



Contains May 2020 P&T Changes
Noted in Red Font that Become Effective July 16, 2020

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 150 Morphine Milligram Equivalents (MME) per day to 120 Morphine Milligram Equivalents (MME) per day. (beginning September 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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ACNE AGENTS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ű	<u> </u>	
AZELEX (azelaic acid) benzoyl peroxide (BPO) GEL, WASH, LOTION OTC clindamycin/BPO (generic Duac) clindamycin phosphate SOLUTION	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) NR benzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide FOAM (generic 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-BPO (generic for Benzamycin) EVOCLIN (clindamycin) FABIOR (tazarotene foam) NEUAC (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (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 In the second

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ALZHEIMER'S AGENTS

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANALGESICS, OPIOID LONG-ACTING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) QL PATCH fentanyl 25, 50, 75, 100 mcg PATCH 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 PATCH QL 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020 **ANALGESICS, OPIOID SHORT-ACTING**^{QL}

acetaminophen/codeine ELIXIR, TABLET codeine TABLET APADAZ (benzhydrocodon Apadaz,CL	ne/APAP/codeine	 Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the last 12 months
morphine CONC SOLUTION, SOLUTION, TABLET oxycodone TABLET, SOLUTION oxycodone/APAP PROLATE (oxycodone/acetaminophen) tramadol TABLET ^{AL} BUDONE (hydrolevorphanol meperidine (genemorphine SUPPO) NALOCET (oxycodone/APAP) NALOCET (oxycodone/APAP) NALOCET (oxycodone/APAP) NALOCET (oxycodone/APAP) OXAYDO (oxycodone/APAP) oxycodone/APAP oxycodone/APAP oxycodone/APAP oxycodone/ibupro oxymorphone IR pentazocine/nalo PRIMLEV (oxycodone/nalo	SA/caffeine/codeine) apound-codeine al/ASA/codeine) APAP/caffeine spirin/caffeine EINE (butalbital/ e/caffeine) LIQUID, DRY (generic Dilaudid) acodone/ibuprofen) PRICE Demerol DISTORIES	 Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will consist of -prescriptions limited to a 7 day supply, AND -initial opiate prescription fill limited to maximum of 50 Morphine Milligram Equivalents (MME) per day These limits may only be exceeded with patient specific documentation of medical necessity, with examples such as, cancer diagnosis, end-of-life care, palliative care, Sickle Cell Anemia, or prescriber attestation that patient is not recently opiate naive Drug-specific criteria: Apadaz: Approval for 14 days or less Nucynta®: Approved only for diagnosis of acute pain, for 30 days or less Tramadol/APAP: Clinical reason

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020 **ANALGESICS, OPIOID SHORT-ACTING**^{QL} (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

ANDROGENIC AGENTS (Topical)

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®: Approved for Males only Natesto®: Approved for Males on with diagnosis of: Primary hypogonadism (congenital or acquired) Hypogonadotropic hypogonadism (congenital or acquired)

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ANGIOTENSIN MODULATORS

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

Nebraska Medicaid **Preferred Drug List**

with Prior Authorization Criteria

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANGIOTENSIN MODULATORS (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria	
ANGIOTENSIN RECEPTOR BLOCKER/DIURETIC COMBINATIONS		Non-preferred agents will be approved for petionts who have	
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 withis drug class within the last 12 months Non-preferred combination products may be covered as individual prescriptions without prior authorization 	thin 2
	I MODULATOR/ OCKER COMBINATIONS	Angiotensin Modulator/Calciu Channel Blocker Combination	um ns:
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)	Combination agents may be approved if there has been a tri and failure of preferred agent	
DIRECT RENI	N INHIBITORS	Direct Renin Inhibitors/Direct Renin Inhibitor Combinations	
	aliskiren (generic Tekturna) ^{QL}	May be approved witha history TWO preferred ACE Inhibitors of	
DIRECT RENIN INHIB	ITOR COMBINATIONS	Angiotensin Receptor Blockers within the last 12 months	
	TEKTURNA/HCT (aliskiren/HCTZ)	- Within the last 12 months	
	TOR COMBINATION		
ENTRESTO (sacubitril/valsartan) ^{CL}			
ANGIOTENSIN RECEPTOR BLOCKI	ER/BETA-BLOCKER COMBINATIONS		
	BYVALSON (nevibolol/valsartan)		

