



DEPT. OF HEALTH AND HUMAN SERVICES

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

January 2021 PDL Contains November 2020 P&T Changes Noted in Red Font that Become Effective January 21, 2021

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

• Opioids- The maximum opioid dose covered will decrease from 120 Morphine Milligram Equivalents (MME) per day to 90 Morphine Milligram Equivalents (MME) per day. (beginning December 1, 2020)

Non-Preferred Drug Coverage

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

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

Specific Class Prior Authorization forms can be found within the PDL class listings and at:

https://nebraska.fhsc.com/priorauth/paforms.asp

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

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

Documentation of Medical Necessity PA Form

For a complete list of Claims Limitations visit: https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) AKLIEF (trifarotene) AL ALTRENO (tretinoin) AL AMZEEQ (minocycline) ARAZLO (tazarotene) AL ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) BENZACLIN PUMP (clindamycin/BPO) BENZEFOAM (benzoyl peroxide) Penzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzeyro) benzoyl peroxide GEL Rx benzoyl peroxide TOWELETTE OTC clindamycin FOAM, LOTION clindamycin GEL clindamycin/BPO (generic Acanya, Benzaclin) GEL clindamycin/tretinoin (generic Veltin, Ziana) dapsone (generic Aczone) EPIDUO FORTE GEL PUMP (adapalene/BPO) erythromycin GEL, PLEDGET erythromycin-BPO (generic for Benzamycin) EVOCLIN (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONACE PLUS (sulfacetamide sodium) PLIXDA (adapalene) SWAB RETIN-A GEL, CREAMAL (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 Output Description: Output Desc

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ALZHEIMER'S AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHOLINESTER	CHOLINESTERASE INHIBITORS	
donepezil (generic for Aricept) donepezil ODT (generic for Aricept ODT) EXELON Transdermal (rivastigmine)	donepezil 23 (generic for Aricept 23) galantamine (generic for Razadyne) SOLUTION, TABLET galantamine ER (generic for Razadyne ER) rivastigmine (generic for Exelon) OR ANTAGONIST	approved for patients who have failed a 120-day trial of ONE preferred agent within this drug class within the last 6 months OR Current, stabilized therapy of the non-preferred agent within the previous 45 days
memantine (generic for Namenda)	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)

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) OXYCONTINCL (oxycodone ER)	ARYMO ER (morphine sulfate) QL BELBUCA (buprenorphine) CL buccal buprenorphine PATCH (generic Butrans) QL EMBEDA (morphine sulfate/ naltrexone) DURAGESIC MATRIX (fentanyl) QL fentanyl 37.5, 62.5, 87.5 mcg PATCHQL hydrocodone bitartrate ER (generic for Zohydro ER) hydromorphone ER (generic for 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	The Center for Disease Control (CDC) does not recommend long acting opioids when beginning opioid treatment. Preferred agents require previous use of a long acting opioid or documentation of a trial on a short acting agent within 90 days Non-preferred agents will be approved with failure on, or intolerance to TWO preferred agents within this drug class Drug-specific criteria: Methadone: Will only be approved for use in pain control or end of life care. Trial of preferred agent not required for end of life care Oxycontin®: Pain contract required for maximum quantity authorization

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANALGESICS, OPIOID SHORT-ACTINGQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acetaminophen/codeine ELIXIR, TABLET codeine TABLET nydrocodone/APAP SOLUTION, TABLET nydrocodone/ibuprofen nydromorphone TABLET morphine CONC SOLUTION, SOLUTION, TABLET oxycodone TABLET, SOLUTION oxycodone/APAP PROLATE (oxycodone/acetaminophen) tramadol TABLET ^{AL}	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/APAP SOLUTION oxycodone/APAP SOLUTION oxycodone/aspirin oxycodone/aspirin oxycodone/ibuprofen oxymorphone IR (generic Opana) pentazocine/naloxone PRIMLEV (oxycodone/acetaminophen) ROXICODONE TABLET (oxycodone) ROXYBOND (oxycodone/APAP)	 Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the las 12 months Note: for short acting opiate tablets 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

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 –

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANALGESICS, OPIOID SHORT-ACTINGQL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NA	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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
estosterone PUMP (generic Androgel) ^{CL}	ANDRODERM (testosterone) ^{CL} NATESTO (testosterone) ^{CL} testosterone PACKET (generic Androgel) ^{CL} testosterone GEL, PACKET, PUMP (generic Vogelxo) 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 la 6 months Drug-specific criteria: Androderm®/Androgel®:

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

AL Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANGIOTENSIN MODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ACE INHIBITORS		Non-preferred agents will be
	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 tablet in not operatorists.
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 REC	EPTOR BLOCKERS	
irbesartan (generic Avapro) losartan (generic Cozaar) valsartan (generic Diovan)	candesartan (generic Atacand) EDARBI (azilsartan) eprosartan (generic Teveten) olmesartan (generic Benicar) telmisartan (generic Micardis)	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANGIOTENSIN MODULATORS (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANGIOTENSIN RECEPTOR BLO	CKER/DIURETIC COMBINATIONS	Non-preferred agents will be
irbesartan/HCTZ (generic Avalide) losartan/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
ANGIOTENSIN	N MODULATOR/	 Angiotensin Modulator/Calcium Channel Blocker Combinations:
CALCIUM CHANNEL BL	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) amlodipine/valsartan/HCTZ (generic Exforge HCT) PRESTALIA (perindopril/amlodipine) trandolapril/verapamil (generic Tarka)	approved if there has been a trial and failure of preferred agent
DIRECT REM	IN INITIDITORS	 Direct Renin Inhibitors/Direct Renin Inhibitor Combinations:
DIRECT REN	IN INHIBITORS aliskiren (generic Tekturna) ^{QL}	May be approved witha history of TWO preferred ACE Inhibitors or Angiotensin Receptor Blockers
DIRECT RENIN INHIE	SITOR COMBINATIONS	within the last 12 months
	TEKTURNA/HCT (aliskiren/HCTZ)	
NEPRILYSIN INHIBITOR COMBINATION		
ENTRESTO (sacubitril/valsartan)QL		
ANGIOTENSIN RECEPTOR BLOCKER/BETA-BLOCKER COMBINATIONS		
	BYVALSON (nevibolol/valsartan)	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTHELMINTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
albendazole (generic for Albenza) BILTRICIDE (praziquantel) ivermectin (generic for Stromectol)	ALBENZA (albendazole) 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 agent within this drug class within the last 6 months Drug-specific criteria: Emverm: Approval will be considered for indications not covered by preferred agents

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTI-ALLERGENS, ORAL

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIBIOTICS, GASTROINTESTINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FIRVANQ (vancomycin) SOLUTION metronidazole TABLET neomycin	ALINIA (nitazoxanide) ^{CL} SUSPENSION DIFICID (fidaxomicin) ^{CL} TABLET, SUSP ^{NR} FLAGYL ER (metronidazole) ^{CL} Metronidazole ^{CL} CAPSULE nitazoxanide (generic Alinia) TABLET ^{AL, CL,NR, QL} 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®/Metronidazole 375mg capsules and Flagyl ER®/ Metronidazole 750mg ER tabs: Clinical reason why the 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 Lomotif® AND Imodium®

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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} tobramycin (generic for Bethkis) ^{NR} 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
		 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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) metronidazole, vaginal	Non-preferred agents will be approved for patients who have failed a therapeutic trial (duration = 3 days) with ONE preferred agent within this drug class within the last 6 months

ANTICOAGULANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) noxaparin (generic Lovenox) PRADAXA (dabigatran) varfarin (generic Coumadin) (ARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg (ARELTO (rivaroxaban) 2.5 mg ^{CL,QL} (ARELTO 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 thrombos (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, myocardia infarction, and stroke) in patients with chronic coronary artery disease or peripheral artery disease

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIEMETICS/ANTIVERTIGO AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BINOIDS	Non-preferred agents will be approved for patients who have
dronabinol (generic Marinol) ^{AL}	CESAMET (nabilone)	failed ONE preferred agent within this drug class within the same
5HT3 RECEPTO	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	Regimens include: AC combination (Doxorubicin or Epirubicin with Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine,
TRADITIONAL	ANTIEMETICS	Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine, 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
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 SUPPOSITORY (generic Compazine) promethazine SUPPOSITORY 50mg scopolamine TRANSDERMAL trimethobenzamide TABLET (generic Tigan)	

