cannot be used Savaysa[®]: Approved diagnoses include: Stroke and systemic embolism (SE) risk reduction in nonvalvular atrial fibrillation (NVAF) OR Treatment of deep vein thrombosis (DVT) and pulmonary embolism (PE) following 5-10 days of parenteral anticoagulant therapy Xarelto 2.5mg: Use limited to reduction of risk of major cardiovascular events (cardiovascular death, myocardial infarction, and stroke) in patients with chronic coronary artery disease or peripheral artery disease

PDL Updated November 3, 2020 Highlights indicated change from previous posting **ANTIEMETICS/ANTIVERTIGO AGENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CANNABINOIDS		Non-preferred agents will be
dronabinol (generic Marinol) ^{AL}	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) ^{QL}	ANZEMET (dolasetron) granisetron (generic Kytril) SANCUSO (granisetron) ^{CL} ZUPLENZ (ondansetron)	 Drug-specific criteria: Akynzeo®/Emend®/Varubi®: Approved for Moderately/Highly emetogenic chemotherapy with dexamethasone and a
NK-1 RECEPTO	R ANTAGONIST	 5-HT3 antagonist WITHOUT trial of preferred agents
	aprepitant (generic Emend) ^{QL,CL} AKYNZEO (netupitant/palonosetron) ^{CL} VARUBI (rolapitant) TABLET ^{CL}	<u>Regimens include</u> : AC combination (Doxorubicin or Epirubicin with Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine,
TRADITIONAL ANTIEMETICS		Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine,
DICLEGIS (doxylamine/pyridoxine) ^{CL,QL} 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 SUPPOSITOR Y (generic Compazine) promethazine SUPPOSITOR Y 50mg scopolamine TRANSDERMAL trimethobenzamide TABLET (generic Tigan)	 Dacarbazine, Dactinomycin, Daunorubicin, Epirubicin, Etoposide, Hexamethylmelamine, Idarubicin, Ifosfamide, Imatinib, Interferon α, Irinotecan, Mechlorethamine, Melphalan, Methotrexate, Oxaliplatin, Procarbazine, Streptozotocin, Temozolomide Diclegis®/Bonjesta: Approved only for treatment of nausea and vomiting of pregnancy Metozolv ODT®: Documentation of inability to swallow or Clinical reason oral liquid cannot be used Sancuso®/Zuplenz®: Documentation of oral dosage form intolerance

PDL Updated November 3, 2020 Highlights indicated change from previous posting ANTIFUNGALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche) luconazole SUSPENSION, TABLET (generic Diflucan) griseofulvin microsized TABLET hystatin SUSPENSION, TABLET erbinafine (generic Lamisil)	CRESEMBA (isavuconazonium) ^{CL} flucytosine (generic Ancobon) ^{CL} griseofulvin ultramicrosize (generic GRIS-PEG) itraconazole (generic Nizoral) nystatin POWDER ONMEL (itraconazole) ORAVIG (miconazole) posaconazole (generic Noxafil) ^{AL,CL} TOLSURA (itraconazole) ^{CL} voriconazole (generic VFEND) ^{CL}	 Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class Drug-specific criteria: Cresemba®: Approved for diagnosis of invasive aspergillosis or invasive mucomycosis Flucytosine: Approved for diagnosis of: Candida: Septicemia, endocarditis, UTIs Cryptococcus: Meningitis, pulmonary infections 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 Noxafil® Suspension: Oropharyngeal/esophageal candidiasi refractory to itraconazole and/or fluconazole Onmel®: Requires trial and failure or contraindication to terbinafine Sporanox®/Itraconazole: Approved for diagnosis, Blastomycosis, Histoplasmosis, Onychomycosis due to terbinafine-resistant dematophytes, Oropharyngeal/ esophageal candidiasis refractory to fluconazole Sporanox®: Requires trial and failure of generic itraconazole Vfend®: No trial for diagnosis of Aspergillosis, Blastomycosis, and Histoplasmosis and requires a trial and failure of generic itraconazole 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 <i>Fusarium spp.</i>, Oropharyngeal/esophageal candidiasi refractory to fluconazole

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTIFUNGAL		Non-preferred agents will be
Clotrimazole CREAM (generic Lotrimin) RX, OTC clotrimazole SOLN OTC ketoconazole CREAM, SHAMPOO (generic Nizoral) LAMISIL (terbinafine) SPRAY OTC LAMISIL AT CREAM (terbinafine) OTC miconazole CREAM, POWDER OTC nystatin terbinafine OTC (generic Lamisil AT) tolnaftate POWDER, CREAM, POWDER OTC (generic Tinactin)	ALEVAZOL (clotrimazole) OTC ciclopirox CREAM, GEL, SUSPENSION (generic Ciclodan, Loprox) ciclopirox NAIL LACQUER (generic	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months Drug-specific criteria: Extina: Requires trial and failure or contraindication to other ketoconazole forms Jublia: Approved diagnoses includ Onychomycosis of the toenails due to <i>T.rubrum OR T. Mentagrophytes</i> nystatin/triamcinolone: Indivudual ingredients available without prior authorization ciclopirox nail lacquer: No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine
ANTIFUNGAL/STER	OID COMBINATIONS	

clotrimazole/betamethasone CREAM (generic Lotrisone)

clotrimazole/betamethasone LOTION (generic Lotrisone) nystatin/triamcinolone (generic Mycolog)

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AL – Age Limit

PDL Updated November 3, 2020 Highlights indicated change from previous posting ANTIHISTAMINES, MINIMALLY SEDATING

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

ANTIHYPERTENSIVES, SYMPATHOLYTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CATAPRES-TTS (clonidine) clonidine TABLET (generic for Catapres) guanfacine (generic for Tenex) methyldopa	clonidine TRANSDERMAL 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

ANTIHYPERURICEMICS

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

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

ANTIMIGRAINE AGENTS, OTHER

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

cannot be used

PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **ANTIMIGRAINE AGENTS**, **TRIPTANS**^{QL}

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

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 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

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

AL__ Age Limit

PDL Updated November 3, 2020 Highlights indicated change from previous posting ANTIPARKINSON'S AGENTS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOL benztropine (generic for Cogentin) trihexyphenidyl (generic for Artane)	INERGICS	 Non-preferred agents will be approved for patients who have failed ONE preferred agents within this drug class
COMT INF	HIBITORS	
DOPAMINE bromocriptine (generic for Parlodel) pramipexole (generic for Mirapex) ropinirole (generic for Requip)	entacapone (generic for Comtan) <i>ONGENTYS (Opicapone)</i> ^{NR,QL} tolcapone (generic for Tasmar) AGONISTS ropinirole ER <i>(generic for Requip ER)</i> ^{CL} NEUPRO (rotigotine) ^{CL} pramipexole ER (generic for Mirapex ER) ^{CL}	 Drug-specific criteria: Carbidopa/Levodopa ODT: Approved for documented swallowing disorder COMT Inhibitors: Approved if using as add-on therapy with levodopa-containing drug Gocovri: Required diagnosis of Parkinson's disease and had trial of or is intolerant to amantadine AND must be used as an add-on therapy with levodopa-containing drug
MAO-B INI	HIBITORS	Inbrija: Approval upon diagnosis of
selegiline CAPSULE, TABLET (generic for Eldepryl)	rasagiline (generic for Azilect) ^{QL} XADAGO (safinamide) ZELAPAR (selegiline) ^{CL}	 Parkinson's disease and concurrent treatment with carbidopa/levodopa agent Neupro[®]: For Parkinsons: Clinical reason
OTHER ANTIPAR	KINSON'S DRUGS	required why preferred agent
amantadine CAPSULE, SYRUP TABLET (generic for Symmetrel) carbidopa/levodopa (generic for Sinemet) carbidopa/levodopa ER (generic for Sinemet CR) levodopa/carbidopa/entacapone (generic for Stalevo)	carbidopa (generic for Lodosyn) carbidopa/levodopa ODT (generic for Parcopa) DUOPA (carbidopa/levodopa) GOCOVRI (amantadine) ^{QL} INBRIJA (levodopa) INHALER ^{CL,QL} <i>KYNMOBI (apomorphine)^{QL/NR}</i> <i>NOURIANZ (istradefylline)^{NR,CL,QL}</i> OSMOLEX ER (amantadine) ^{QL} RYTARY (carbidopa/levodopa) STALEVO (levodopa/carbidopa/entacapone)	

ANTIPSORIATICS, ORAL

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

ANTIPSORIATICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcipotriene CREAM, OINTMENT, SOLUTION,	calcitriol (generic for Vectical) calcipotriene/betamethasone (generic for Taclonex) calcipotriene/betamethasone SUSP (generic for Taclonex Scalp) ^{NR} CALCITRENE (calcipotriene) DOVONEX CREAM (calcipotriene) DUOBRII (halobetasol prop./tazarotene ENSTILAR (calcipotriene/betamethasone) SORILUX (calcipotriene) TACLONEX SCALP (calcipotriene/betamethasone)	 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
ANTI-HERPE	ANTI-HERPETIC DRUGS	
acyclovir (generic Zovirax) famciclovir (generic Famvir) valacyclovir (generic Valtrex)	acyclovir SUSPENSION (generic for Zovirax) SITAVIG (acyclovir buccal) ^{CL}	approved for patients who have failed a 10-day trial of ONE preferred agent within the same group
ANTI-INFLUE	ANTI-INFLUENZA DRUGS	
oseltamivir (generic Tamiflu) ^{qL}	rimantadine (generic Flumadine) RELENZA (zanamivir) ^{QL} TAMIFLU (oseltamivir) ^{QL} XOFLUZA (baloxavir marboxil) ^{AL,CL,QL}	 Drug-specific criteria: Sitavig[®]: Approved for recurrent herpes labialis (cold sores) in immunocompetent adults Xofluza: Requires clinical, patient specific reason that a preferred agent cannot be used