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ANTHELMINTICS

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

ANTI-ALI FRGENS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ORALAIR (sweet vernal/orchard/rye/ timothy/kentucky blue grass mixed pollen allergen extract) PALFORZIA ^{NR,AL} (peanut allergen powder-dnfp)	 Class Criteria: Approved for immunotherapy for the treatment of grass pollen-induced allergic rhinitis with or without conjunctivitis. Patient has had treatment failure with or contraindication to: antihistamines AND montelukast Clinical reason as to why allergy shots cannot be used. Drug-specific criteria: ORALAIR Confirmed by positive skin teror in vitro testing for pollenspecific IgE antibodies for Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass Mixed Pollens. For use in patients 10 through 65 years of age.

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIBIOTICS, GASTROINTESTINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FIRVANQ (vancomycin) SOLUTION netronidazole TABLET neomycin	ALINIA (nitazoxanide) ^{CL} SUSPENSION DIFICID (fidaxomicin) ^{CL} FLAGYL ER (metronidazole) ^{CL} Metronidazole ^{CL} CAPSULE paromomycin SOLOSEC (secnidazole) tinidazole (generic Tindamax) ^{CL} vancomycin CAPSULE (generic Vancocin) ^{CL} XIFAXAN (rifaximin) ^{CL}	 Note: Although azithromycin, ciprofloxacin and trimethoprim/ sulfmethoxazole are not included in this review, they are available without prior authorization Drug-specific criteria: Alinia®: Trial and failure with metronidazol is required for a diagnosis of giardiasis Dificid®: Trial and failure with oral vancomycin is required for a diagnosis of difficile diarrhea (pseudomembranous colitis) Flagyl ER®: Trial and failure with metronidazole is required Flagyl®/Metronidazole 375mg capsules and Flagyl ER®/ Metronidazole 750mg Etabs: 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 Lomotil® AND Imodium®

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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 Tobi)	 Diagnosis of Cystic Fibrosis is required for all agents ICD10 Group = E84, ICD9 = 277.00, 277.01, 277.02, 277.03, 277.09 Drug-specific criteria: Arikayce: Requires diagnosis of refractory MAC lung disease defined as patients who did not achieve negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy Cayston®: Trial of tobramycin via nebulizer and demonstration of TOBI® compliance required Tobi Podhaler®: Requires trial of tobramycin via nebulizer or documentation why nebulized tobramycin cannot be used

ANTIBIOTICS, TOPICAL

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg ^{CL,QL}	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:

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIEMETICS/ANTIVERTIGO AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CANNABINOIDS		Non-preferred agents will be
dronabinol (generic Marinol) ^{AL}	CESAMET (nabilone)	approved for patients who have failed ONE preferred agent within this drug class within the same
5HT3 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 5-HT3 antagonist WITHOUT trial of
NK-1 RECEPTO	R ANTAGONIST	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,
TRADITIONAL	ANTIEMETICS	Carbplatin, Cisplatin, Clofarabine,
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)	Bendamustine, Busulfan, Carmustine, 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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche) fluconazole SUSPENSION, TABLET (generic Diflucan) griseofulvin SUSPENSION griseofulvin microsized TABLET nystatin SUSPENSION, TABLET terbinafine (generic Lamisil)	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 approve for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class Cresemba®: Approved for diagnosis 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), Neutropenia hematologic malignancies, Graft vs. Host disease (GVHD), Immunosuppression secondary to hematopoietic stem cell transplant Noxafil® Suspension:

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIFUNGALS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole CREAM (generic Lotrimin) RX, OTC clotrimazole SOLN OTC ketoconazole CREAM, SHAMPOO (generic Nizoral) LAMISIL (terbinafine) SPRAY OTC LAMISIL AT CREAM (terbinafine) OTC miconazole CREAM, POWDER OTC nystatin terbinafine OTC (generic Lamisil AT) tolnaftate POWDER, CREAM, POWDER OTC (generic Tinactin)	ciclopirox SHAMPOO (generic Loprox) clotrimazole SOLUTION RX (generic Lotrimin) DESENEX POWDER OTC (miconazole) econazole (generic Spectazole) ERTACZO (sertaconazole) EXELDERM (sulconazole) FUNGOID OTC JUBLIA (efinaconazole) KERIDYN (tavaborole) ketoconazole FOAM (generic Extina, Ketodan) LAMISIL AT GEL, SPRAY (terbinafine) OTC LOPROX (ciclopirox) SUSPENSION, SHAMPOO, CREAM LOTRIMIN AF CREAM OTC (clotrimazole) LOTRIMIN ULTRA (butenafine) luliconazole (generic Luzu) MENTAX (butenafine) miconazole OTC OINTMENT, SPRAY miconazole/zinc oxide/petrolatum (generic Vusion) naftifine CREAM, GEL (generic Naftin) oxiconazole (generic Bensal HP) tolnaftate SPRAY, OTC	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months Drug-specific criteria: Extina: Requires trial and failure or contraindication to other ketoconazole forms Jublia: Approved diagnoses includ Onychomycosis of the toenails due to <i>T.rubrum OR T. Mentagrophytes</i> nystatin/triamcinolone: Indivudual ingredients available without prior authorization ciclopirox nail lacquer: No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine
clotrimazole/betamethasone CREAM (generic Lotrisone)	clotrimazole/betamethasone LOTION (generic Lotrisone) nystatin/triamcinolone (generic Mycolog)	

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIHISTAMINES, MINIMALLY SEDATING

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

ANTIHYPERTENSIVES, SYMPATHOLYTICS

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

ANTIHYPERURICEMICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
allopurinol (generic for Zyloprim) colchicine CAPSULE (generic for Mitigare) probenecid probenecid/colchicine (generic for Col- Probenecid)	colchicine TABLET (generic for Colcrys) ^{CL} febuxostat (generic for Uloric) ^{CL} GLOPERBA SOLN (colchicine) ^{NR,CL,QL}	 Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class colchicine tablet®: Approved without trial for familial Mediterranean fever OR pericarditis Gloperba: Approved for documenter 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 – Age Limit

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ANTIMIGRAINE AGENTS, OTHER

Preferred Agents Non-Preferred Agents	
AJOVY (fremanezumab-vfrm) CL, QL EMGALITY 120 mg/mL (galcanezumab- gnlm) CL, QL PEN, SYRINGE NURTEC ODT (rimegepant) AL, CL, QL EMGALITY 100 mg (galcanezumab- gnlm) CL, QL SYRINGE REGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL REYVOW (lasmiditan) AL, CL, QL TABLET UBRELVY (ubrogepant) AL, CL, QL TABLET	 Non-preferred agents will be approved for patients who have a contraindication OR trial failure of a triptan 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 or patient specific documentation of why Emgality 120mg is not appropriate for patient Nurtec ODT, Reyvow, and Ubrelvy will be approved for patients who have a failed trial or contraindication to 2 triptan agents

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020 **ANTIMIGRAINE AGENTS, TRIPTANS**^{QL}

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

ANTIPARASITICS, TOPICAL

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIPARKINSON'S AGENTS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOL benztropine (generic for Cogentin) trihexyphenidyl (generic for Artane)	INERGICS	 Non-preferred agents will be approved for patients who have failed ONE preferred agents within this drug class
		Drug-specific criteria: Carbidopa/Levodopa ODT: Approved for documented swallowing disorder COMT Inhibitors: Approved if using as add-on therapy with levodopacontaining 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 levodopacontaining drug Inbrija: Approval upon diagnosis of Parkinson's disease and concurrent treatment with carbidopa/levodopa agent Neupro®:
amantadine CAPSULE, SYRUP TABLET (generic for Symmetrel) carbidopa/levodopa (generic for Sinemet) carbidopa/levodopa ER (generic for Sinemet CR) levodopa/carbidopa/entacapone (generic for Stalevo)	carbidopa (generic for Lodosyn) carbidopa/levodopa ODT (generic for Parcopa) DUOPA (carbidopa/levodopa) GOCOVRI (amantadine) ^{QL} INBRIJA (levodopa) INHALER ^{CL,QL} KYNMOBI (apomorphine) ^{QL/NR} NOURIANZ (istradefylline) ^{NR,CL,QL} OSMOLEX ER (amantadine) ^{QL} RYTARY (carbidopa/levodopa) STALEVO (levodopa/carbidopa/entacapone)	For Parkinsons: Clinical reason required why preferred agent cannot be used For Restless Leg (RLS): Requires trial OR Contraindication to ropinirole AND pramipexole Nourianz: Approval upon diagnosis of 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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ANTIPSORIATICS, ORAL