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIFUNGALS, ORAL

Preferred Agents Non-Preferred Agents Prior Authorization/Class Criteria

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

clotrimazole (mucous membrane, troche)

fluconazole SUSPENSION, TABLET (generic Diflucan) griseofulvin SUSPENSION griseofulvin microsized TABLET nystatin SUSPENSION, TABLET terbinafine (generic Lamisil) CRESEMBA (isavuconazonium)^{CL} flucytosine (generic Ancobon)^{CL} griseofulvin ultramicrosize (generic GRIS-PEG)

itraconazole (generic Sporanox)^{CL} ketoconazole (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 candidiasis refractory to itraconazole and/or fluconazole
- Onmel[®]: Requires trial and failure or contraindication to terbinafine
- Sporanox®/Itraconazole: Approved for diagnosis of Aspergillosis, Blastomycosis, Histoplasmosis, Onychomycosis due to terbinafineresistant dermatophytes, Oropharyngeal/ esophageal candidiasis refractory to fluconazole
- Sporanox[®]: Requires trial and failure of generic itraconazole
- Sporanox® Liquid: Clinical reason solid oral cannot be used
- Tolsura: Approved 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 Fusarium spp., Oropharyngeal/esophageal candidiasis refractory to fluconazole

ANTIFUNGALS, TOPICAL

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

CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL – Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
·	FUNGAL	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	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
	ROID COMBINATIONS	
clotrimazole/betamethasone CREAM (generic Lotrisone)	clotrimazole/betamethasone LOTION (generic Lotrisone) nystatin/triamcinolone (generic Mycolog)	

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIHISTAMINES, MINIMALLY SEDATING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
cetirizine TABLET, SOLUTION (Rx only) (generic for Zyrtec) loratadine TABLET, SOLUTION (generic for Claritin) levocetirizine TABLET (generic for Xyzal)	cetirizine CHEWABLE (generic for Zyrtec) cetirizine SOLUTION (OTC) desloratadine (generic for Clarinex) desloratadine ODT (generic for Clarinex Reditabs) fexofenadine (generic for Allegra) fexofenadine 180mg (generic for Allegra 180mg) levocetirizine (generic for Xyzal) SOLUTION loratadine CAPSULE, CHEWABLE, ODT (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) MITIGARE (colchicine) probenecid probenecid/colchicine (generic for Col- Probenecid)	colchicine TABLET (generic for Colcrys) ^{CL} colchicine CAPSULE (generic for Mitigare) febuxostat (generic for Uloric) ^{CL} GLOPERBA SOLN (colchicine) ^{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 AL –

AL Age Limit

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIMIGRAINE AGENTS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
JOVY (fremanezumab-vfrm) CL, QL JOVY AUTOINJECTOR (fremanezumab-vfrm)CL, QL MGALITY 120 mg/mL (galcanezumab-gnlm) CL, QL PEN, SYRINGE IURTEC 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, Emgaility 100mg is recommended dosing for Episodic Cluster Headache Aimovig, Ajovy and Emgality 120mg: Require ≥ 4 migraines per month for ≥ 3 months and has tried and failed a ≥ 1 month trial of two medications listed in the 2012 American Academy of Neurology/American Headache Society guidelines (examples include: antidepressants (amitriptyline, venlafaxine), Beta blockers (propranolol, metroprolol, timolol, atenolol), anti-epileptics (valproate, topiramate), ACE/ARB (lisinopril, candesartan) In addition, Aimovig requires a trial of Emgality 120mg or Ajovy or clinical, patient specific reason that a preferred agent cannot be used

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIMIGRAINE AGENTS, TRIPTANSQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORAL		Non-preferred agents will be
rizatriptan (generic Maxalt)	almotriptan (generic Axert)	approved for patients who have failed ALL preferred agents within
rizatriptan ODT (generic Maxalt MLT)	eletriptan (generic Relpax)	this drug class
sumatriptan	frovatriptan (generic Frova)	.
	IMITREX (sumatriptan)	Drug-specific criteria:
	naratriptan (generic Amerge)	 Sumavel® Dosepro: Requires clinical reason sumatriptan
	RELPAX (eletriptan) ^{QL}	injection cannot be used
	sumatriptan/naproxen (generic Treximet)	 Onzetra, Zembrace: approved for patients who have failed ALL
	zolmitriptan (generic Zomig/Zomig ZMT)	preferred agents
NA	SAL	
sumatriptan	IMITREX (sumatriptan)	
	ONZETRA XSAIL (sumatriptan)	
	TOSYMRA (sumatriptan)	
	ZOMIG (zolmitriptan)	
INJECTABLE		
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 ivermectin (generic Sklice) ^{NR} lindane malathion (generic Ovide) SKLICE (ivermectin) spinosad (generic Natroba) VANALICE (piperonyl butoxide/pyrethrins)	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ANTIPARKINSON'S AGENTS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOLINERGICS		Non-preferred agents will be
benztropine (generic for Cogentin) trihexyphenidyl (generic for Artane)		approved for patients who have failed ONE preferred agent within this drug class
COMT INI	HIBITORS	triis drug class
	entacapone (generic for Comtan) ONGENTYS (Opicapone) ^{NR,QL} tolcapone (generic for Tasmar)	 Drug-specific criteria: Carbidopa/Levodopa ODT: Approved for documented swallowing disorder COMT Inhibitors: Approved if using
DOPAMINE	AGONISTS	as add-on therapy with levodopa-
pramipexole (generic for Mirapex) ropinirole (generic for Requip)	bromocriptine (generic for Parlodel) ropinirole ER (generic for Requip ER) ^{CL} NEUPRO (rotigotine) ^{CL} pramipexole ER (generic for Mirapex ER) ^{CL} ropinirole ER (generic for Requip XL) ^{CL} ropinirole ER (generic for Requip XL) ^{CL}	 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 Inbrija: Approval upon diagnosis of Parkinson's disease and concurrent treatment with carbidopa/levodopa agent Neupro®:
MAO-B IN	HIBITORS	For Parkinsons: Clinical reason
selegiline CAPSULE, TABLET (generic for Eldepryl)	rasagiline (generic for Azilect) QL XADAGO (safinamide) ZELAPAR (selegiline)CL	required why preferred agent cannot be used For Restless Leg (RLS): Requires trial OR Contraindication to ropinirole AND pramipexole
OTHER ANTIPARI	KINSON'S DRUGS	Nourianz: Approval upon diagnosis of

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

amantadine CAPSULE, SYRUP
TABLET (generic for Symmetrel)
carbidopa/levodopa (generic for
Sinemet)
carbidopa/levodopa ER (generic for
Sinemet CR)
levodopa/carbidopa/entacapone
(generic for Stalevo)

APOKYN (apomorphine) SUB-Q
carbidopa (generic for Lodosyn)
carbidopa/levodopa ODT (generic for
Parcopa)
DUOPA (carbidopa/levodopa)
GOCOVRI (amantadine)^{QL}
INBRIJA (levodopa) INHALER^{CL,QL}
KYNMOBI (apomorphine)^{QL, KIT,}
SUBLINGUAL
NOURIANZ (istradefylline)^{CL,QL}
OSMOLEX ER (amantadine)^{QL}

RYTARY (carbidopa/levodopa)
STALEVO
(levodopa/carbidopa/entacapone)

Parkinson's disease and concurrent treatment with carbidopa/levodopa agent

- Osmolex ER: Required diagnosis of Parkinson's disease or drug-induced extrapyramidal reactions and had trial of or is intolerant to amantadine IR
- Pramipexole ER: Required diagnosis of Parkinson's along with preferred agent trial
- Ropinerole ER: Required diagnosis of Parkinson's along with preferred agent trial
- Zelapar®: Approved for documented swallowing disorder

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
--	------------------	----------------------	------------------------------------

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

ANTIVIRALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTI-HERPETIC DRUGS		Non-preferred agents will be
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-INFLUENZA DRUGS		- Drug apositio oritorio:
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

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

Page **23** of **93**

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) orazepam INTENSOL , TABLET (generic for Ativan)	alprazolam ER (generic for Xanax XR) alprazolam ODT alprazolam INTENSOL ^{CL} clorazepate (generic for Tranxene-T) diazepam INTENSOL ^{CL} lorazepam 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 trial of diazepam solution OR lorazepam Intensol®

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

BETA BLOCKERS, ORAL

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

BILE SALTS

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

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

BONE RESORPTION SUPRESSION AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BISPHOSPHONATES		Non-preferred agents will be
alendronate (generic Fosamax) TABLET ibandronate (generic Boniva) ^{QL}	alendronate SOLUTION (generic Fosamax) ^{QL} ATELVIA DR (risedronate)	approved for patients who have failed a trial of ONE preferred agent within the same group
	BINOSTO (alendronate)	Drug-specific criteria:
	etidronate disodium (generic Didronel) FOSAMAX PLUS DQL	individual agents without prior authorization
	risedronate (generic Actonel) ^{QL}	 Atelvia DR®: Requires clinical reason alendronate cannot be taken on an empty stomach
		Binosto®: Requires clinical reason why
	PRESSION AND RELATED DRUGS	alendronate tablets OR Fosamax® solution cannot be used
calcitonin-salmon NASAL raloxifene (generic Evista)	EVISTA (raloxifene) FORTEO (teriparatide) ^{QL}	Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification
	Teriparatide ^{QL}	Forteo®: Covered for high risk of fracture
	TYMLOS (abaloparatide)	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
		o Rheumatoid Arthritis
		 Postmenopausal women with BMD T- score ≤ -2.5 at any site with any clinical risk factors
		 More than 2 units of alcohol per day
		 Current smoker Men with primary or hypogonadal osteoporosis
		 Osteoporosis associated with sustained systemic glucocorticoid therapy
		Trial of calcitonin-salmon not required