ANTIVIRALS, TOPICAL

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

ANXIOLYTICS

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

 Alprazolam Intensol[®]: Requires trial of diazepam solution OR lorazepam Intensol[®]

PDL Updated November 3, 2020 Highlights indicated change from previous posting **BETA BLOCKERS, ORAL**

BILE SALTS

sotalol (generic Betapace)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ursodiol CAPSULE 300mg (generic for Actigall) ursodiol 250mg TABLET (generic for URSO)	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
ursodiol 500mg TABLET (generic for URSO FORTE)		

SOTYLIZE (sotalol)

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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) flavoxate MYRBETRIQ (mirabegron) OXYTROL (oxybutynin) tolterodine IR, ER (generic Detrol/ Detrol LA) trospium IR, ER (generic Sanctura/ Sanctura XR) VESICARE (solifenacin)	 Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class Drug-specific criteria: Myrbetrig[®]: Covered without trial in contraindication to anticholinergic agents

PDL Updated November 3, 2020 Highlights indicated change from previous posting BONE RESORPTION SUPRESSION AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BISPHOSF	PHONATES	Non-preferred agents will be
alendronate (generic Fosamax) TABLET ibandronate (generic Boniva) ^{QL}	alendronate SOLUTION (generic Fosamax) ^{QL} ATELVIA DR (risedronate) BINOSTO (alendronate) etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL}	 approved for patients who have failed a trial of ONE preferred agent within the same group Drug-specific criteria: Actonel[®] Combinations: Covered as individual agents without prior authorization Atelvia DR[®]: Requires clinical reason alendronate cannot be taken on an empty stomach
OTHER BONE RESORPTION SUP	PRESSION AND RELATED DRUGS	Binosto [®] : Requires clinical reason why alendronate tablets OR Fosamax [®] solution
calcitonin-salmon NASAL raloxifene (generic Evista)	EVISTA (raloxifene) FORTEO (teriparatide) ^{QL} Teriparatide ^{QL} TYMLOS (abaloparatide)	 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 at 7.5 dose of prednisolone equivalent Rheumatoid Arthritis Postmenopausal women with BMD T-score ≤ -2.5 at any site 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

PDL Updated November 3, 2020 Highlights indicated change from previous posting **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-REDUCTASE (5AR) INHIBITORS		• Alfuzosin/dutasteride/finasteride
dutasteride (generic for Avodart) finasteride (generic for Proscar)	dutasteride/tamsulosin (generic for Jalyn)	 Covered for males only Cardura XL[®]: Requires clinical reason generic IR form cannot be used Flomax[®]: Females covered for a 7 day supply with diagnosis of acute kidney stones Jalyn[®]: Requires clinical reason why individual agents cannot be used

BRONCHODILATORS, BETA AGONIST

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INHALERS – Short Acting		Non-preferred agents will be
PROAIR HFA (albuterol) PROVENTIL HFA (albuterol)	albuterol HFA (generic for ProAir HFA, Proventil HFA, Ventolin HFA) levalbuterol HFA (generic for Xopenex HFA) <i>PROAIR DIGIHALER (albuterol)</i> ^{NR} PROAIR RESPICLICK (albuterol)	 approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Ventolin HFA[®]: Requires trial and failure on Proventil HFA® AND Proair HFA® OR allergy/ contraindication/side effect to
INHALERS -	- Long Acting	BOTH
SEREVENT (salmeterol)	ARCAPTA NEOHALER (indacaterol) STRIVERDI RESPIMAT (olodaterol)	Xopenex [®] : Covered for cardiac diagnoses or side effect of tachycardia with albuterol product
INHALATIO	N 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)	BROVANA (arformoterol) levalbuterol (generic for Xopenex) PERFOROMIST (formoterol)	
ORAL		
albuterol SYRUP	albuterol TABLET albuterol ER (generic for Vospire ER) metaproterenol (formerly generic for Alupent) terbutaline (generic for Brethine)	

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting CALCIUM CHANNEL BLOCKERS, ORAL

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

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	6 – First Generation	_
cefadroxil CAPSULE, SUSPENSION (generic Duricef) cephalexin CAPSULE, SUSPENSION (generic Keflex)	cefadroxil TABLET (generic Duricef) cephalexin TABLET 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) VIAL	GRANIX (tbo-filgrastim) NEUPOGEN DISP SYR (filgrastim) NIVESTYM SYR,VIAL (filgrastim-aafi) ZARXIO (filgrastim-sndz) ZIEXTENZO SYR (pegfilgrastim- bmez) ^{NR}	 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	hailey fe 1/20 (norethindrone acetate/ ethinyl estradiol-iron) ^{NR} charlotte 24 fe (norethindrone acetate/ethinyl estradiol-iron) ^{NR}	
Specific agents can be looked up using the Drug Look-up Tool at: <u>https://druglookup.fhsc.com/drug</u> lookupweb/?client=nestate		

PDL Updated November 3, 2020 Highlights indicated change from previous posting COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
č	ANORO ELLIPTA (umeclidinium/vilanterol) DUAKLIR PRESSAIR (aclidinium br and formoterol fum) ^{NR} 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
albuterol/ipratropium (generic for Duoneb) ipratropium SOLUTION (generic for Atrovent)	N SOLUTION LONHALA (glycopyrrolate inhalation soln) YUPELRI (revefenacin) AGENT	Requires trial of a bronchodilator Requires documentation of one exacerbation in last year upon initial review
	AGENT DALIRESP (roflumilast) ^{CL, QL}	-

COUGH AND COLD, OPIATE COMBINATION

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting **CYSTIC FIBROSIS, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	KALYDECO PACKET, TABLET (ivacaftor) ^{QL, AL} ORKAMBI (lumacaftor/ivacaftor) PACKET, TABLET ^{QL, AL} SYMDEKO (tezacaftor/ivacaftor) ^{QL, AL} TRIKAFTA (elexacaftor, tezacaftor, ivacaftor) ^{AL, CL}	 Drug-specific criteria: Kalydeco®: Diagnosis of CF and documentation of the drug-specific, FDA-approved mutation of CFTR gene Orkambi®: Diagnosis of CF and documentation of presence of two copies of the F580del mutation (homozygous) of CFTR gene Symdeko: Diagnosis of CF and documentation of the drug specific, FDA approved mutation of CFTR gene. Trikafta: Diagnosis of CF and documentation of at least one F508del mutation in the CFTR gene

CYTOKINE & CAM ANTAGONISTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ENBREL (etanercept) KIT, MINI CART, PEN ^{QL} HUMIRA (adalimumab) ^{QL} OTEZLA (apremilast) ORAL^{CL,QL}	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIMZIA (certolizumab pegol) ^{QL} COSENTYX (secukinumab) ^{GL} <i>ENBREL (entanercept) VIAL^{NR,QL}</i> ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYRINGE KINERET (anakinra) OLUMIANT (baricitinib) ORAL^{CL,QL} ORENCIA (abatacept) SUB-Q RINVOQ ER (upadacitinib, ^{CL,QL} SILIQ (brodalumab) SIMPONI (golimumab) SKYRIZI (risankizamab-rzaa) STELARA (ustekinumab) SUB-Q TALTZ (ixekizumab) ^{AL} TREMFYA (guselkumab) ^{QL} XELJANZ (tofacitinib) ORAL^{CL,QL}	 Preferred agents will be approved with FDA-approved indication – ICD-10 diagnosis code is required. 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. Drug-specific criteria: Otezla: Requires a trial of Humira Olumiant: Requires documentation of inadequate response or intolerance to methotrexate and an inadequate response to one or more TNF antagonist therapies. Rinvoq, Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to methotrexate

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PDL Updated November 3, 2020 Highlights indicated change from previous posting DIURETICS

Preferred Agents	Non-Preferred Agents		Prior Authorization/Class Criteria
SINGLE-AGEN	IT PRODUCTS	•	Non-preferred agents will be
amiloride TABLET bumetanide TABLET chlorothiazide TABLET 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) methyclothiazide TABLET triamterene (generic Dyrenium)	_	approved for patients who have failed a trial of TWO preferred agents within this drug class
COMBINATIO	N PRODUCTS		
amiloride/HCTZ TABLET spironolactone/HCTZ TABLET (generic Aldactazide) triamterene/HCTZ CAPSULE, TABLET (generic Dyazide, Maxzide)			

ENZYME REPLACEMENT, GAUCHERS DISEASE

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

EPINEPHRINE, SELF-INJECTEDQL

	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
İ	binephrine (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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^{NR} – Product was not reviewed - New Drug criteria will apply Page **31** of **79**