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

ANTIPSORIATICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcipotriene CREAM, OINTMENT, SOLUTION,	calcitriol (generic for Vectical) calcipotriene/betamethasone	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-HERPI	ETIC 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-INFLUE	NZA DRUGS	 Drug-specific criteria:
oseltamivir (generic Tamiflu) ^{QL}	rimantadine (generic Flumadine) RELENZA (zanamivir) ^{QL} TAMIFLU (oseltamivir) ^{QL} XOFLUZA (baloxavir marboxil) ^{AL,CL,QL}	 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

ANTIVIRALS, TOPICAL

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

ANXIOLYTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
alprazolam TABLET (generic for Xanax) buspirone (generic for Buspar) chlordiazepoxide diazepam TABLET , SOLUTION (generic for Valium) lorazepam INTENSOL , TABLET (generic for Ativan)	alprazolam ER (generic for Xanax XR) alprazolam ODT alprazolam INTENSOL ^{CL} clorazepate (generic for Tranxene-T) diazepam INTENSOL ^{CL} 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®

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

BETA BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atenolol (generic Tenormin) atenolol/chlorthalidone (generic Tenoretic) bisoprolol (generic Zebeta) bisoprolol/HCTZ (generic Ziac) metoprolol (generic Lopressor) metoprolol ER (generic Toprol XL) propranolol (generic Inderal) propranolol ER (generic Inderal LA)	acebutolol (generic Sectral) betaxolol (generic Kerlone) BYSTOLIC (nebivolol) HEMANGEOL (propranolol) SOLUTION INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) LEVATOL (penbutolol) metoprolol/HCTZ (generic Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide pindolol (generic Viskin) propranolol/HCTZ (generic Inderide) timolol (generic Blocadren) TOPROL XL (metoprolol ER)	 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 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)	

BILE SALTS

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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: Myrbetriq®: Covered without trial in contraindication to anticholinergic agents