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS

Preferred Agents Non-Preferred Agents	Prior Authorization/Class Criteria
ALPHA BLOCKERS alfuzosin (generic Uroxatral) doxazosin (generic Cardura) tamsulosin (generic Flomax) CARDURA XL (doxazosin) silodosin (generic Rapaflo)	Prior Authorization/Class Criteria Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class ug-specific criteria: Alfuzosin/dutasteride/finasteride 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

BRONCHODILATORS, BETA AGONIST

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INHALERS -	Short Acting	Non-preferred agents will be
PROAIR HFA (albuterol) INHALERS -	albuterol HFA (generic for ProAir HFA, Proventil HFA, Ventolin HFA) levalbuterol HFA (generic for Xopenex HFA) PROAIR DIGIHALER (albuterol) PROAIR RESPICLICK (albuterol) PROVENTIL HFA (albuterol) - Long Acting	approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Xopenex®: Covered for cardiac diagnoses or side effect of tachycardia with albuterol product
SEREVENT (salmeterol)	ARCAPTA NEOHALER (indacaterol) STRIVERDI RESPIMAT (olodaterol)	
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)	
OF	RAL	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

albuterol TABLET albuterol ER (generic for Vospire ER) metaproterenol (formerly generic for Alupent) terbutaline (generic for Brethine)

CALCIUM CHANNEL BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SHORT-ACTING		Non-preferred agents will be
Dihydro	pyridines	approved for patients who have failed a trial of ONE preferred
	isradipine (generic Dynacirc)	agent within this drug class
	nicardipine (generic Cardene)	s s
	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	pyridines	
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)	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BETA LACTAM/BETA-LACTAM	ASE INHIBITOR COMBINATIONS	
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
CEPHALOSPORIN	S – 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) Nyvepria (pegfilgrastim-apgf) ^{NR} ZARXIO (filgrastim-sndz) ZIEXTENZO SYR (pegfilgrastim-bmez)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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 Specific agents can be looked up using the Drug Look-up Tool at: https://druglookup.fhsc.com/druglookupweb/?client=nestate	charlotte 24 fe (norethindrone	

COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

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

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

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

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

Page **31** of **93**

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

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}	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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

CYTOKINE & CAM ANTAGONISTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ENBREL (etanercept) KIT, MINI CART, PENQL HUMIRA (adalimumab) QL ENBREL (entanercept) VIAL QL DTEZLA (apremilast) ORAL CL, QL	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIMZIA (certolizumab pegol) ^{QL} COSENTYX (secukinumab) ^{EL} ENSPRYNG (satralizumab-mwge) SUB-Q 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} XELJANZ XR (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: Requires documentation of inadequate response or intolerance to methotrexate Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to methotrexate. Diagnosis of Juvenile Idiopathic Arthritis for ages 2 years old and older does not require documentation of treatment failure with methotrexate. Diagnosis of moderate to severe ulcerative colitis (UC) requires documentation of treatment failure with a Tumor Necrosis Factor blocker agent; does not require documentation of treatment failure with methotrexate.

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

DIURETICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SINGLE-AGEN	IT PRODUCTS	Non-preferred agents will be
amiloride TABLET bumetanide TABLET chlorothiazide TABLET chlorthalidone TABLET (generic Diuril) furosemide SOLUTION, TABLET (generic Lasix) hydrochlorothiazide CAPSULE, TABLET (generic Microzide) indapamide TABLET metolazone TABLET spironolactone TABLET (generic Aldactone) torsemide TABLET	CAROSPIR (spironolactone) SUSPENSION eplerenone TABLET (generic Inspra) ethacrynic acid CAPSULE (generic Edecrin) 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
epinephrine (AUTHORIZED GENERIC for Epipen/ Epipen Jr.) AUTOINJECTOR	epinephrine (generic for Adrenaclick) epinephrine (generic for Epipen/ Epipen Jr.) AUTOINJECTOR EPIPEN (epinephrine) AUTOINJ EPIPEN JR. (epinephrine) AUTOINJ SYMJEPI (epinephrine) PFS	Non-preferred agents require clinical documentation why the preferred product within this drug class is not appropriate Brand name product may be authorized in event of documented national shortage of generic product.

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

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

Page **34** of **93**

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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
ciprofloxacin TABLET (generic Cipro) 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 Drug-specific criteria: Baxdela: Coverable with documented intolerance or failure of preferred MRSA agents (clindamycin, doxycycline, linezolic 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 (nongonorrhea)

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) Iubiprostone (generic Amitiza) CAPSULE AL, NR, QL 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 Lotronex®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate Relistor®: Covered for diagnosis of opioid-induced constipation in adults with chronic, non-cancer pain after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.) and failure of Movantik Symproic: Covered for diagnosis of opioid-induced constipation in adult patients with chronic non-cancer pain after trial on at least TWO OTC laxatives and failure of Movantik Trulance®: Covered for diagnosis of either chronic idiopathic constipation or IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.) Viberzi®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate

GLUCAGON AGENTSQL

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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) GLUCOCORTICOID/BRONCH ADVAIR DISKUS (fluticasone/ salmeterol) QL ADVAIR HFA (fluticasone/salmeterol) QL DULERA (mometasone/formoterol) SYMBICORT (budesonide/ formoterol)	AEROSPAN (flunisolide) ALVESCO (ciclesonide) ^{AL,CL} ARMONAIR DIGIHALER (fluticasone) ^{AL,NR,QL} ARMONAIR RESPICLICK (fluticasone) ^{AL} ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) ^{CL,AL,QL} FLOVENT DISKUS (fluticasone) QVAR (beclomethasone) QVAR Redihaler (beclomethasone) IODILATOR COMBINATIONS AIRDUO DIGIHALER (fluticasone/salmeterol) ^{AL,QL} BREO ELLIPTA (fluticasone/vilanterol) BREZTRI (budesonide/formoterol/glycopyrrolate) ^{QL} Budesonide/formoterol (generic for Symbicort) fluticasone/salmeterol (generic for Advair Diskus) ^{QL} fluticasone/salmeterol) TRELEGY ELLIPTA (fluticasone/umeclidinium/vilanterol) WIXELA INHUB (generic for Advair Diskus) ^{QL}	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 failed a trial of two preferred agents within this drug class, within the last 6 months.
INHALATION	SOLUTION	
	budesonide RESPULES (generic for Pulmicort)	

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

GLUCOCORTICOIDS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
budesonide EC CAPSULE (generic for Entocort EC) dexamethasone TABLET dexamethasone ELIXIR, SOLN hydrocortisone TABLET methylprednisolone tablet (generic for Medrol) prednisolone SOLUTION prednisolone sodium phosphate prednisone DOSE PAK prednisone TABLET	ALKINDI (hydrocortisone) GRANULES ^{AL/NR} CORTEF (hydrocortisone) cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSPENSION, TABLET ^{CL} ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg, 32mg ORTIKOS ER (budesonide) ^{AL,QL} PEDIAPRED (prednisolone sodium phosphate) prednisolone sodium phosphate (generic for Millipred/Veripred) prednisone SOLUTION prednisone INTENSOL RAYOS DR (prednisone) TABLET	 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

GROWTH HORMONES

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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 ORLADEYO (berotralstat) CAPAL, NR,QL 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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HEMOPHILIA TREATMENTS

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

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

CL – Prior Authorization / Class Criteria apply

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HEPATITIS B TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
entecavir TABLET lamivudine 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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HEPATITIS C TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIRECT ACTIN	IG ANTI-VIRAL	Hepatitis C Treatments PA Form
MAVYRET (glecaprevir/pibrentasvir) ^{CL} VOSEVI (sofosbuvir/velpatasvir/ voxilaprev) ^{CL}	DAKLINZA (daclatasvir) CL HARVONI 200/45MG, TABLET,	Non-preferred products require 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 ribavirin
RIBA	VIRIN	Harvoni:
	REBETOL (ribavirin)	For genotype 1 with decompensated cirrhosis along with ribavirin Post liver transplant for genotype 1 or 4 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 nonresponse after previous treatment course of Direct Acting Anti-viral agent (DAA) for genotype 1-6 without cirrhosis or with compensated cirrhosis