ERYTHROPOIESIS STIMULATING PROTEINS

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

FLUOROQUINOLONES, ORAL

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

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GI MOTILITY, CHRONIC

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

GLUCAGON AGENTS^{QL}

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) ^{AL} NASAL GLUCAGON EMERGENCY (glucagon) INJ KIT (Lilly) glucagon INJECTION PROGLYCEM (diazoxide) SUSP	diazoxide SUSP (generic Proglycem) GLUCAGON EMERGENCY (glucagon) INJ KIT (Fresenius) GVOKE (glucagon) ^{AL} PEN , SYRINGE	 Non-preferred agents will be approved for patients who have failed a trial of 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

GLUCOCORTICOIDS, INHALED

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

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GLUCOCORTICOIDS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
budesonide EC CAPSULE (generic for	ALKINDI (hydrocortisone) GRANULES ^{AL/NR} CORTEF (hydrocortisone) cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSPENSION, TABLET ^{CL} ENTOCORT EC (budesonide) <i>HEMADY (dexamethasone)^{NR}</i> methylprednisolone 8mg, 16mg <i>ORTIKOS ER (budesonide)^{AL,NR,QL}</i> PEDIAPRED (prednisolone sodium phosphate) prednisolone sodium phosphate (generic for Millipred/Veripred) prednisolone sodium phosphate ODT prednisolone SOLUTION	 Prior Authorization/Class Criteria 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 Drug-specific criteria: Emflaza: Indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients 2 years of age and older Intensol Products: Patient specific documentation of why the less concentrated solution is not appropriate for the patient
	prednisone INTENSOL RAYOS DR (prednisone) TABLET	

GROWTH HORMONES

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

PDL Opdated November 3, 2020 Highlights indicated change from previo

H. PYLORI TREATMENTS

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

HAE TREATMENTSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BERINERT (C1 esterase inhibitor, human) INTRAVENOUS FIRAZYR (icatibant acetate) ^{AL} SUB-Q HAEGARDA (C1 esterase inhibitor, human) ^{AL} SUB-Q	CINRYZE (C1 esterase inhibitor, human)AL INTRAVENOUS icatibant acetate (generic for FIRAZYR) ^{AL} SUB-Q KALBITOR (ecallantide) ^{AL} SUB-Q RUCONEST (recombinant human C1 inhibitor) ^{AL} INTRAVENOUS TAKHZYRO (lanadelumab-flyo) ^{AL} SUB-Q	 HAE Treatments PA Form 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, and estrogencontaining products is contraindicated All prophylaxis agents (Haegarda, Takhzyro and Ciryze) require a history of two or more HAE attacks monthly, and trial and failure or contraindication to oral danazol Non-preferred agents will be approved for patients who have a failed trial or a contraindication to ONE preferred agent within this drug class

HEMOPHILIA TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FACTOR VIII		 Non-preferred agents will be
ADVATE ALPHANATE HELIXATE FS HUMATE-P KOATE-DVI VIAL KOVALTRY NOVOEIGHT NUWIQ RECOMBINATE XYNTHA KIT, SOLOFUSE	ADYNOVATE AFSTYLA ELOCTATE <i>ESPEROCT^{NR,}</i> HEMOFIL-M JIVI ^{AL} KOATE-DVI KIT KOGENATE FS OBIZUR	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class
FACT		
BENEFIX MONONINE PROFILNINE SD	ALPHANINE SD ALPROLIX IDELVION IXINITY REBINYN RIXUBIS	
FACTOR VIIa AND PROTHROME	SIN COMPLEX-PLASMA DERIVED	
NOVOSEVEN RT	FEIBA NF	
	XIII PRODUCTS	
CORIFACT	COAGADEX ^{CL} TRETTEN ^{CL}	
	AND PRODUCTS	
VONVENDI WILATE		
BISPECIFIC	CFACTORS	
	HEMLIBRA ^{CL}	

HEPATITIS B TREATMENTS

	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
entecavir 1 Iamivudine	TABLET hbv TABLET	adefovir dipivoxil BARACLUDE (entecavir) SOLUTION, TABLET EPIVIR HBV (lamivudine) TABLET , SOLUTION HEPSERA (adefovir dipivoxil) VEMLIDY (tenofovir alafenamide fumarate)	 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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PDL Updated November 3, 2020 Highlights indicated change from previous posting **HEPATITIS C TREATMENTS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIRECT ACTIN MAVYRET (glecaprevir/pibrentasvir) ^{CL} VOSEVI (sofosbuvir/velpatasvir/	IG ANTI-VIRAL DAKLINZA (daclatasvir) ^{CL} HARVONI 200/45MG, TABLET,	 <u>Hepatitis C Treatments PA Form</u> <u>Hepatitis C Criteria</u> Non-preferred products require trial of preferred agents within the
voxilaprev) ^{CL}	 HAR VONI 200/4300G, TABLET, (sofosbuvir/ledipasvir)^{CL} HAR VONI (ledipasvir/sofosbuvir)^{CL,NR} PELLET sofosbuvir/ledipasvir (generic Harvoni)^{CL} sofosbuvir/velpatasvir (generic Epclusa)^{CL} SOVALDI (sofosbuvir)^{CL,NR} PELLET SOVALDI TABLET (sofosbuvir)^{CL} VIEKIRA PAK (ombitasvir/ paritaprevir/ritonavir/dasabuvir)^{CL} ZEPATIER (elbasvir/grazoprevir)^{CL} 	 trial of preferred agents within the same group and 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 Mavyret not required in the following: Epclusa: For genotype 1-6 with decompensated cirrhosis along with
RIBA	VIRIN	ribavirin ■ Harvoni:
	REBETOL (ribavirin)	 For genotype 1 with decompensated cirrhosis along with ribavirin
PEGASYS (pegylated interferon alfa-	FERON	 Post liver transplant for genotype 1 or 4
2a) ^{CL} PEG-INTRON (pegylated interferon alfa-2b) ^{CL}		 For pediatric patients ages 3 to 11 years old with FDA indications
		 Sovaldi: For pediatric patients ages 3 to 11 years old with genotype 2 or 3 chronic HCV infection without cirrhosis or with compensated cirrhosis in combination with ribavirin Vosevi: Requires documentation of non-response after previous treatment course of Direct Acting Anti-viral agent (DAA) for genotype 1-6 without cirrhosis or with compensated cirrhosis

PDL Updated November 3, 2020 Highlights indicated change from previous posting **HISTAMINE II RECEPTOR BLOCKERS**