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

BONE RESORPTION SUPRESSION AND RELATED DRUGS

risedronate (generic Actonel) ^{QL} other Bone resorption suppression and related Drugs calcitonin-salmon NASAL raloxifene (generic Evista) EVISTA (raloxifene) FORTEO (teriparatide) ^{QL} Treiparatide ^{QL} TYMLOS (abaloparatide) Forteo®: Covered for high risk of fracture High risk of fracture: BMD -3 or worse Postmenopausal women with history on non-traumatic fractures Postmenopausal women with 2 or more clinical reason why alendronate tablets OR Fosamax® solution cannot be used Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification Forteo®: Covered for high risk of fracture High risk of fracture: BMD -3 or worse Postmenopausal women with 12 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 Reumatoid Arthritis Postmenopausal women with BMD T-	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
TABLET ibandronate (generic Fosamax) TABLET ibandronate (generic Boniva) ^{QL} ATELVIA DR (risedronate) etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL} OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS calcitonin-salmon NASAL raloxifene (generic Evista) EVISTA (raloxifene) FORTEO (teriparatide) ^{QL} TyMLOS (abaloparatide) TYMLOS (abaloparatide) Fortoe: Covered for high risk of fracture High risk of fracture: BMD -3 or worse Postmenopausal women with history on more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Requires clinical reason and betternophic ossification Fortoe: Covered for high risk of fracture: BMD -3 or worse Postmenopausal women with bistory on more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Postmenopausal women with 2 or more clinical risk factors Requires clinical reason and the similar fractures Drug-specific criteria: Actionale® Combinations: Covered as individual agents without prior authorization Activate Pisconese Requires clinical reason and empty stomach Binosto®: Requires clinical reason and empty stomach Etidronate disodium: Trial not required for diagnosis of heterrophic ossification Fortoe: Covered for high risk of fracture High risk of fracture: BMD -3 or worse Postmenopausal women with BMD T-score ≤ -2.5 at any site with any clinical risk factors Activate Pisconese S-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	BISPHOSE	PHONATES	
BINOSTO (alendronate) etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL} etidronate (generic Actonel) ^{QL} risedronate (generic Actonel) ^{QL} OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS calcitonin-salmon NASAL raloxifene (generic Evista) EVISTA (raloxifene) FORTEO (teriparatide) ^{QL} Teriparatide ^{QL} TYMLOS (abaloparatide) TYMLOS (abaloparatide) Fostmenopausal women with history on non-traumatic fractures Postmenopausal women with bistory on non-traumatic fractures Postmenopausal women with Driscore ≤ -2.5 at any site Glucocorticoid use ≥ 6 months at 7.5 dose of prednisolone equivalent Report Services Servi	TABLET	Fosamax) ^{QL}	failed a trial of ONE preferred
etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL} other BONE RESORPTION SUPPRESSION AND RELATED DRUGS calcitonin-salmon NASAL raloxifene (generic Evista) EVISTA (raloxifene) FORTEO (teriparatide) ^{QL} TYMLOS (abaloparatide) TYMLOS (abaloparatide) Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnosis of hetertrophic ossification Fortse ^{QL} Submendate disodium: Trial not required for diagnos	ibandronate (generic Boniva)	,	Drug-specific criteria:
TymLOS (abaloparatide) TymLo		etidronate disodium (generic Didronel) FOSAMAX PLUS D^QL	 Actonel® Combinations: Covered as individual agents without prior authorization Atelvia DR®: Requires clinical reason alendronate cannot be taken on an empty
Teriparatide of the post menor with pistory of non-traumatic fractures Postmenopausal women with history of non-traumatic fractures Postmenopausal women with bistory of non-traumatic fractures DXA BMD T-score ≤ -2.5 at any site factors Reumatoid Arthritis Postmenopausal women with BMD T-score ≤ -2.5 at any site factors Reumatoid Arthritis Postmenopausal women with BMD T-score ≤ -2.5 at any site factors Current smoker Men with primary or hypogonadal			
calcitonin-salmon NASAL raloxifene) FORTEO (teriparatide) ^{QL} Teriparatide ^{QL} TYMLOS (abaloparatide) Teriparatide one (generic Evista) Touriparatide one (generic Evista) Teriparatide one of the properties of the properties of the properties of the properties of the diagnosis of hetertrophic ossification one of the diagnosis of hetertrophic ossification one of the properties of the p	OTHER BONE RESORPTION SUPP	PRESSION AND RELATED DRUGS	alendronate tablets OR Fosamax® solution
Osteoporosis associated with sustained systemic glucocorticoid therapy		FORTEO (teriparatide) ^{QL} Teriparatide ^{QL}	 Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification Forteo®: Covered for high risk of fracture High risk of fracture: BMD -3 or worse Postmenopausal women with history of non-traumatic fractures Postmenopausal women with 2 or more clinical risk factors Family history of non-traumatic fractures DXA BMD T-score ≤ -2.5 at any site Glucocorticoid use ≥ 6 months at 7.5 dose of prednisolone equivalent Rheumatoid Arthritis Postmenopausal women with BMD T-score ≤ -2.5 at any site with any clinical risk factors More than 2 units of alcohol per day Current smoker Men with primary or hypogonadal osteoporosis Osteoporosis associated with sustained systemic glucocorticoid

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
alfuzosin (generic Uroxatral) doxazosin (generic Cardura) tamsulosin (generic Flomax) terazosin (generic Hytrin)	Non-Preferred Agents LOCKERS CARDURA XL (doxazosin) silodosin (generic Rapaflo) SE (5AR) INHIBITORS dutasteride/tamsulosin (generic for Jalyn)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-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
		 Homax[®]: Females covered for a 7 day supply with diagnosis of acute kidney stones Jalyn[®]: Requires clinical reason why individual agents cannot be used