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HISTAMINE II RECEPTOR BLOCKERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
famotidine TABLET (generic for Pepcid) nizatidine SOLUTION (generic for Axid)	cimetidine TABLET, SOLUTION ^{CL} (generic for Tagamet) famotidine SUSPENSION nizatidine CAP (generic for Axid) ranitidine CAPSULE, (generic for Zantac) ranitidine OTC, SYRUP, TABLET (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 M. contagiosum or common wart V. Vulgaris treatment Famotidine susp/cimetidine solution: Requires clinical reason why nizatidine solution cannot be used

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HIV / AIDSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CCR5 AN	ITAGONISTS	 Non-preferred agents will be
SELZENTRY SOLN, TAB (maraviroc)		approved for patients who hav diagnosis of HIV/AIDS and particular approach
FUSION INHIBITORS		specific documentation of why the preferred products within this drug
FUZEON SUB-Q (enfuvirtide) ^{QL}		class are not appropriate for
HIV-1 ATTACK	MENT INHIBITOR	patient, including, but not limit to, drug resistance or concomi
SENTRESS (raltegravir) ^{QL}	TIVICAY PD (dolutegravir) ^{NR}	conditions not recommended very preferred agents
SENTRESS HD (raltegravir)		Patients undergoing treatment
ΓΙVICAY (dolutegravir)		the time of any preferred statu change will be allowed to cont
NON-NUCLEOSIDE REVERSE TR	ANSCRIPTASE INHIBITORS (NNRTIS)	therapy
EDURANT (rilpivirine)	efavirenz (generic Sustiva)	Diagnosis of HIV/AIDS require OR
NTELENCE (etravirine) ^{QL}	nevirapine IR, ER (generic	Pre and Post Exposure
PIFELTRO (doravirine) ^{QL}	Viramune/Viramune XR)	Prophylaxis
SUSTIVA CAPSULE, TABLET	RESCRIPTOR (delavirdine)	
(efavirenz)	VIRAMUNE (nevirapine) SUSP	
NUCL FOSIDE REVERSE TRA	NSCRIPTASE INHIBITORS (NRTIs)	
	NSCRIPTASE INHIBITORS (NRTIs) didanosine DR (generic Videx EC)	
abacavir SOLN, TABLET (generic Ziagen)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR}	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir) NUCLEOTIDE REVERSE TRA	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir) NUCLEOTIDE REVERSE TRA	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) amivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir) NUCLEOTIDE REVERSE TRA tenofovir TABLET (generic Viread)	didanosine DR (generic Videx EC) emtricitabine CAPSULE (generic for Emtriva) ^{NR} EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HIV / AIDS^{CL} (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROTEASE	INHIBITORS	
atazanavir CAPSULE (generic Reyataz)	APTIVUS CAPSULE , SOLN	
LEXIVA SUSP, TABLET	(tipranavir)	
(fosamprenavir)	CRIXIVAN (indinavir)	
NORVIR (ritonavir) TAB	fosamprenavir TAB (generic Lexiva)	
PREZISTA (darunavir) SUSP, TABLET	INVIRASE (saquinavir)	
	NORVIR POWDER , SOLN (ritonavir)	
	REYATAZ POWDER (atazanavir)	
	ritonavir TABLET (generic Norvir)	
	VIRACEPT (nelfinavir)	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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 TAB (lopinavir/ritonavir) PREZCOBIX (darunavir/cobicistat) ^{QL} lopinavir/ritonavir SOLN (generic Kaletra)	KALETRA SOLN (lopinavir/ritonavir)	
COMBINATION NUCLEOS(T)IDE RI	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) emtricitabine/tenofovir (generic Truvada) ^{CL,NR} EPZICOM (abacavir sulfate/lamivudine) TEMIXYS (lamivudine/tenofovir) ^{QL} TRIZIVIR (abacavir/lamivudine/zidovudine)	
COMBINATION PRODU	CTS - MULTIPLE CLASSES	
ATRIPLA (tenofovir/emtricitabine/efavirenz) BIKTARVY (bictegravir/emtricitabine/tenofovir) COMPLERA (rilpivirine/emtricitabine/tenofovir) DELSTRIGO (doravirine/lamivudine/tenofovir) GENVOYA (elvitegravier/cobicistat/emtricitabine/tenofovir) ODEFSEY (emtricitabine/rilpivirine/tenofovir) STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir) STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir) SYMFI (efavirenz/lamivudine/tenofovir) CSYMFI LO (efavirenz/lamivudine/tenofovir) CSYMFI LO (efavirenz/lamivudine/tenofovir) CSYMFI LO (dolutegravir/abacavir/lamivudine)	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}	

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

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

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 PEN (exenatide ER) subcutaneous BYETTA (exenatide) subcutaneous VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) BYDUREON BCISE PEN (exenatide) ^{QL} OZEMPIC (semaglutide) RYBELSUS (semaglutide) TANZEUM (albiglutide) TRULICITY (dulaglutide)	trial and diagnosis of diabetes Non-preferred agents will be approved for patients who have: Failed a trial of TWO preferred agents within GLP-1 RA AND Diagnosis of diabetes with HbA1C ≥ 7 AND
INSULIN/GLP-1 R	A COMBINATIONS	Trial of metformin, or contraindication or intolerance to
	SOLIQUA (insulin glargine/lixisenatide) XULTOPHY (insulin degludec/liraglutide)	metformin
AMYLIN	ANALOG	ALL criteria must be met
DIDEDTIDVI DEDTIDAS	SYMLIN (pramlintide) subcutaneous	 Concurrent use of short-acting mealtime insulin Current therapy compliance No diagnosis of gastroparesis HbA1C ≤ 9% within last 90 days Fingerstick monitoring of glucose during initiation of therapy
DIPERTIDIL PERTIDAS	E-4 (DPP-4) INHIBITOR ^{QL}	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

Preferred Agents Prior Authorization/Class Criteria **Non-Preferred Agents** Non-preferred agents will be HUMALOG (insulin lispro) U-100 ADMELOG (insulin lispro) PEN, VIAL approved for patients who have CARTRIDGE, PEN, VIAL AFREZZA (regular insulin) failed a trial of ONE preferred HUMALOG JR. (insulin lispro) U-100 **INHALATION** agent within this drug class **PEN** APIDRA (insulin alulisine) **HUMALOG MIX VIAL** (insulin BASAGLAR (insulin glargine, rec) Drug-specific criteria: lispro/lispro protamine) **PEN** Afrezza®: Approved for T1DM on HUMALOG MIX PEN (insulin long-acting insulin with no current FIASP (insulin aspart) CARTRIDGE, lispro/lispro protamine) history of smoking or chronic lung PEN, VIAL disease HUMULIN (insulin) VIAL HUMALOG (insulin lispro) U-200 PEN Humulin® R U-500 Kwikpen: **HUMULIN 70/30 VIAL** insulin lispro (generic for Humalog) Approved for physical reasons -**HUMULIN U-500 VIAL** PEN, VIAL such as dexterity problems and vision impairment HUMULIN R U-500 KWIKPEN^{CL} insulin aspart (generic for Novolog) Usage must be for self-LYUMJEV KWIKPEN, VIAL(insulin **HUMULIN OTC PEN** administration, not only lispro-aabc)NR **HUMULIN 70/30 OTC PEN** convenience NOVOLIN (insulin) LANTUS SOLOSTAR PEN (insulin Patient requires >200 units/day NOVOLIN 70/30 VIAL(insulin) glargine) Safety reason patient can't use TOUJEO SOLOSTAR (insulin LANTUS (insulin glargine) VIAL vial/syringe glargine) LEVEMIR (insulin detemir) PEN, VIAL SEMGLEE (insulin glargine)NR PEN, NOVOLOG (insulin aspart) VIAL CARTRIDGE, PEN, VIAL TRESIBA (insulin degludec) NOVOLOG MIX PEN, VIAL (insulin aspart/aspart protamine)

HYPOGLYCEMICS, MEGLITINIDES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria

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

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

QL – Quantity/Duration Limit

AL – Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

repaglinide (generic for Prandin)	nateglinide (generic for Starlix) repaglinide/metformin (generic for Prandimet)	•	Non-preferred agents will be approved for patients with: Failure of a trial of ONE preferred agent in another Hypoglycemic class OR T2DM and inadequate glycemic control
-----------------------------------	---	---	--

HYPOGLYCEMICS, METFORMINS

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

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}	 Preferred agents are Approved for diagnosis of diabetes AND a trial of metformin Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

HYPOGLYCEMICS, SULFONYLUREAS

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

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

HYPOGLYCEMICS, TZD

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

IDIOPATHIC PULMONARY FIBROSIS

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

IIWINONONODULATORS, ASTHINA		
Preferred Agents		Prior Authorization/Class Criteria