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

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PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **HIV / AIDS^{CL}**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CCR5 ANT	AGONISTS	 Non-preferred agents will be
SELZENTRY SOLN, TAB (maraviroc)		 approved for patients who have a diagnosis of HIV/AIDS and patie
EUSION IN	NHIBITORS	 specific documentation of why th
FUZEON SUB-Q (enfuvirtide) ^{QL}		 preferred products within this dru class are not appropriate for
``````````````````````````````````````		patient, including, but not limited
		to, drug resistance or concomita conditions not recommended wit
SENTRESS (raltegravir) ^{QL}	TIVICAY PD (dolutegravir) ^{NR}	preferred agents
SENTRESS HD (raltegravir)		<ul> <li>Patients undergoing treatment a the time of any preferred status</li> </ul>
TIVICAY (dolutegravir)		change will be allowed to continu
NON-NUCLEOSIDE REVERSE TRA	NSCRIPTASE INHIBITORS (NNRTIS)	therapy
EDURANT (rilpivirine)	efavirenz (generic Sustiva)	<ul> <li>Diagnosis of HIV/AIDS required</li> <li>OR</li> </ul>
NTELENCE (etravirine) ^{QL}	nevirapine IR, ER (generic	<ul> <li>Pre and Post Exposure</li> </ul>
PIFELTRO (doravirine) ^{QL}	Viramune/Viramune XR)	Prophylaxis
SUSTIVA CAPSULE, TABLET	RESCRIPTOR (delavirdine)	
(efavirenz)	VIRAMUNE (nevirapine) <b>SUSP</b>	
	SCRIPTASE INHIBITORS (NRTIS)	-
	. ,	_
abacavir <b>SOLN, TABLET</b> (generic Ziagen)	didanosine DR (generic Videx EC)	
EMTRIVA CAPSULE, SOLN	emtricitabine <b>CAPSULE</b> (generic for Emtriva) ^{NR}	
(emtricitabine)	EPIVIR (lamivudine)	
amivudine SOLN, TABLET (generic	RETROVIR (zidovudine)	
Epivir)	stavudine CAPSULE (generic Zerit)	
idovudine CAPSULE, SYRUP,	VIDEX (didanosine) <b>SOLN</b>	
<b>TABLET</b> (generic Retrovir)	ZIAGEN (abacavir)	
NUCLEOTIDE REVERSE TRAN	SCRIPTASE INHIBITORS (NRTIS)	
enofovir TABLET (generic Viread)		_
		_
	ETIC ENHANCER	
FRANINACURIN		

TYBOST (cobicistat)QL

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### HIV / AIDS^{CL} (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROTEASE	INHIBITORS	
atazanavir <b>CAPSULE</b> (generic Reyataz) LEXIVA <b>SUSP, TABLET</b> (fosamprenavir) NORVIR (ritonavir) <b>TAB</b> PREZISTA (darunavir) <b>SUSP, TABLET</b>	(tipranavir) CRIXIVAN (indinavir) fosamprenavir <b>TAB</b> (generic Lexiva)	
	VIRACEPT (nelfinavir)	

# PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **HIV / AIDS^{CL} (Continued)**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	E INHIBITORS (PIs) or PIs plus NETIC ENHANCER	
EVOTAZ (atazanavir/cobicistat) ^{QL} KALETRA <b>TAB</b> (lopinavir/ritonavir) PREZCOBIX (darunavir/cobicistat) ^{QL} lopinavir/ritonavir <b>SOLN</b> (generic Kaletra)	KALETRA <b>SOLN</b> (lopinavir/ritonavir)	
COMBINATION NUCLEOS(T)IDE RE	EVERSE TRANSCRIPTASE INHIBITORS	
abacavir/lamivudine (generic Epzicom) abacavir/lamivudine/zidovudine (generic Trizivir) CIMDUO (lamivudine/tenofovir) ^{QL} DESCOVY (emtricitabine/tenofovir) ^{QL} lamivudine/zidovudine (generic Combivir) TRUVADA (emtricitabine/tenofovir)	COMBIVIR (lamivudine/zidovudine) <i>emtricitabine/tenofovir (generic</i> <i>Truvada)</i> ^{CL,NR} EPZICOM (abacavir sulfate/lamivudine) TEMIXYS (lamivudine/tenofovir) ^{QL} TRIZIVIR (abacavir/lamivudine/zidovudine)	
COMBINATION PRODU	CTS – MULTIPLE CLASSES	
BIKTARVY (bictegravir/emtricitabine/ tenofovir) ^{QL} COMPLERA (rilpivirine/emtricitabine/tenofovir)	DOVATO (dolutegravir/lamivudine) ^{QL} efavirenz/emtricitabine/tenofovir (generic Atripla) ^{CL,NR} efavirenz/lamivudine/tenofovir (generic for Symfi) ^{NR,QL} efavirenz/lamivudine/tenofovir (generic for Symfi Lo) ^{NR,QL} JULUCA (dolutegravir/rilpivirine) ^{QL} SYMTUZA (darunavir/cobicistat/ emtricitabine/tenofovir) ^{QL}	

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### PDL Updated November 3, 2020 Highlights indicated change from previous posting HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acarbose (generic for Precose) Glyset (miglitol)	miglitol (generic for Glyset)	<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>

### HYPOGLYCEMICS. INCRETIN MIMETICS/ENHANCERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GLUCAGON-LIKE PEPTIDE-1 RE	CEPTOR AGONIST (GLP-1 RA) ^{CL}	Preferred agents require metformin
BYDUREON (exenatide ER) subcutaneous BYDUREON <b>PEN</b> (exenatide ER) subcutaneous BYETTA (exenatide) subcutaneous VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) BYDUREON BCISE <b>PEN</b> (exenatide) ^{QL} OZEMPIC (semaglutide) RYBELSUS (semaglutide) TANZEUM (albiglutide) TRULICITY (dulaglutide) A COMBINATIONS SOLIQUA (insulin glargine/lixisenatide) XULTOPHY (insulin	<ul> <li>trial and diagnosis of diabetes</li> <li>Non-preferred agents will be approved for patients who have:</li> <li>Failed a trial of TWO preferred agents within GLP-1 RA</li> <li>AND</li> <li>Diagnosis of diabetes with HbA1C ≥ 7 AND</li> <li>Trial of metformin, or contraindication or intolerance to metformin</li> </ul>
	degludec/liraglutide)	
AMYLIN	SYMLIN (pramlintide) subcutaneous	<ul> <li>ALL criteria must be met</li> <li>Concurrent use of short-acting mealtime insulin</li> <li>Current therapy compliance</li> <li>No diagnosis of gastroparesis</li> <li>HbA1C ≤ 9% within last 90 days</li> <li>Fingerstick monitoring of glucose during initiation of therapy</li> </ul>
DIPEPTIDYL PEPTIDAS	E-4 (DPP-4) INHIBITOR ^{QL}	
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) ^{AL}	Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within DPP-4

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NR – Product was not reviewed - New Drug criteria will apply Page 44 of 79

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### HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
JMALOG (insulin lispro) U-100 CARTRIDGE, PEN, VIAL JMALOG JR. (insulin lispro) U-100 PEN JMALOG MIX VIAL (insulin lispro/lispro protamine) JMALOG MIX PEN (insulin lispro/lispro protamine) JMULIN (insulin) VIAL JMULIN 70/30 VIAL JMULIN 70/30 VIAL JMULIN U-500 VIAL JMULIN R U-500 KWIKPEN ^{CL} JMULIN OTC PEN JMULIN 70/30 OTC PEN ANTUS SOLOSTAR PEN (insulin glargine) ANTUS (insulin glargine) VIAL EVEMIR (insulin detemir) PEN, VIAL DVOLOG (insulin aspart) CARTRIDGE, PEN, VIAL (insulin aspart/aspart protamine)	ADMELOG (insulin lispro) <b>PEN, VIAL</b> AFREZZA (regular insulin) <b>INHALATION</b> APIDRA (insulin glulisine) BASAGLAR (insulin glargine, rec) <b>PEN</b> FIASP (insulin aspart) <b>CARTRIDGE,</b> <b>PEN, VIAL</b> HUMALOG (insulin lispro) U-200 <b>PEN</b> insulin lispro (generic for Humalog) <b>PEN, VIAL</b> insulin aspart (generic for Novolog) <i>LYUMJEV KWIKPEN, VIAL(insulin</i> <i>lispro-aabc)</i> ^{NR} NOVOLIN (insulin) NOVOLIN (insulin) NOVOLIN 70/30 <b>VIAL</b> (insulin) TOUJEO SOLOSTAR (insulin glargine) <i>SEMGLEE (insulin glargine)</i> ^{NR} <b>PEN,</b> <i>VIAL</i> 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: Approved for physical reasons – such as dexterity problems and vision impairment</li> <li>Usage must be for self- administration, not only convenience</li> <li>Patient requires &gt;200 units/day</li> <li>Safety reason patient can't use vial/syringe</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)	<ul> <li>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</li> </ul>

### 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) ^{AL}	<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>

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CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

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

### HYPOGLYCEMICS, SGLT2

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) ^{QL,CL} INVOKAMET (canagliflozin/metformin) ^{QL, CL} INVOKANA (canagliflozin) ^{CL} JARDIANCE (empagliflozin) ^{QL, CL} XIGDUO XR (dapagliflozin/metformin) ^{QL,CL}	INVOKAMET XR (canagliflozin/metformin) ^{QL} SEGLUROMET (ertugliflozin/metformin) ^{QL} STEGLATRO (ertugliflozin) ^{QL} SYNJARDY (empagliflozin/metformin) SYNJARDY XR (empagliflozin/ metformin) ^{QL}	<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>
SULFONYLUREA	COMBINATIONS	
glipizide/metformin glyburide/metformin (generic Glucovance)		

### HYPOGLYCEMICS, TZD

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
THIZAOLIDIN	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)	<ul> <li>Combination products: Require clinical reason why individual ingredients cannot be used</li> </ul>

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### **IDIOPATHIC PULMONARY FIBROSIS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OFEV (nintedanib esylate) ^{CL}	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, ATOPIC DERMATITISAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIDEL (pimecrolimus)	DUPIXENT (dupilumab) ^{AL,CL} DUPIXENT <b>PEN^{AL,NR}</b> EUCRISA (crisaborole) pimecrolimus (generic for Elidel) tacrolimus (generic for Protopic) ^{CL}	<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: For 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> </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)	<ul> <li>Non-preferred agents require clinical reason why preferred agent within this drug class cannot be used</li> </ul>