BRONCHODILATORS, BETA AGONIST

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INHALERS –	Short Acting	Non-preferred agents will be
PROAIR HFA (albuterol)	albuterol HFA (generic for ProAir	approved for patients who have failed a trial of ONE preferred
PROVENTIL HFA (albuterol)	HFA, Proventil HFA, Ventolin HFA)	agent within this drug class
	levalbuterol HFA (generic for Xopenex	
	HFA)	Drug-specific criteria:
	PROAIR DIGIHALER (albuterol) ^{NR}	 Ventolin HFA®: Requires trial and failure on Proventil HFA® AND
	PROAIR RESPICLICK (albuterol)	Proair HFA® OR allergy/
		contraindication/side effect to
INHALERS – Long Acting		BOTH
SEREVENT (salmeterol)	ARCAPTA NEOHALER (indacaterol)	Xopenex®: Covered for cardiac diagnoses or side effect of
	STRIVERDI RESPIMAT (olodaterol)	tachycardia with albuterol product
INHALATIO	N SOLUTION	-
albuterol (2.5mg/3ml premix or	BROVANA (arformoterol)	
2.5mg/0.5ml)	levalbuterol (generic for Xopenex)	
albuterol 100 mg/20 mL	PERFOROMIST (formoterol)	
albuterol low dose (0.63mg/3ml &		
1.25mg/3ml)		
	RAL	
albuterol SYRUP	albuterol TABLET albuterol ER (generic for Vospire ER)	
	metaproterenol (formerly generic for	
	Alupent)	
	terbutaline (generic for Brethine)	
	- (3	

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

CALCIUM CHANNEL BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SHORT-ACTING		Non-preferred agents will be approved for patients who have
Dihydro	Dihydropyridines	
	isradipine (generic Dynacirc)	failed a trial of ONE preferred agent within this drug class
	nicardipine (generic Cardene)	
	nifedipine (generic Procardia)	Drug-specific criteria:
	nimodipine (generic Nimotop)	Nifedipine: May be approved without trial for diagnosis of
	NYMALIZE (nimodipine) SOLUTION	Preterm Labor or Pregnancy
Non-dihyd	ropyridines	Induced Hypertension (PIH)
diltiazem (generic Cardizem)		Nimodipine: Covered without trial for diagnosis of subarachnoid
verapamil (generic Calan/Isoptin)		hemorrhage
LONG-ACTING		 Katerzia: May be approved with documented swallowing difficulty
Dihydro	Dihydropyridines	
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)	

Nebraska Medicaid **Preferred Drug List**

with Prior Authorization Criteria

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

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

COLONY STIMULATING FACTORS

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

CONTRACEPTIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
All reviewed agents are recommended preferred at this time Only those products for review are listed. Brand name products may be subject to Maximum Allowable Cost (MAC) pricing or require substitution with a generic equivalent	hailey FE 1/20 (norethindrone acetate and ethinyl estradiol tablets USP and ferrous fumarate) ^{NR}	
Specific agents can be looked up using the Drug Look-up Tool at: https://druglookup.fhsc.com/druglookupweb/?client=nestate		

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ATROVENT HFA (ipratropium) BEVESPI AEROSPHERE (glycopyrolate/formoterol) COMBIVENT RESPIMAT (albuterol/ipratropium) SPIRIVA (tiotropium) STIOLTO RESPIMAT (tiotropium/olodaterol)	ANORO ELLIPTA (umeclidinium/vilanterol) DUAKLIR PRESSAIR (aclidinium br and formoterol fum) ^{NR} INCRUSE ELIPTA (umeclidnium) SEEBRI NEOHALER (glycopyrolate) SPIRIVA RESPIMAT (tiotropium) TUDORZA PRESSAIR (aclidinium br) UTIBRON NEOHALER (indacaterol/glycopyrolate)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class OR Patient specific documentation of inability to use traditional inhaler device. Drug-specific criteria: Daliresp®: Covered for diagnosis of severe COPD associated with chronic bronchitis 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}	