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

FASENRA (benralizumab) ^{AL} PEN	NUCALA (mepolizumab) ^{AL} AUTO-INJ, SYR,	Drug Specific Criteria: Dupixent: See criteria listed under Immunomodulator, Atopic Dermatitis class Fasenra: is indicated for patient 12 years and older for add on
		maintenance treatment of severe asthma, and with an eosinophilic phenotype
		Nucala: is indicated for -Patients 6 years and older for add on maintenance treatment of severe asthma, and with an eosinophilic phenotype
		-Patients 12 years and older with hypereosinophilic syndrome (HES) for ≥6 months without identifiable non-hematologic secondary cause
		-Adult patients with eosinophilic granulomatosis with polyangiitis

IMMUNOMODULATORS, ATOPIC DERMATITISAL

Preferred Agents Non-Preferred Agents Prior Authorization/Class Criteria

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

circuive January 21, 2021		
ELIDEL (pimecrolimus) EUCRISA (crisaborole) ^{CL,QL}	DUPIXENT PEN ^{AL} pimecrolimus (generic for Elidel) tacrolimus (generic for Protopic) ^{CL}	 Non-preferred agents require: Trial of a topical steroid AND Trial of one preferred product within this drug class Drug-specific criteria: Dupixent: Indicated for moderate to severe atopic dermatitis, must have trial of Eucrisa; For moderate to severe asthma, must have eosinophilic phenotype or oral corticosteroid dependent asthma uncontrolled with maintenance controller medication; For adults with chronic rhinosinusitis with nasal polyposis, must document inadequate control on current treatment regimen and be used as add-on maintenance treatment with intranasal steroid Eucrisa: Requires use and failure of 1 topical steroid or Elidel.

IMMUNOMODULATORS, TOPICAL

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

IMMUNOSUPPRESSIVES, ORAL

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

AL Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathiaprine (generic Imuran) cyclosporine, modified CAPSULE (generic Neoral) mycophenolate CAPSULE, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLUTION 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	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Patients established on existing therapy will be allowed to continue

INTRANASAL RHINITIS DRUGS

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

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

LEUKOTRIENE MODIFIERS

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

LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

LIPOTROPICS, OTHER

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

LIPOTROPICS, OTHER (continued)

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

LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atorvastatin (generic Lipitor) ^{QL} lovastatin (generic Mevacor) pravastatin (generic Pravachol) rosuvastatin (generic Crestor) simvastatin (generic Zocor)	ALTOPREV (lovastatin ER) ^{CL} EZALLOR SPRINKLE (rosuvastatin) ^{QL} fluvastatin IR/ER (generic Lescol/ Lescol XL) LIVALO (pitavastatin)	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 Drug-specific criteria:
, , , , , , , , , , , , , , , , , , ,	ZYPITAMAG (pitavastatin) //BINATIONS	 Altoprev®: One of the TWO trials must be IR lovastatin Combination products: Require clinical reason why individual ingredients cannot be

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

atorvastatin/amlodipine (generic Caduet) simvastatin/ezetimibe (generic Vytorin)	used fluvastatin ER: Requires trial of TWO preferred agents AND trial of IR fluvastatin OR clinical reason IR cannot be used simvastatin/ezetimibe: Approved for 3-month continuous trial of ONE standard dose statin
---	---

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

MACROLIDES AND KETOLIDES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MACROLIDES		Require clinical reason why
azithromycin (generic Zithromax) clarithromycin TABLET, SUSPENSION (generic Biaxin)	clarithromycin ER (generic Biaxin XL) E.E.S. SUSPENSION, TABLET (erythromycin ethylsuccinate) ERY-TAB (erythromycin) ERYPED SUSPENSION (erythromycin) ERYTHROCIN (erythromycin) erythromycin base TABLET, CAPSULE erythromycin ethylsuccinate SUSPENSION	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 PF VIAL, TABLET, VIAL	OTREXUP (methotrexate) SUB-Q RASUVO (methotrexate) SUB-Q TREXALL (methotrexate) TABLET XATMEP (methotrexate) SOLUTION	 Non-preferred agents will be approved for FDA-approved indications Drug-specific criteria: XatmepTM:Indicated for pediatric patients only

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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 XENAZINE (tetrabenazine) ^{CL}	Non-preferred agent requires trial of Austedo All drugs require an FDA approved indication – ICD-10 diagnosis code required. Drug-specific criteria: • Austedo: Diagnosis of Tardive Dyskinesia or chorea associated with Huntington's Disease; Requires a Step through tetrabenazine with the diagnosis of chorea associated with Huntington's Disease • Ingrezza: Diagnosis of Tardive Dyskinesia in adults and trial of Austedo • tetrabenazine:Diagnosis of chorea with Huntington's Disease

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) BAFIERTAM (monomethyl fumarate) ^{NR,QL} dalfampridine (generic Ampyra) ^{QL} dimethyl fumarate (generic for Tecfidera) ^{NR} EXTAVIA (interferon beta-1b) ^{QL} glatiramer (generic Copaxone) ^{QL} KESIMPTA ((Ofatumumab) ^{NR,QL} MAVENCLAD (cladribine) MAYZENT (siponimod) ^{QL} PLEGRIDY (peginterferon beta-1a) ^{QL} REBIF (interferon beta-1a) ^{QL} VUMERITY (diroximel) ^{QL} ZEPOSIA (ozanimod) ^{AL,NR,QL}	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Ampyra®: Approved for diagnosis of gait disorder associated with MS AND EDSS score ≤ 7 Plegridy: Approved for diagnosis of relapsing MS

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

NITROFURAN DERIVATIVES

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

NSAIDs, ORAL

Preferred Agents	Non-Preferred Agents		Prior Authorization/Class Criteria
COX-I SELECTIVE		•	Non-preferred agents within COX-

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

diclofenac sodium (generic for Voltaren)
ibuprofen OTC, Rx (generic for Advil, Motrin) CHEW, DROPS, SUSPENSION, TABLET
indomethacin CAPSULE (generic for Indocin)
ketorolac (generic for Toradol)
meloxicam TABLET (generic for Mobic)
nabumetone (generic for Relafen)
naproxen Rx, OTC (generic for Naprosyn)
naproxen enteric coated

sulindac (generic for Clinoril)

diclofenac potassium (generic for Cataflam, Zipsor) diclofenac SR (generic for Voltaren-XR) diflunisal (generic for Dolobid) etodolac & SR (generic for Lodine/XL) fenoprofen (generic for Nalfon) flurbiprofen (generic for Ansaid) ibuprofen OTC (generic for Advil, Motrin) CAPSULE indomethacin ER (generic for Indocin) INDOCIN RECTAL, SUSPENSION ketoprofen & ER (generic for Orudis) meclofenamate (generic for Meclomen) mefenamic acid (generic for Ponstel) meloxicam CAP (generic Vivlodex)CL, NR,QL naproxen CR (generic for Naprelan) naproxen SUSPENSION (generic for Naprosyn) naproxen sodium (generic for

naproxen-esomeprazole (generic for

oxaprozin (generic for Daypro) piroxicam (generic for Feldene) QMIIZ ODT (meloxicam) QL RELAFEN DS (nabumetone) tolmetin (generic for Tolectin)

Anaprox)

Vimovo)

1 SELECTIVE group will be approved for patients who have failed no less than 30-day trial of TWO preferred agents within this drug class

Drug-specific criteria:

- Arthrotec®: Requires clinical reason why individual ingredients cannot be used
- Duexis®/Vimovo®: Requires clinical reason why individual agents cannot be used
- meclofenamate: Approvable without trial of preferred agents for menorrhagia

NSAIDs, ORAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COX-I SELECTIVE (continued)		

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

effective January 21, 2021			
NSAID/GI PROTECT/	ALL BRAND NAME NSAIDs including: CAMBIA (diclofenac oral solution) DUEXIS (ibuprofen/famotidine) SPRIX (ketorolac nasal spray) NASALQL, CL TIVORBEX (indomethacin) VIVLODEX (meloxicam submicronized) ZIPSOR (diclofenac) ZORVOLEX (diclofenac)	Drug-specific criteria: Sprix®: Approved for patients unable to tolerate, swallow OR absorb oral NSAIDs OR contraindication OR trial of TWO preferred oral NSAIDs Tivorbex®: Requires clinical reason why indomethacin capsules cannot be used Zorvolex®: Requires trial of oral diclofenac OR clinical reason why diclofenac potassium/sodium cannot be used	
	diclofenac/misoprostol (generic for Arthrotec)	•	
COX-II SE celecoxib (generic for Celebrex)	ELECTIVE		