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AL – Age Limit

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

### **IMMUNOSUPPRESSIVES, ORAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathiaprine (generic Imuran) cyclosporine, modified <b>CAPSULE</b> (generic Neoral) mycophenolate <b>CAPSULE, TABLET</b> (generic Cellcept) RAPAMUNE (sirolimus) <b>SOLUTION</b> tacrolimus	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 RAPAMUNE (sirolimus) TABLET SANDIMMUNE (cyclosporine) CAPSULE, SOLUTION sirolimus SOLUTION, TABLET (generic Rapamune) everolimus (generic for Zortress) ^{AL}	<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>Patients established on existing therapy will be allowed to continue</li> </ul>

### **INTRANASAL RHINITIS DRUGS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOL	INERGICS	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	<b>FAMINES</b>	drug class
azelastine 0.1% (generic for Astelin)	azelastine 0.15% (generic for Astepro) azelastine/fluticasone (generic for Dymista) olopatadine (generic for Patanase)	<ul> <li>Drug-specific criteria:</li> <li>mometasone: Prior authorization NOT required for children ≤ 12 years</li> <li>budesonide: Approved for use in Pregnancy (Pregnancy Category</li> </ul>
CORTICOS	STEROIDS	- B)
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)	<ul> <li>Veramyst®: Prior authorization NOT required for children ≤ 12 years</li> <li>Xhance: Indicated for treatment of nasal polyps in ≥ 18 years only</li> </ul>

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^{NR} – Product was not reviewed - New Drug criteria will apply Page **48** of **79** 

### PDL Updated November 3, 2020 Highlights indicated change from previous posting

### **LEUKOTRIENE MODIFIERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
montelukast <b>TABLET/CHEWABLE</b> (generic for Singulair) ^{AL}	montelukast <b>GRANULES</b> (generic for Singulair) ^{CL, AL} 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: PA not required for age &lt; 2 years</li> </ul>

### LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clindamycin <b>CAPSULE</b> clindamycin palmitate <b>SOLUTION</b> linezolid <b>TABLET</b>	CLEOCIN (clindamycin ) CAPSULE CLEOCIN PALMITATE (clindamycin) linezolid SUSPENSION SIVEXTRO (tedizolid phosphate) ZYVOX (linezolid) SUSPENSION, TABLET	<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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PDL Updated November 3, 2020 Highlights indicated change from previous posting LIPOTROPICS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BILE ACID SEQUESTRANTS		<ul> <li>Non-preferred agents will be</li> </ul>
cholestyramine (generic Questran) colestipol <b>TABLETS</b> (generic Colestid)	colesevelam (generic Welchol) <b>TABLET, PACKET</b> colestipol <b>GRANULES</b> (generic Colestid) QUESTRAN LIGHT (cholestyramine)	<ul> <li>approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Colesevelam: Trial not required for diabetes control and monotherapy with</li> </ul>
TREATMENT OF HOMOZYGOUS FA	MILIAL HYPERCHOLESTEROLEMIA	metformin, sulfonylurea, or insulin has been inadequate
	JUXTAPID (lomitapide) ^{CL} KYNAMRO (mipomersen) ^{CL}	<ul> <li>Juxtapid[®]/ Kynamro[®]:</li> <li>Approved for diagnosis of homozygous</li> </ul>
FIBRIC ACID	DERIVATIVES	familial hypercholesterolemia (HoFH)
fenofibrate (generic Tricor) gemfibrozil (generic Lopid)	fenofibrate (generic Antara/Fenoglide/ Lipofen/Lofibra/Triglide) fenofibric acid (generic Fibricor/Trilipix)	<ul> <li>OR</li> <li>Treatment failure/maximized dosing/contraindication to ALL the following: statins, ezetimibe, niacin, fibric acid derivatives, omega-3 agents,</li> </ul>
NIA	CIN	bile acid sequestrants
niacin ER (generic for Niaspan)	NIACOR (niacin IR) NIASPAN (niacin ER)	• Require faxed copy of REMS PA form
OMEGA-3 F	ATTY ACIDS	<ul> <li>Lovaza[®]: Approved for TG ≥ 500</li> <li>Several other forms of OTC Niacin and fish</li> </ul>
CHOLESTEROL ABS ezetimibe (generic for Zetia)	omega-3 fatty acids (generic for Lovaza) ^{CL} VASCEPA (icosapent) ^{CL} <b>DRPTION INHIBITORS</b> NEXLIZET (bempedoic acid/ezetimibe) ^{NR,QL}	<ul> <li>Several other forms of OFC Machinand fish oil are also covered without prior authorization under Medicaid with a prescription</li> <li>Vascepa[®]: Approved for TG ≥ 500</li> </ul>

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PDL Updated November 3, 2020 Highlights indicated change from previous posting LIPOTROPICS, OTHER (continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ROPROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 9 (PCSK9) INHIBITORS	
	PRALUENT (alorocumab) ^{CL}	(ASCVD)
	REPATHA (evolocumab) ^{CL}	<ul> <li>heterozygous familial hypercholesterolemia (HeFH)</li> </ul>
		AND
		Maximized high-intensity statin WITH     ezetimibe for at 3 continuous months
		<ul> <li>Failure to reach target LDL-C levels: ASCVD - &lt; 70 mg/dL, HeFH - &lt; 100 mg/dL</li> </ul>
		Repatha [®] : Approved for:
		adult diagnoses of atherosclerotic cardiovascular disease (ASCVD)
		<ul> <li>heterozygous familial hypercholesterolemia (HeFH)</li> </ul>
		<ul> <li>homozygous familial hypercholesterolemia (HoFH) in age ≥ 13</li> </ul>
		statin-induce rhabdomyolysis
		AND
		Maximized high-intensity statin WITH     ezetimibe for 3+ continuous months
		<ul> <li>Failure to reach target LDL-C levels: ASCVD - &lt; 70 mg/dL, HeFH - &lt; 100 mg/dL</li> </ul>
		Concurrent use of maximally-tolerated statin must continue

### LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
STA atorvastatin (generic Lipitor) ^{QL} lovastatin (generic Mevacor) pravastatin (generic Pravachol)	TINS ALTOPREV (lovastatin ER) ^{CL} EZALLOR SPRINKLE (rosuvastatin) ^{QL} fluvastatin IR/ER (generic Lescol/	<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> </ul>
rosuvastatin (generic Crestor) simvastatin (generic Zocor)	Lescol XL) LIVALO (pitavastatin) ZYPITAMAG (pitavastatin)	<ul> <li>Drug-specific criteria:</li> <li>Altoprev[®]: One of the TWO trials must be IR lovastatin</li> <li>Combination products: Require clinical</li> </ul>
STATIN CON	<b>IBINATIONS</b>	reason why individual ingredients cannot be
	atorvastatin/amlodipine (generic Caduet) simvastatin/ezetimibe (generic Vytorin)	<ul> <li>used</li> <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>

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

PDL Updated November 3, 2020 *Highlights* indicated change from previous posting **MACROLIDES AND KETOLIDES, ORAL** 

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

#### METHOTREXATE

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting **MOVEMENT DISORDERS** 

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) ^{CL} tetrabenazine (generic for Xenazine) ^{CL}	INGREZZA (valbenazine) ^{CL} CAP, INITIATION PACK	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
		<ul> <li>Ingrezza: Diagnosis of Tardive Dyskinesia in adults and trial of Austedo</li> </ul>
		<ul> <li>tetrabenazine:Diagnosis of chorea with Huntington's Disease</li> </ul>

### **MULTIPLE SCLEROSIS DRUGS**

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

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### NITROFURAN DERIVATIVES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
nitrofurantoin macrocrystals <b>CAPSULE</b> (generic for Macrodantin) nitrofurantoin monohydrate- macrocrystals <b>CAPSULE</b> (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 <b>ONE</b> preferred agent within this drug class</li> </ul>

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PDL Updated November 3, 2020 Highlights indicated change from previous posting NSAIDs, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COX-I SE diclofenac sodium (generic for Voltaren) ibuprofen OTC, Rx (generic for Advil,	diclofenac potassium (generic for Cataflam, Zipsor) diclofenac SR (generic for Voltaren-XR)	<ul> <li>Non-preferred agents within COX- 1 SELECTIVE group will be approved for patients who have failed no less than 30-day trial of TWO preferred agents within this</li> </ul>
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 sulindac (generic for Clinoril)	diflunisal (generic for Dolobid) etodolac & SR (generic for Lodine/XL) fenoprofen (generic for Nalfon) flurbiprofen (generic for Ansaid) ibuprofen OTC (generic for Advil, Motrin) <b>CAPSULE</b> indomethacin ER (generic for Indocin) INDOCIN <b>RECTAL</b> , <b>SUSPENSION</b> ketoprofen & ER (generic for Orudis) meclofenamate (generic for Orudis) meclofenamate (generic for Ponstel) naproxen CR (generic for Naprelan) naproxen SUSPENSION (generic for Naprosyn) naproxen sodium (generic for Anaprox) <i>naproxen-esomeprazole (generic for</i> <i>Vimovo)</i> oxaprozin (generic for Daypro) piroxicam (generic for Feldene) QMIIZ ODT (meloxicam) ^{QL} RELAFEN DS (nabumetone) tolmetin (generic for Tolectin) Ketorolac Nasal ^{QL} (generic for Sprix)	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 menorrhagia

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PDL Updated November 3, 2020 *Highlights* indicated change from previous posting

### NSAIDs, ORAL (Continued)

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