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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} <i>TRIKAFTA</i> (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

CYTOKINE & CAM ANTAGONISTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ENBREL (etanercept) KIT, MINI CART, PEN ^{QL} HUMIRA (adalimumab) ^{QL} OTEZLA (apremilast) ORAL ^{CL,QL}	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIMZIA (certolizumab pegol)QL COSENTYX (secukinumab)GL ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYRINGE KINERET (anakinra) OLUMIANT (baricitinib) ORALCL,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) ORALCL,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 Humiral documentation of inadequate response or intolerance to methotrexate and an inadequate response to one or more TNF antagonist therapies. Rinvoq, Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to methotrexate

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

DIURETICS

Preferred Agents	Non-Preferred Agents		Prior Authorization/Class Criteria
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)	•	Non-preferred agents will be 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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, linezolid, sulfamethoxazole/trimethoprim) Ciprofloxacin/Levofloxacin Suspension: Coverable with documented swallowing disorders Ofloxacin: Trial of preferred not required for diagnoses of Pelvic Inflammatory Disease OR Acute Epididymitis (nongonorrhea)

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

GI MOTILITY, CHRONIC

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

GLUCAGON AGENTSQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) ^{AL,CL} 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 Drug Specific Criteria: Baqsimi approved for patients who failed a trial of ONE preferred injectable agent in this class

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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 RESPICLICK (fluticasone) ^{AL} ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) ^{CL,AL,QL} FLOVENT DISKUS (fluticasone) QVAR (beclomethasone) QVAR Redihaler (beclomethasone) BODILATOR COMBINATIONS BREO ELLIPTA (fluticasone/vilanterol) Budesonide/formoterol (generic for Symbicort) fluticasone/salmeterol (generic for Advair Diskus) ^{QL} fluticasone/salmeterol (generic for Airduo Respiclick) TRELEGY ELLIPTA (fluticasone/ umeclidinium/vilanterol) WIXELA INHUB (generic for Advair Diskus) ^{QL}	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)	

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

GLUCOCORTICOIDS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
budesonide EC CAPSULE (generic for Entocort EC) dexamethasone SOLN, TABLET dexamethasone ELIXIR, SYRUP hydrocortisone TABLET methylprednisolone DOSE PAK methylprednisolone tablet (generic for Medrol) prednisolone SOLUTION prednisolone sodium phosphate prednisone DOSE PAK prednisone TABLET	cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSPENSION, TABLETCL ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

H. PYLORI TREATMENTS

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

HAE TREATMENTSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BERINERT (C1 esterase inhibitor, human) INTRAVENOUS FIRAZYR (icatibant acetate) ^{AL} SUB-Q HAEGARDA (C1 esterase inhibitor, human) ^{AL} SUB-Q	CINRYZE (C1 esterase inhibitor, human)AL INTRAVENOUS icatibant acetate (generic for FIRAZYR)AL SUB-Q KALBITOR (ecallantide)AL SUB-Q RUCONEST (recombinant human C1 inhibitor)AL INTRAVENOUS TAKHZYRO (lanadelumab-flyo)AL SUB-Q	 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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

HEMOPHILIA TREATMENTS

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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, (sofosbuvir/ledipasvir)CL HARVONI (ledipasvir/sofosbuvir)CL,NR PELLET sofosbuvir/ledipasvir (generic Harvoni)CL sofosbuvir/velpatasvir (generic Epclusa)CL SOVALDI (sofosbuvir)CL,NR PELLET SOVALDI TABLET (sofosbuvir)CL VIEKIRA PAK (ombitasvir/ paritaprevir/ritonavir/dasabuvir)CL ZEPATIER (elbasvir/grazoprevir)CL	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) FERON	 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 non-response after previous treatment course of Direct Acting Anti-viral agent (DAA) for genotype 1-6 without cirrhosis or with compensated cirrhosis