NSAIDs, TOPICAL

Preferred Agents Non-Preferred Agents Prior Authorization/Class Criteria

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

diclofenac sodium GEL (OTC only)	diclofenac (generic for Pennsaid Solution) ^{CL} FLECTOR PATCH (diclofenac) ^{CL} <i>LICART PATCH</i> (diclofenac) ^{CL} PENNSAID PACKET , PUMP (diclofenac) ^{CL} VOLTAREN GEL (diclofenac) ^{CL}	 Flector®/Licart: Approved for diagnosis of acute pain due to sprain/strain/contusion AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form Pennsaid®: Approved for osteoarthritis of the knees AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form Pennsaid® Pump: Requires clinical reason why 1.5% solution cannot be used Voltaren®: Approved for diagnosis of osteoarthritis AND trial of oral diclofenac OR clinical resaon patient cannot use oral dosage form
----------------------------------	--	---

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

ONCOLOGY AGENTS, ORAL, BREAST

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

ONCOLOGY AGENTS, ORAL, HEMATOLOGIC

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
mercaptopurine A	LL PURIXAN (mercaptopurine)	 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
Α	ML	from current treatment guidelines
IMBRUVICA (irutinib)	DAURISMO (glasdegib maleate) ^{QL} IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) ^{QL} XOSPATA (gilteritinib) ^{QL} ELL COPIKTRA (duvelisib) ^{QL}	 Drug-specific critera Hydrea®: Requires clinical reason why generic cannot be used melphalan: Requires trial of Alkeran or clinical reason Alkeran cannot be used Tabloid: Prior authorization not required for age <19
LEUKERAN (chlorambucil) VENCLEXTA (venetoclax)	ZYDELIG (idelalisib)	 Tasigna: Patients receiving Tasigna, which changed from preferred to non-preferred on 1-17- 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}	 therapy Xpovio: Indicated for relapsed or refractory multiple myeloma. Requires concomitant therapy with dexamethasone
M	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 ^{QL} CALQUENCE (acalabrutinib) ^{QL} INREBIC (fedratinib dihydrochloride) ^{QL} INQOVI (decitabine/cedazuridine) ZOLINZA (vorinostat)	

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

ONCOLOGY AGENTS, ORAL, LUNG

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALK		Non-preferred agents DO NOT require a trial of a preferred agent
ALECENSA (alectinib)	ALUNBRIG (brigatinib) LORBRENA (lorlatinib) QL ZYKADIA (ceritinib) CAPSULE, TABLET	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 _Drug-Specific Criteria
ALK / ROS1 / NTRK		 Iressa/ Xalkori: Patients receiving Iressa or Xalkori prior to 1/21/21 (which changed from preferred to non-preferred) will be allowed to continue current treatment
	ROZLYTREK (entrectinib) AL,QL XALKORI (crizotinib)	
EC	GFR SFR	
TAGRISSO (osimertinib) OT	erlotinib (generic for Tarceva) GILOTRIF (afatinib) IRESSA (gefitinib) TARCEVA (erlotinib) VIZIMPRO (dacomitinib) ^{QL} HER GAVRETO (pralsetinib) ^{QL} HYCAMTIN (topotecan) RETEVMO (selpercatinib) ^{AL} TABRECTA (capmatinib) ^{QL}	

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ONCOLOGY AGENTS, ORAL, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CAPRELSA (vandetanib) GLEOSTINE (lomustine) LYNPARZA (olaparib) temozolomide (generic for Temodar) ZEJULA (niraparib)	BALVERSA (erdafitinib) COMETRIQ (cabozantinib) HEXALEN (altretamine) KOSELUGO (selumetinib) ^{AL} LONSURF (trifluridine/tipiracil) PEMAZYRE (pemigatinib) ^{QL} RUBRACA (rucaparib) STIVARGA (regorafenib) TAZVERIK (tazemetostat) ^{AL} TURALIO (pexidartinib) ^{QL} VITRAKVI (larotrectinib) CAPSULE, SOLUTION ^{QL}	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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

ONCOLOGY AGENTS, ORAL, PROSTATE

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

ONCOLOGY AGENTS, ORAL, RENAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INLYTA (axitinib) LENVIMA (lenvatinib) SUTENT (sunitinib) VOTRIENT (pazopanib	AFINITOR DISPERZ (everolimus)CL CABOMETYX (cabozantinib) everolimus (generic for Afinitor) NEXAVAR (sorafenib)	 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 Drug-specific critera Afinitor: Patients receiving Afinitor, which changed from preferred to non-preferred on 1-17-19 will be allowed to continue therapy

ONCOLOGY AGENTS, ORAL, SKIN

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ERIVEDGE (vismodegib)	ODOMZO (sonidegib) ^{CL}	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
BRAF MUTATION		

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

Page **69** of **93**

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

MEKINIST (trametinib) TAFINLAR (dabrafenib)	BRAFTOVI (encorafenib) COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)	 Drug-specific critera Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17- 19 will be allowed to continue therapy
---	--	---

OPHTHALMICS, ALLERGIC CONJUNCTIVITIS

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

OPHTHALMICS, ANTIBIOTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FLUOROQUINOLONES		 Non-preferred agents will be
ciprofloxacin SOLUTION (generic for Ciloxan) ofloxacin (generic for Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic for Zymaxid) levofloxacin MOXEZA (moxifloxacin) moxifloxacin (generic for Vigamox) moxifloxacin (generic for Moxeza) VIGAMOX (moxifloxacin)	 approved for patients who have failed a one-month trial of TWO preferred agent within this drug class Azasite®: Approval only requires trial of erythromycin Drug-specific criteria: Natacyn®: Approved for documented fungal infection
MACROLIDES		3
erythromycin	AZASITE (azithromycin) ^{CL}	
AMINOGLYCOSIDES		
gentamicin OINTMENT	TOBREX OINTMENT (tobramycin)	
gentamicin SOLUTION		
tobramycin (generic for Tobrex drops)		
OTHER OPHTH	ALMIC AGENTS	
bacitracin/polymyxin B (generic	bacitracin	
Polysporin)	NATACYN (natamycin) ^{CL}	
polymyxin B/trimethoprim (generic for Polytrim)	neomycin/bacitracin/polymyxin B OINTMENT	
	neomycin/polymyxin B/gramicidin	
	NEOSPORIN (neomycin/polymyxin B/gramcidin)	
	sulfacetamide SOLUTION (generic for Bleph-10)	
	sulfacetamide OINTMENT	

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

OPHTHALMICS, ANTIBIOTIC-STEROID COMBINATIONS

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

OPHTHALMICS, ANTI-INFLAMMATORIES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CORTICO	STEROIDS	Non-preferred agents will be
fluorometholone 0.1% (generic for FML) OINTMENT LOTEMAX SOLUTION (loteprednol 0.5%) MAXIDEX (dexamethasone) PRED MILD (prednisolone 0.12%)	dexamethasone (generic for Maxidex) DUREZOL (difluprednate) FLAREX (fluorometholone) FML (fluorometholone 0.1% SOLUT.) FML FORTE (fluorometholone 0.25%) FML S.O.P. (fluorometholone 0.1%) INVELTYS(loteprednol etabonate) LOTEMAX OINTMENT, GEL (loteprednol) loteprednol 0.5% SOLUTION (generic for Lotemax SOLUTION) prednisolone acetate 1% (gen. for Omnipred, Pred Forte) prednisolone sodium phosphate	 approved for patients who have failed a trial of TWO preferred agents within this drug class NSAID class: Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent
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
------------------	----------------------	------------------------------------

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

CL - QL - Quantity/Duration Limit AL - Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

RESTASIS (cyclosporine)
RESTASIS MULTIDOSE
(cyclosporine)
XIIDRA (lifitegrast)

CEQUA (cyclosporine) QL EYSUVIS (loteprednol etabonate)NR,QL Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

OPHTHALMICS, GLAUCOMA

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIO	TICS	Non-preferred agents will be
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide)	approved for patients who have failed a trial of ONE preferred agent within this drug class
SYMPATHO	MIMETICS	· ·
brimonidine 0.2% (generic for	Alphagan P (brimonidine 0.1%)	
Alphagan)	Alphagan P (brimonidine 0.15%)	
	apraclonidine (generic for lopidine)	
BETA BLO	OCKERS	
levobunolol (generic for Betagan)	betaxolol (generic for Betoptic)	
timolol (generic for Timoptic)	BETIMOL (timolol)	
	BETOPTIC S (betaxolol)	
	carteolol (generic for Ocupress)	
	timolol (generic for Istalol)	
	timolol (generic for Timoptic	
	Ocudose) ^{NR}	
	TIMOPTIC OCUDOSE	
	TIMOPTIC XE (timolol gel forming	
OARRONIO ANILIVEI	solution)	-
CARBONIC ANHYDI	KASE INHIBITORS	
dorzolamide (generic for Trusopt)	AZOPT (brinzolamide)	
PROSTAGLAND	OIN ANALOGS	