### **NSAIDs, TOPICAL**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	diclofenac (generic for Pennsaid Solution) ^{CL} FLECTOR <b>PATCH</b> (diclofenac) ^{CL} <i>LICART <b>PATCH</b> (diclofenac)</i> PENNSAID <b>PACKET, PUMP</b> (diclofenac) ^{CL} VOLTAREN <b>GEL</b> (diclofenac) ^{CL}	<ul> <li>Drug Specific Criteria</li> <li>Flector[®]: 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</li> <li>Pennsaid[®]: Approved for osteoarthritis of the knees AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form</li> <li>Pennsaid[®] Pump: Requires clinical reason why 1.5% solution cannot be used</li> <li>Voltaren[®]: Approved for diagnosis of osteoarthritis AND trial of oral diclofenac OR clinical reason patient cannot be used</li> </ul>

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AL____Age Limit

PDL Updated November 3, 2020 *Highlights* indicated change from previous posting NOTE: Other oral oncology agents not listed here may also be available. See <u>https://nebraska.fhsc.com/default.asp</u> for coverage information and prior authorization status for products not listed.

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CDK 4/6 I IBRANCE (palbociclib)	NHIBITOR KISQALI (ribociclib)	Non-preferred agents DO NOT require a trial of a preferred agent,
	KISQALI FEMARA <b>CO-PACK</b> VERZENIO (abemaciclib)	but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
СНЕМОТ	HERAPY	- Drug-specific critera
cyclophosphamide XELODA (capecitabine)	capecitabine (generic for Xeloda) ^{CL}	<ul> <li>anastrozole: May be approved for malignant neoplasm of male breast (male breast cancer)</li> </ul>
HORMONE	BLOCKADE	capecitabine: Requires trial of Xeloda or clinical reason Xeloda
anastrozole (generic for Arimidex) exemestane (generic for Aromasin) letrozole (generic for Femara) tamoxifen citrate (generic for Nolvadex)	SOLTAMOX <b>SOLN</b> (tamoxifen) ^{CL} toremifene (generic for Fareston) ^{CL}	<ul> <li>Fareston[®]: Require clinical reason why tamoxifen cannot be used</li> <li>Ietrozole: 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) <i>lapatinib (generic Tykerb)^{CL,NR}</i> TALZENNA (talazoparib tosylate) ^{QL} TUKYSA(tucatinib) ^{NR,QL}	<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
	LL	<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent,</li> </ul>
mercaptopurine	PURIXAN (mercaptopurine)	but DO require an FDA-approved indication OR documentation submitted supporting off-label use
Α	ML	from current treatment guidelines
C IMBRUVICA (irutinib) LEUKERAN (chlorambucil) VENCLEXTA (venetoclax)	DAURISMO (glasdegib maleate) ^{QL} IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) ^{QL} XOSPATA (gilteritinib) ^{QL} LL COPIKTRA (duvelisib) ^{QL} 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>Tabloid: Prior authorization not required for age &lt;19</li> <li>Tasigna: Patients receiving Tasigna, which changed from preferred to non-preferred on 1-17-</li> </ul>
С	ML	19 will be allowed to continue
hydroxyurea (generic for Hydrea) imatinib (generic for Gleevec) ^{GL} MYLERAN (busulfan) SPRYCEL (dasatinib)	BOSULIF (bosutinib) GLEEVEC (imatinib) HYDREA (hydroxyurea) ICLUSIG (ponatinib) TASIGNA (nilotinib) ^{CL}	<ul> <li>therapy</li> <li>Xpovio: Indicated for relapsed or refractory multiple myeloma. Requires concomitant therapy with dexamethasone</li> </ul>
Μ	PN	_
JAKAFI (ruxolitinib)		
MYE	LOMA	
ALKERAN (melphalan) REVLIMID (lenalidomide)	FARYDAK (panobinostat) melphalan (generic for Alkeran) NINLARO (ixazomib) POMALYST (pomalidomide) THALOMID (thalidomide) XPOVIO (selinexor) ^{CL}	
от	HER	
MATULANE (procarbazine) TABLOID (thioguanine) tretinoin (generic for Vesanoid)	BRUKINSA (zanubrutinib) ^{NR,QL} CALQUENCE (acalabrutinib) ^{QL} INREBIC (fedratinib dihydrochloride) ^{QL} INQOVI (decitabine/cedazuridine) ^{NR} ONUREG (azacytidine) ^{NR} ZOLINZA (vorinostat)	

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
A	LK	<ul> <li>Non-preferred agents DO NOT</li> </ul>
ALECENSA (alectinib)	ALUNBRIG (brigatinib) LORBRENA (lorlatinib) ^{QL} ZYKADIA (ceritinib) <b>CAPSULE,</b> <i>TABLET</i>	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
ALK / RO	S1 / NTRK	
XALKORI (crizotinib)	ROZLYTREK (entrectinib) AL,QL	
EC	<b>F</b> R	
IRESSA (gefitinib) TAGRISSO (osimertinib)	erlotinib (generic for Tarceva) GILOTRIF (afatinib) TARCEVA (erlotinib) VIZIMPRO (dacomitinib) ^{QL}	
от	HER	-
	GAVRETO (pralsetinib) ^{NR,QL} HYCAMTIN (topotecan) RETEVMO (selpercatinib) ^{NR,AL} TABRECTA (capmatinib) ^{NR,QL}	_

### **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) <i>KOSELUGO (selumetinib)^{NR,AL}</i> LONSURF (trifluridine/tipiracil) <i>PEMAZYRE (pemigatinib)^{NR,QL}</i> RUBRACA (rucaparib) STIVARGA (regorafenib) TURALIO (pexidartinib) ^{QL} VITRAKVI (larotrectinib) <b>CAPSULE,</b> <b>SOLUTION</b> ^{QL}	<ul> <li>Non-preferred agents DO NOT require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines</li> </ul>

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AL__ Age Limit

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

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

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

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

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BASA ERIVEDGE (vismodegib)	L <b>CELL</b> ODOMZO (sonidegib) ^{CL}	<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>
BRAF M MEKINIST (trametinib) TAFINLAR (dabrafenib)	UTATION BRAFTOVI (encorafenib) COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)	<ul> <li>from current treatment guidelines</li> <li>Drug-specific critera</li> <li>Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17- 19 will be allowed to continue therapy</li> </ul>

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

CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

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

### **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) EMADINE (emedastine) epinastine (generic for Elestat) LASTACAFT (alcaftadine) olopatadine 0.1% (generic for Patanol) olopatadine 0.2% (generic for Pataday) PATADAY OTC (olopatadine 0.2%) ZERVIATE (certirizine) ^{AL,NR}	<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>

### **OPHTHALMICS, ANTIBIOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FLUOROQU	FLUOROQUINOLONES	
ciprofloxacin <b>SOLUTION</b> (generic for Ciloxan) MOXEZA (moxifloxacin) ofloxacin (generic for Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic for Zymaxid) levofloxacin 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</li> </ul>
MACRO	DLIDES	documented fungal infection
erythromycin	AZASITE (azithromycin) ^{CL}	
AMINOGL	COSIDES	
gentamicin SOLUTION	gentamicin OINTMENT	
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) ^{CL} neomycin/bacitracin/polymyxin B <b>OINTMENT</b> neomycin/polymyxin B/gramicidin NEOSPORIN (neomycin/polymyxin B/gramcidin) sulfacetamide <b>SOLUTION</b> (generic for Bleph-10) sulfacetamide <b>OINTMENT</b>	

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
neomycin/polymyxin/dexamethasone (generic for Maxitrol) sulfacetamide/prednisolone TOBRADEX <b>SUSPENSION,</b> <b>OINTMENT</b> (tobramycin and dexamethasone)	<ul> <li>BLEPHAMIDE (prednisolone and sulfacetamide)</li> <li>BLEPHAMIDE S.O.P.</li> <li>neomycin/polymyxin/HC</li> <li>neomycin/bacitracin/poly/HC</li> <li>PRED-G SUSPENSION, OINTMENT (prednisolone/gentamicin)</li> <li>tobramycin/dexamethasone</li> <li>SUSPENSION (generic for Tobradex)</li> <li>TOBRADEX S.T. (tobramycin and dexamethasone)</li> <li>ZYLET (loteprednol, tobramycin)</li> </ul>	<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>

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

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

### **OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
RESTASIS (cyclosporine) RESTASIS MULTIDOSE (cyclosporine)	CEQUA (cyclosporine) ^{QL} XIIDRA (lifitegrast)	<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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AL – Age Limit

### **OPHTHALMICS, GLAUCOMA**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIO	TICS	<ul> <li>Non-preferred agents will be</li> </ul>
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	DCKERS	
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) TIMOPTIC OCUDOSE TIMOPTIC XE (timolol gel forming solution)	
CARBONIC ANHYDRASE INHIBITORS		
dorzolamide (generic for Trusopt)	AZOPT (brinzolamide)	_
PROSTAGLANDIN ANALOGS		
latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATI	ON DRUGS	-
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine	
OTHER		•
RHOPRESSA (netarsudil) ^{CL} ROCKLATAN (netarsudil and latanoprost) ^{CL}		Drug-specific criteria: • Rhopressa and Rocklatan: Electronically approved for patients who have a trial of ONE generic agent, within ophthalmics- glaucoma within 60 days

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### PDL Updated November 3, 2020 Highlights indicated change from previous posting

### **OPIOID DEPENDENCE TREATMENTS**

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

### **OPIOID-REVERSAL TREATMENTS**

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

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

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

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^{NR} – Product was not reviewed - New Drug criteria will apply Page 65 of 79

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### **OTIC ANTIBIOTICS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CIPRODEX (ciprofloxacin/dexamethasone) neomycin/polymyxin/hydrocortisone (generic for Cortisporin) ofloxacin (generic for Floxin)	CIPRO HC (ciprofloxacin/ hydrocortisone) ciprofloxacin <i>ciprofloxacin/dexamethasone (generic</i> <i>for CIPRODEX)</i> ^{NR} 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
ADCIRCA (tadalafil) ^{CL} ambrisentan (generic Letairis) sildenafil <b>TABLET</b> (generic Revatio) ^{CL} TRACLEER <b>TABLET</b> (bosentan) TYVASO <b>INHALATION</b> (treprostinil) VENTAVIS <b>INHALATION</b> (iloprost)	ADEMPAS (riociguat) ^{CL} bosentan <b>TABLET</b> (generic Tracleer) LETAIRIS (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) sildenafil <b>SUSPENSION</b> (generic Revatio) ^{CL} tadalafil (generic for Adcirca) ^{CL} TRACLEER <b>TABLETS FOR</b> <b>SUSPENSION</b> (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:</li> <li>Adcirca®/Revatio®: Approved for diagnosis of Pulmonary Arterial Hypertension (PAH)</li> <li>Adempas®: PAH: Requires clinical reason preferred agent cannot be used CTEPH: Approved for persistent/recurrent diagnosis after surgical treatment or inoperable CTEPH NOT for use in Pregnancy</li> <li>sildenafil suspension: Requires clinical reason why sildenafil tablets cannot be used</li> </ul>

#### **PANCREATIC ENZYMES**

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

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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 – A

AL – Age Limit

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting

### PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD LITTLE ANIMALS VITAMINS CHEW OTC (pedi multivit 91/iron fum) CHEW 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)	AQUADEKS (pedi multivit 40/phytonadione) ESCAVITE (pedi multivit 47/iron/fluoride) ESCAVITE D (pedi multivit 78/iron/fluoride) <b>CHEW</b> ESCAVITE LQ (pedi multivit 86/iron/fluoride) <b>CHEW</b> FLORIVA (pedi multivit 85/fluoride) <b>CHEW</b> FLORIVA PLUS OTC and Rx (pedi multivit 130/fluoride) <b>DROPS</b> multivit A, B, D, E, K, ZN (pediatric multivit 153/D3/K) POLY-VI-FLOR (pedi multivit 33/fluoride) <b>CHEW</b> POLY-VI-FLOR (pedi multivit 33/fluoride) <b>DROPS</b> POLY-VI-FLOR w/IRON (pedi multivit 33/fluoride/iron) <b>CHEW</b> POLY-VI-FLOR w/IRON (pedi multivit 37/fluoride/iron) <b>DROPS</b> QUFLORA OTC and Rx (pedi multivit 84/fluoride) QUFLORA FE (pedi multivit 142/iron/fluoride) TRI-VI-FLOR (ped multivit A, C, D3, 38/fluoride)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> <li>Drug specific criteria:         <ul> <li>Aquadeks: Approved for diagnosi of Cystic Fibrosis</li> </ul> </li> </ul>

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### PDL Updated November 3, 2020 *Highlights* indicated change from previous posting

### PENICILLINS

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

#### **PHOSPHATE BINDERS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcium acetate <b>TABLET, CAPSULE</b> CALPHRON OTC (calcium acetate) sevelamer carbonate (generic Renvela)	AURYXIA (ferric citrate) ELIPHOS (calcium acetate) Ianthanum (generic FOSRENOL) PHOSLO (calcium acetate) PHOSLYRA (calcium acetate) RENAGEL (sevelamer HCI) sevelamer HCI (generic Renagel) 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) ^{CL}	<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</li> </ul>

peripheral artery disease (PAD) Use with aspirin and/or clopidogrel

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AL – Age Limit

#### **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 <b>SOFTGEL</b> complete natal dha (pnv2/iron b-g suc-p/fa/omega-3) calcium-pnv 28-1-250mg <b>SOFTGEL</b> classic prenatal <b>TABLET</b> (prenatal vit/fe furn/fa) COMPLETENATE <b>CHEWABLE</b> CONCEPT DHA <b>CAPSULE</b> elite-ob <b>CAPLET</b> (fe c/fa) MARNATAL-F <b>CAPSULE</b> PRENATA TAB <b>CHEW</b> pnv with ca, #72/iron/fa pnv-ob+dha combo pack (pnv22/iron cbn&gluc/fa/dss/dha) pnv-vp-u <b>CAPSULE</b> prenaissance <b>CAPSULE</b> (pnv80/iron furn/fa/dss/dha) prenaissance plus <b>SOFTGEL</b> (pnv69/iron/fa/dss/dha) prenatal vitamin <b>TABLET</b> (pnv#124/iron/fa) prenatal vitamin <b>TABLET</b> (pnv#78/iron/fa) PUREFE PLUS PUREFE OB PLUS TARON-PREX PRENATAL TRINATAL RX 1 triveen-duo dha combo pack (pnv53/iron b-g hcl-p/fa/omega3) trust natal dha (pnv2/iron b-g suc-p/fa/omega-3) virtprex <b>CAPSULE</b> (pnv66/iron furn/fa/dss/dha) virt-nate dha <b>SOFTGEL</b> (pnv7a78/iron/fa1/dha) virt-pn plus <b>SOFTGEL</b> (pnv80/iron furn/fa/dss/dha) virt-pn <b>TABLET</b> (pnv#1-iron furn/fa.cmb no.1) virt-pn plus <b>SOFTGEL</b> (pnv80/iron furn/fa/dss/dha) virt-pn <b>TABLET</b> (pnv80/iron furn/fa/dss/dha) virt-pn plus <b>SOFTGEL</b> (pnv/ca no.68/iron/fa1/dha)	folivane-ob CAPSULE (pnv#15/iron fum & ps cmp/fa) niva-plus TABLET (pnv with ca,no.74/iron/fa) pnv-dha SOFTGEL (pnv combo#47/iron/fa #1/dha) taron-c dha CAPSULE (pnv#16/iron fum &ps/fa/om-3) virt-c dha SOFTGEL (pnv combo#47/iron/fa #1/dha) zatean-pn dha CAPSULE (pnv #47/iron/fa #1/dha)	<ul> <li>Non-preferred agents will be approved for patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class</li> </ul>

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

### PROGESTERONE (hydroxyprogesterone caproate)

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

### **PROTON PUMP INHIBITORS**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
omeprazole (generic Prilosec) <b>RX</b> pantoprazole (generic Protonix) ^{QL}	DEXILANT (dexlansoprazole) esomeprazole magnesium (generic Nexium) esomeprazole strontium lansoprazole (generic Prevacid) NEXIUM <b>SUSPENSION</b> (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) <i>pantoprazole GRANULES</i> ^{NR,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 compounded suspension.</li> <li>Patients ≥ 5 years if age- Only approve non-preferred for GI diagnosis if:                 <ul> <li>Child can not swallow whole generic omeprazole capsules OR,</li> <li>Documentation that contents of capsule may not be sprinkled in applesauce</li> <li>Paper Solutab and the sprinkled in applesauce</li> <li>Privacid Solutab and the sprinkled in applesauce</li> <li>Prevacid Solutab and the sprinkled in applesauce</li> <li>Non-preferred for GI compounded suspension.</li> <li>Patients prove and the sprinkled in applesauce</li> <li>Provementation that contents of capsule may not be sprinkled in applesauce</li> <li>Privation applesauce</li> <li>Provementation that contents of capsule may not be sprinkled in applesauce</li> <li>Provementation that contents of capsule may not be sprinkled in applesauce</li></ul></li></ul></li></ul>

### SEDATIVE HYPNOTICS

BENZODIAZEPINES       -         temazepam 15mg, 30mg (generic for Restorii)       estazolam (generic for ProSom) flurazepam (generic for Restorii) 7.5mg, 22.5mg triazolam (generic for Restorii) 7.5mg, 22.5mg       -         oTHERS       BELSOMRA (suvorexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,QL} doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopiclone (generic for Cameto Lunesta) HETLUOZ (tasimeteon) ^{CL} rameteon (generic for Ambien)       BELSOMRA (suvorexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,QL} doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopiclone (generic for Ambien)       -         VIETUOZ (tasimeteon) ^{CL} rameteon (generic for Ambien)       -       -         VIETUOZ (tasimeteon) ^{CL} rameteon (generic for Intermezzo)       -       -         VIETUOZ (tasimeteon) ^{CL} rameteon (generic for Intermezzo)       -       -         Solpidem SL (generic for Intermezzo)       -       -       -         Order and comparison of solution of solution of solution is solution is solution is solution of solution is solution is solution is solutis solution is	temazepam 15mg, 30mg (generic for Restoril)       estazolam (generic for ProSom) flurazepam (generic for Dalmane) temazepam (generic for Restoril) 7.5mg, 22.5mg       ER: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used         OTHERS         zaleplon (generic for Sonata) zolpidem (generic for Ambien)       BELSOMRA (suvorexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,MR,QL} doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopicione (generic for Lunesta) HETLIOZ (tasimelteon) ^{CL} ramelteon (generic for Rozerem) zolpidem SL (generic for Intermezzo)       flurazepam/triazolam: Requires trial of preferred benzodiazepine sublingual) eszopicione (generic for Intermezzo)         0       Kequires a trial with generic zolpidem within the last 12 months AND clinical reason why zaleplon and preferred benzodiazepine fur action of swallowing disorder         0       Contraindication of swallowing disorder         0       Requires trial with generic zolpidem within last 12 months AND clinical reason why zaleplon AND preferred benzodiazepine cannot be used         0       Contraindication to preferred oral