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

CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL – Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATIO	ON DRUGS	
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine	
ОТН	IER	•
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

OPIOID DEPENDENCE TREATMENTS

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

OPIOID-REVERSAL TREATMENTS

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

CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

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

OTIC ANTIBIOTICS

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

PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria

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

CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL – Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ADCIRCA (tadalafil)CL ambrisentan (generic Letairis) sildenafil TABLET (generic Revatio)CL TRACLEER TABLET (bosentan) TYVASO **INHALATION** (treprostinil) VENTAVIS **INHALATION** (iloprost)

ADEMPAS (riociquat)CL bosentan TABLET (generic Tracleer) Revatio)CL tadalafil (generic for Adcirca)CL

LETAIRIS (ambrisentan) **OPSUMIT** (macitentan) **ORENITRAM ER (treprostinil)** sildenafil SUSPENSION (generic

TRACLEER TABLETS FOR **SUSPENSION** (bosentan)

UPTRAVI (selexipag)

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

Drug-specific criteria:

- Adcirca®/Revatio®: Approved for diagnosis of Pulmonary Arterial Hypertension (PAH)
- Adempas®:

PAH: Requires clinical reason preferred agent cannot be used CTEPH: Approved for persistent/recurrent diagnosis after surgical treatment or inoperable CTĔPH

NOT for use in Pregnancy

sildenafil suspension: Requires clinical reason why sildenafil tablets cannot be used

PANCREATIC ENZYMES

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

PEDIATRIC VITAMIN PREPARATIONS

•				
	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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)

no.164/ferrous sulfate drops)

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

multivitamins with fluoride (pedi multivit 2/fluoride) DROPS

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

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

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

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

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

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

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

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

AQUADEKS (pedi multivit 40/phytonadione)

ESCAVITE (pedi multivit 47/iron/fluoride)

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

ESCAVITE LQ (pedi multivit 86/iron/fluoride)

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

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

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

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

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

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

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

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

QUFLORA FE (pedi multivit 142/iron/fluoride)

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

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

Drug specific criteria:

Aquadeks: Approved for diagnosis of Cystic Fibrosis

PENICILLINS

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

PHOSPHATE BINDERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcium acetate TABLET , CAPSULE CALPHRON OTC (calcium acetate) sevelamer carbonate (generic Renvela)	AURYXIA (ferric citrate) ELIPHOS (calcium acetate) lanthanum (generic FOSRENOL) PHOSLO (calcium acetate) PHOSLYRA (calcium acetate) RENAGEL (sevelamer HCI) sevelamer HCI (generic Renagel) VELPHORO (sucroferric oxyhydroxide)	 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

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}	 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 Zontivity®: Approved for reduction of thrombotic cardiovascular events in history of MI or with peripheral artery disease (PAD) Use with aspirin and/or clopidogrel

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

PRENATAL VITAMINS

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
c-nate dha SOFTGEL complete natal dha (pnv2/iron b-g suc-p/fa/omega-3) calcium-pnv 28-1-250mg SOFTGEL classic prenatal TABLET (prenatal vit/fe fum/fa) COMPLETENATE CHEWABLE CONCEPT DHA CAPSULE CONCEPT DHA CAPSULE elite-ob CAPLET (fe c/fa) MARNATAL-F CAPSULE PRENATA TAB CHEW pnv with ca, #72/iron/fa pnv-ob+dha combo pack (pnv22/iron cbn&gluc/fa/dss/dha) pnv-vp-u CAPSULE prenaissance CAPSULE (pnv80/iron fum/fa/dss/dha) prenaissance plus SOFTGEL (pnv69/iron/fa/dss/dha) prenatal vitamin TABLET (pnv#124/iron/fa) prenatal no.137/iron/fa OTC pretab 29mg-1 TABLET (pnv#78/iron/fa) PUREFE PLUS PUREFE OB PLUS TARON-PREX PRENATAL TRINATAL RX 1 triveen-duo dha combo pack (pnv53/iron b-g hcl-p/fa/omega-3) virtprex CAPSULE (pnv66/iron fum/fa/dss/dha) virt-nate dha SOFTGEL (pnv 11-iron fum-fa-om3) virt-pn TABLET (pnv w-ca no.40/iron fum/fa cmb no.1) virt-pn plus SOFTGEL (pnv/ca no.68/iron/fa1/dha) virt-select CAPSULE (pnv80/iron fum/fa/dss/dha) virt-vite gt TABLET (prenatal vit 16/iron cb/fa/dss) VOL-PLUS TABLET vp-ch-pnv prenatal SOFTGEL vp-heme ob TABLET (pnv#21/iron/ps& heme polyp/fa) zatean-pn plus SOFTGEL (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 dha SOFTGEL (pnv combo#47/iron/fa #1/dha) WESTGEL DHA (PRENATAL 93/IRON/FOLATE 9/DHA) ^{VR} zatean-pn dha CAPSULE (pnv #47/iron/fa #1/dha)	 Non-preferred agents will be approved for patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

PROGESTERONE (hydroxyprogesterone caproate)

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

PROTON PUMP INHIBITORS

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

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

SEDATIVE HYPNOTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
temazepam 15mg, 30mg (generic for Restoril)	estazolam (generic for ProSom) flurazepam (generic for Dalmane) temazepam (generic for Restoril) 7.5mg, 22.5mg triazolam (generic for Halcion) ERS BELSOMRA (suvorexant) ^{AL,QL} DAYVIGO (lemborexant) ^{AL,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) zolpidem SL (generic for Intermezzo)	 Lunesta®/ Rozerem®/zolpidem ER: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used Edluar®: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used and Requires documentation of swallowing disorder flurazepam/triazolam: Requires trial of preferred benzodiazepine Hetlioz®: Requires trial with generic zolpidem within last 12 months AND clinical reason why zaleplon AND preferred benzodiazepine cannot be used Silenor®: Must meet ONE of the following: 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 ER® 6.25mg zolpidem SL: Requires clinical reason why half of zolpidem tablet cannot be used

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

SICKLE CELL ANEMIA TREATMENTAL

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

SINUS NODE INHIBITORS

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

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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 TABLET (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 CAPSULE ZANAFLEX (tizanidine) CAPSULE, TABLET	 Non-preferred agents will be approved for patients who have failed a 1-week trial of TWO preferred agents within this drug class Drug-specific criteria: cyclobenzaprine ER: Requires clinical reason why IR cannot be used Approved only for acute muscle spasms NOT approved for chronic use carisoprodol: Approved for Acute, musculoskeletal pain - NOT for chronic pain Use is limited to no more than 30 days Additional authorizations will not be granted for at least 6 months following the last dayy of previous course of therapy Dantrolene: Trial NOT required for treatment of spasticity from spinal cord injury Lorzone®: Requires clinical reason why chlorzoxazone cannot be used Soma® 250mg: Requires clinical reason why 350mg generic strength cannot be used Zanaflex® Capsules: Requires clinical reason generic cannot be used

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

STEROIDS TOPICAL

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

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

CL – Prior Authorization / Class Criteria apply

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

STEROIDS, TOPICAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HIGH POTENCY		 High Potency Non-preferred
triamcinolone acetonide OINTMENT, CREAM	amcinonide CREAM, LOTION, OINTMENT	agents will be approved for patients who have failed a trial of TWO preferred agents within this
triamcinolone LOTION	betamethasone dipropionate betamethasone / propylene glycol	drug class
	betamethasone valerate	
	desoximetasone	
	diflorasone diacetate fluocinonide SOLUTION	
	fluocinonide CREAM, GEL, OINTMENT	
	fluocinonide emollient	
	halcinonide CREAM (generic for Halog)	
	HALOG (halcinonide) CREAM, OINT, SOLN	
	KENALOG AEROSOL (triamcinolone)	
	SERNIVO (betamethasone dipropionate)	
	triamcinolone SPRAY (generic for Kenalog spray)	
	TRIANEX OINTMENT (triamcinolone)	
	VANOS (fluocinonide)	
VERY HIG	H POTENCY	 Very High Potency Non-preferred

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

clobetasol emollient (generic for Temovate-E) clobetasol propionate CREAM, GEL, OINTMENT, SOLUTION halobetasol propionate (generic for Ultravate) APEXICON-E (diflorasone)
BRYHALI (halobetasol prop) LOTION
clobetasol SHAMPOO, LOTION
clobetasol propionate FOAM, SPRAY
CLOBEX (clobetasol)
halobetasol propionate FOAM (generic for Lexette) AL,QL
IMPEKLO (clobetasol) LOTIONAL,NR
LEXETTE(halobetasol propionate) AL,QL
OLUX-E /OLUX/OLUX-E CP

(clobetasol)