sedative hypnotics         0       Contraindication to preferred oral sedative hypnotics         0       Requires clinical reason why t5mg/30mg cannot be used	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	<ul> <li>zolpidem SL: Requires clinical reason why half of zolpidem tablet</li> </ul>	BENZODI temazepam 15mg, 30mg (generic for Restoril) OTH zaleplon (generic for Sonata)	AZEPINES estazolam (generic for ProSom) flurazepam (generic for Dalmane) temazepam (generic for Restoril) 7.5mg, 22.5mg triazolam (generic for Halcion) ERS BELSOMRA (suvorexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,NR,QL} doxepin (generic for Silenor) EDLUAR (zolpidem sublingual) eszopiclone (generic for Lunesta) HETLIOZ (tasimelteon) ^{CL} ramelteon (generic for Rozerem) zolpidem ER (generic for Ambien CR)	Lunesta®/ Rozerem®/zolpidem ER: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used Edluar®: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used and Requires documentation of swallowing disorder flurazepam/triazolam: Requires trial of preferred benzodiazepine Hetlioz®: Requires trial with generic zolpidem within last 12 months AND clinical reason why zaleplon AND preferred benzodiazepine cannot be used Silenor®: Must meet ONE of the following: • Contraindication to preferred oral sedative hypnotics • Medical necessity for doxepin dose < 10mg • Age greater than 65 years old or hepatic impairment (3mg dose will be approved if this criteria is met) temazepam 7.5mg/22.5mg: Requires clinical reason why 15mg/30mg cannot be used zolpidem/zolpidem ER: Maximum daily dose for females: Zolpidem 5mg; Zolpidem SL: Requires clinical reason why half of zolpidem tablet

### PDL Updated November 3, 2020 Highlights indicated change from previous posting

### 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>

### SKELETAL MUSCLE RELAXANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
baclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) ^{QL} methocarbamol (generic Robaxin) tizanidine <b>TABLET</b> (generic Zanaflex)	carisoprodol (generic Soma) ^{CL,QL} carisoprodol compound cyclobenzaprine ER (generic Amrix) ^{CL} dantrolene (generic Dantrium) FEXMID (cyclobenzaprine ER) LORZONE (chlorzoxazone) ^{CL} metaxalone (generic Skelaxin) NORGESIC FORTE (orphenadrine/ASA/caffeine) orphenadrine ER PARAFON FORTE (chlorzoxazone) tizanidine <b>CAPSULE</b> ZANAFLEX (tizanidine) <b>CAPSULE,</b> <b>TABLET</b>	<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 350mg generic strength cannot be used</li> <li>Soma® 250mg: Requires clinical reason why 350mg generic strength cannot be used</li> </ul>

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AL__ Age Limit

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

### 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 hydrocortisone/aloe OINTMENT, CREAM SCALPICIN OTC (hydrocortisone)	<ul> <li>ALA-CORT (hydrocortisone) CREAM</li> <li>ALA-SCALP HP (hydrocortisone)</li> <li>alclometasone dipropionate (generic for Aclovate)</li> <li>CAPEX SHAMPOO (fluocinolone)</li> <li>DESONATE (desonide) GEL</li> <li>desonide LOTION (generic for Desowen)</li> <li>desonide CREAM, OINTMENT (generic for former products Desowen, Tridesilon)</li> <li>fluocinolone 0.01% OIL (generic for DERMA-SMOOTHE-FS)</li> <li>MICORT-HC (hydrocortisone)</li> <li>TEXACORT (hydrocortisone)</li> </ul>	will be approved for patients who have failed a trial of ONE preferred agent within this drug class
MEDIUM	, , ,	<ul> <li>Medium Potency Non-preferred</li> </ul>
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 <b>LOTION</b> (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 PC	DTENCY	<ul> <li>High Potency Non-preferred</li> </ul>
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, SOLN ^{WR} 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 HIGH	I POTENCY	<ul> <li>Very High Potency Non-preferred</li> </ul>
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} 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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^{NR} – Product was not reviewed - New Drug criteria will apply

### STIMULANTS AND RELATED AGENTSAL

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>
Amphetar	Amphetamine type	
amphetamine salt combination ER (generic for Adderall XR) amphetamine salt combination IR VYVANSE (lisdexamfetamine) CAPSULE, CHEWABLE	<ul> <li>ADDERALL XR (amphetamine salt combo)</li> <li>ADZENYS XR (amphetamine)</li> <li>amphetamine ER (generic for Adzenys ER) SUSPENSION</li> <li>amphetamine sulfate (generic for Evekeo)</li> <li>dextroamphetamine (generic for Dexedrine)</li> <li>dextroamphetamine SOLUTION (generic for Procentra)</li> <li>dextroamphetamine ER (generic for Dexedrine ER)</li> <li>DYANAVEL XR (amphetamine)</li> <li>EVEKEO ODT (amphetamine sulfate)</li> <li>MYDAYIS (amphetamine salt combo)^{QL}</li> <li>methamphetamine (generic for Desoxyn)</li> <li>ZENZEDI (dextroamphetamine)</li> </ul>	<ul> <li>agent within this drug class</li> <li>Drug-specific criteria: <ul> <li>Procentra[®]: May be approved with documentation of swallowing disorder</li> </ul> </li> <li>Zenzedi[®]: Requires clinical reason generic dextroamphetamine IR cannot be used</li> </ul>

### STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents Non-Preferred Agents	Prior Authorization/Class Criteria
Methylphenidate typeAPTENSIO XR (methylphenidate) dexmethylphenidate (generic for Focalin IR)FOCALIN XR (dexmethylphenidate) methylphenidate)METHYLIN SOLUTION (methylphenidate 30/70 (generic for Metadate CD)methylphenidate SOLUTION (generic for Methylin) methylphenidate ER 10mg, 20mg (generic for Ritalin SR, Metadate ER)methylphenidate ER 18mg, 27mg, 36mg, 54mg (generic Concerta)QLQUILLICHEW ER CHEWTAB (methylphenidate)QUILLICHEW ER CHEWTAB (methylphenidate)QUILLICHEW ER CHEWTAB (methylphenidate)QUILLICATEW ER CHEWTAB (methylphenidate)QUILLICATEW ER CHEWTAB (methylphenidate)Methylphenidate)QUILLIVANT XR SUSP (methylphenidate)QUILLIVANT XR SUSP (methylphenidate)QUILLIVANT XR SUSP (methylphenidate)	Non-preferred agents will be approved for patients who have 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 Orug-specific criteria: Daytrana®: May be approved in history of substance use disorder by parent, caregiver, or patient. May be approved with documentation of difficulty swallowing

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### STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

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

PDL Updated November 3, 2020 Highlights indicated change from previous posting

### **TETRACYCLINES**

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR (generic Vibramycin) doxycycline monohydrate <b>50MG</b> , <b>100MG CAPSULE</b> doxycycline monohydrate <b>SUSP</b> , <b>TABLET</b> (generic Vibramycin) minocycline HCI <b>CAPSULE</b> , <b>TABLET</b> (generic Dynacin/ Minocin/Myrac)	<ul> <li>demeclocycline (generic Declomycin)^{CL}</li> <li>DORYX MPC DR (doxycycline pelletized)</li> <li>doxycycline hyclate DR (generic Doryx)</li> <li>doxycycline monohydrate 40MG, 75MG and 150MG CAPSULES (generic for Adoxa/Monodox/ Oracea)</li> <li>minocycline HCI ER (generic Solodyn)</li> <li>NUZYRA (omadacycline)</li> <li>tetracycline</li> <li>VIBRAMYCIN SUSP (doxycycline)</li> <li>XIMINO (minocycline ER)^{QL}</li> </ul>	<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>

### **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>	EUTHYROX (levothyroxine) LEVO-T (levothyroxine) THYROLAR <b>TABLET</b> (liotrix) TIROSINT <b>TABLET</b> (levothyroxine) TIROSINT-SOL (LIQUID) (levothyroxine) ^{CL}	<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	ORAL	
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) GIAZO (balsalazide) mesalamine ER (generic Apriso) mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	<ul> <li>approved for patients who have failed a trial of ONE preferred agent within this drug class</li> <li>Drug-specific criteria:</li> <li>Asacol HD[®]/Delzicol DR[®]/Lialda[®]/Pentasa[®]: Requires clinical reason why preferred mesalamine products cannot be used</li> <li>Giazo[®]: Requires clinical reason why generic balsalazide cannot be</li> </ul>
REC	TAL	used
CANASA (mesalamine)	mesalamine <b>ENEMA</b> (generic Rowasa) mesalamine <b>SUPPOSITORY</b> (generic Canasa) UCERIS (budesonide)	<ul> <li>NOT covered in females</li> </ul>

### UTERINE DISORDER TREATMENT

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORILISSA (elagolix sodium) ^{QL,CL}	ORIAHNN (elagolix/ estradiol/ norethidrone) AL,NR	Drug-specific criteria: Orilissa: Requires an FDA approved indication, must follow FDA dosing guidelines, and have had a trial and failure of an NSAID and oral contraceptive

### **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	<ul> <li>BIDIL (isosorbide dinitrate/ hydralazine)^{CL}</li> <li>GONITRO (nitroglycerin)</li> <li>NITRO-BID <b>OINTMENT</b> (nitroglycerin)</li> <li>NITRO-DUR (nitroglycerin)</li> <li>nitroglycerin <b>TRANSLINGUAL</b> (generic Nitrolingual)</li> <li>NITROMIST (nitroglycerin)</li> </ul>	<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> </ul>

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