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

STIMULANTS AND RELATED AGENTS^{AL}

Preferred Agents	Non-Preferred Agents	Pri	or Authorization/Class Criteria
CNS STIMULANTS		Non	 Non-preferred agents will be approved for patients who have
Amphetamine type		appi	

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ADDERALL XR (amphetamine sa	alt
combo)	

amphetamine salt combination IR VYVANSE (lisdexamfetamine) CAPSULE, CHEWABLE

ADZENYS XR (amphetamine) amphetamine ER (generic for Adzenys ER) **SUSPENSION**

amphetamine salt combination ER (generic for Adderall XR)

amphetamine sulfate (generic for Evekeo)

dextroamphetamine (generic for Dexedrine)

dextroamphetamine **SOLUTION** (generic for Procentra)

dextroamphetamine ER (generic for Dexedrine ER)

DYANAVEL XR (amphetamine)

EVEKEO ODT (amphetamine sulfate)

MYDAYIS (amphetamine salt combo)^{QL}

methamphetamine (generic for Desoxyn)

ZENZEDI (dextroamphetamine)

failed a trial of ONE preferred agent within this drug class

Drug-specific criteria:

- Procentra®: May be approved with documentation of swallowing disorder
- Zenzedi[®]: Requires clinical reason generic dextroamphetamine IR cannot be used

STIMULANTS AND RELATED ADHD DRUGS (Continued)^{AL}

Preferred Agents Non-Preferred Agents Prior Authorization/Class Criteria

Methylphenidate type Non-preferred agents will be

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

Page **88** of **93**

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

CONCERTA (methylphenidate ER)^{QL}
18mg, 27mg, 36mg, 54mg
dexmethylphenidate (generic for
Focalin IR)
FOCALIN XR (dexmethylphenidate)
METHYLIN SOLUTION
(methylphenidate)
methylphenidate (generic for Ritalin)
methylphenidate SOLUTION (generic
for Methylin)
methylphenidate ER
(generic for Ritalin SR)
QUILLICHEW ER CHEWTAB
(methylphenidate)

ADHANSIA XR (methylphenidate) QL APTENSIO XR (methylphenidate) COTEMPLA XR-ODT (methylphenidate)QL DAYTRANA PATCH (methylphenidate)QL dexmethylphenidate XR (generic for Focalin XR) FOCALIN IR (dexmethylphenidate) JORNAY PM (methylphenidate) QL methylphenidate 50/50 (generic for Ritalin methylphenidate 30/70 (generic for Metadate CD) methylphenidate ER 18mg, 27mg, 36mg, 54mg (generic Concerta)QL methylphenidate ER CAP (generic for Aptensio XR)QL Methylphenidate ER (generic for Metadate ER) methylphenidate ER 72mg (generic for RELEXXII)QL

methylphenidate ER (generic for Ritalin

SR)

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

Drug-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

STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents Non-Preferred Agents

Prior Authorization/Class Criteria

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

atomoxetine (generic for Strattera) ^{QL} guanfacine ER (generic for Intuniv) ^{QL} STRATTERA (atomoxetine) ANALEPTICS armodafinil (generic for Nuvigil) ^{CL} modafanil (generic for Provigil) ^{CL} SUNOSI (solriamfetol) CL,QL WAKIX (pitolisant) ^{CL,QL} WAKIX (pitolisant) ^{CL,QL} Clonidine IR are available without prior authorization Drug-specific criteria: armodafinil and Sunosi: Requitial of modafinil armodafinil armodafinil approved only for: Sleep Apnea with documentation/confirm via sleep study and documentation that C-l has been maxed Narcolepsy with	MISCELLANEOUS Note: generic guanfacine IR an				
atomoxetine (generic for Strattera) ^{QL} guanfacine ER (generic for Intuniv) ^{QL} STRATTERA (atomoxetine) Drug-specific criteria: armodafinil and Sunosi: Requiral of modafinil and modafinil approved only for: o Sleep Apnea with documentation/confirm via sleep study and documentation that C-l has been maxed Narcolepsy with	MISCELLANEOUS		dine IR are available without		
ANALEPTICS armodafinil (generic for Nuvigil) ^{CL} modafanil (generic for Provigil) ^{CL} SUNOSI (solriamfetol) ^{CL,QL} WAKIX (pitolisant) ^{CL,QL} WAKIX (pitolisant) ^{CL,QL} armodafinil and Sunosi: Requitival of modafinil and modafinil: approved only for: Sleep Apnea with documentation/confirm via sleep study and documentation that C-lhas been maxed Narcolepsy with		(generic for Kapvay) ^{QL} prior			
armodafinil (generic for Nuvigil) ^{CL} modafanil (generic for Provigil) ^{CL} SUNOSI (solriamfetol) ^{CL,QL} WAKIX (pitolisant) ^{CL,QL} WAKIX (pitolisant) CL,QL WAKIX (pitolisa	ANALEDTICS	a	rmodafinil and Sunosi: Require		
modafanil (generic for Provigil) ^{CL} SUNOSI (solriamfetol) ^{CL,QL} WAKIX (pitolisant) ^{CL,QL} WAKIX (pitolisant) CL,QL					
SUNOSI (solriamfetol) CL,QL WAKIX (pitolisant) CL,QL documentation/confirm via sleep study and documentation that C-l has been maxed Narcolepsy with			• •		
	SUNOSI (riamfetol) ^{CL,QL}	documentation/confirmation via sleep study and documentation that C-PAP has been maxed		
documentation of diagrams of the contraction of the contr			documentation of diagnosis via sleep study o Shift Work Sleep Disorder		
months) with work sche verified and documente Shift work is defined as working the all night sh			months) with work schedule verified and documented. Shift work is defined as working the all night shift		
via sleep study and documentation that C-l has been maxed		- S	 Sleep Apnea with documentation/confirmation via sleep study and documentation that C-PAP has been maxed 		
via sleep study			documentation of diagnosis via sleep study		
daytime sleepiness in adults w.		d n n			

TETRACYCLINES

Non-Preferred Agents **Preferred Agents Prior Authorization/Class Criteria**

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug. $\frac{QL}{QL}$ — Quantity/Duration Limit $\frac{AL}{QL}$ — Age Limit

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

doxycycline hyclate IR (generic Vibramycin)

doxycycline monohydrate **50MG**, **100MG CAPSULE**

doxycycline monohydrate **SUSP**, **TABLET** (generic Vibramycin)

minocycline HCI **CAPSULE**, **TABLET** (generic Dynacin/ Minocin/Myrac)

demeclocycline (generic Declomycin)^{CL}

DORYX MPC DR (doxycycline pelletized)

doxycycline hyclate DR (generic Doryx)

doxycycline monohydrate 40MG, 75MG and 150MG **CAPSULES** (generic for Adoxa/Monodox/ Oracea)

minocycline HCl ER (generic Solodyn) NUZYRA (omadacycline) tetracycline

VIBRAMYCIN **SUSP** (doxycycline) XIMINO (minocycline ER)^{QL}

 Non-preferred agents will be approved for patients who have failed an 3-day trial of TWO preferred agents within this drug class

Drug-specific criteria:

- Demeclocycline: Approved for diagnosis of SIADH
- Doryx®/doxycycline hyclate DR/ Dynacin®/Oracea®/Solodyn®: Requires clinical reason why generic doxycycline, minocycline or tetracycline cannot be used
- doxycycline suspension: May be approved with documented swallowing difficulty

THROMBOPOIESIS STIMULATING PROTEINSCL

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

THYROID HORMONES

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

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

with Prior Authorization Criteria

Highlights indicate changes from previous posting and November 2020 P&T changes, effective January 21, 2021

ULCERATIVE COLITIS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORAL		Non-preferred agents will be
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)	 approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Asacol HD®/Delzicol DR®/Lialda®/Pentasa®: Requires clinical reason why preferred mesalamine products cannot be used Giazo®: Requires clinical reason why generic balsalazide cannot be
RECTAL		used
CANASA (mesalamine)	mesalamine ENEMA (generic Rowasa) mesalamine SUPPOSITORY (generic Canasa) UCERIS (budesonide)	NOT covered in females

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

VASODII ATORS, CORONARY

VAGODILATORO, CORORART		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate TABLET isosorbide dinitrate ER, SA TABLET (generic Dilatrate-SR/Isordil) isosorbide mono IR/SR TABLET nitroglycerin SUBLINGUAL, TRANSDERMAL nitroglycerin ER TABLET	BIDIL (isosorbide dinitrate/ hydralazine) ^{CL} GONITRO (nitroglycerin) NITRO-BID OINTMENT (nitroglycerin) NITRO-DUR (nitroglycerin) nitroglycerin TRANSLINGUAL (generic Nitrolingual) NITROMIST (nitroglycerin)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: BiDil: Approved for the treatment of heart failure as an adjunct therapy to standard therapy in self-identified black patients

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