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

HISTAMINE II RECEPTOR BLOCKERS

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

HIV / AIDSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CCR5 AN	TAGONISTS	 Non-preferred agents will be
SELZENTRY SOLN, TAB (maraviroc)		approved for patients who have a diagnosis of HIV/AIDS and patient
FUSION I	NHIBITORS	 specific documentation of why the preferred products within this drug
FUZEON SUB-Q (enfuvirtide) ^{QL}		class are not appropriate for patient, including, but not limited
INTEGRASE STRAND TRA	NSFER INHIBITORS (INSTIS)	to, drug resistance or concomitant
ISENTRESS (raltegravir) ^{QL} ISENTRESS HD (raltegravir) TIVICAY (dolutegravir)	TIVICAY PD (dolutegravir) ^{NR}	 conditions not recommended with preferred agents Patients undergoing treatment at the time of any preferred status
NON-NUCLEOSIDE REVERSE TRA	NSCRIPTASE INHIBITORS (NNRTIS)	change will be allowed to continue therapy
EDURANT (rilpivirine) INTELENCE (etravirine) ^{QL} PIFELTRO (doravirine) ^{QL} SUSTIVA CAPSULE, TABLET (efavirenz)	efavirenz (generic Sustiva) nevirapine IR, ER (generic Viramune/Viramune XR) RESCRIPTOR (delavirdine) VIRAMUNE (nevirapine) SUSP	 Diagnosis of HIV/AIDS required OR Pre and Post Exposure Prophylaxis
NUCLEOSIDE REVERSE TRAN	SCRIPTASE INHIBITORS (NRTIs)	_
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPSULE, SOLN (emtricitabine) lamivudine SOLN, TABLET (generic Epivir) zidovudine CAPSULE, SYRUP, TABLET (generic Retrovir)	didanosine DR (generic Videx EC) EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPSULE (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
NUCLEOTIDE REVERSE TRAN	ISCRIPTASE INHIBITORS (NRTIs)	_
tenofovir TABLET (generic Viread)		
PHARMACOKIN	IETIC ENHANCER	
TYBOST (cobicistat) ^{QL}		
PROTEASE	INHIBITORS	
atazanavir CAPSULE (generic Reyataz LEXIVA SUSP, TABLET (fosamprenavir) NORVIR (ritonavir) TAB PREZISTA (darunavir) SUSP, TABLET	(tipranavir) CRIXIVAN (indinavir) fosamprenavir TAB (generic Lexiva)	

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

Page **41** of **77**

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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} opinavir/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} amivudine/zidovudine (generic Combivir) TRUVADA (emtricitabine/tenofovir)	COMBIVIR (lamivudine/zidovudine) EPZICOM (abacavir sulfate/lamivudine) TEMIXYS (lamivudine/tenofovir) ^{QL} TRIZIVIR (abacavir/lamivudine/zidovudine)	
	CTS – MULTIPLE CLASSES	
ATRIPLA (tenofovir/emtricitabine/efavirenz) BIKTARVY (bictegravir/emtricitabine/tenofovir)QL COMPLERA (rilpivirine/emtricitabine/tenofovir) DELSTRIGO (doravirine/lamivudine/tenofovir)QL GENVOYA (elvitegravier/cobicistat/emtricitabine/tenofovir)QL, AL ODEFSEY (emtricitabine/rilpivirine/tenofovir)QL STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir)QL STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir)QL SYMFI (efavirenz/lamivudine/tenofovir)QL SYMFI LO (efavirenz/lamivudine/tenofovir)QL TRIUMEQ (dolutegravir/abacavir/lamivudine)	DOVATO (dolutegravir/lamivudine) ^{QL} JULUCA (dolutegravir/rilpivirine) ^{QL} SYMTUZA (darunavir/cobicistat/ emtricitabine/tenofovir) ^{QL}	

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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) <i>RYBELSUS</i> (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 ₹ AND
INSULIN/GLP-1 RA	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
	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
DIPEPTIDYL PEPTIDASI	E-4 (DPP-4) INHIBITOR ^{QL}	
GLYXAMBI (empagliflozin/linagliptin) JANUMET (sitagliptin/metformin) JANUMET XR (sitagliptin/metformin) JANUVIA (sitagliptin) JENTADUETO (linagliptin/metformin) TRADJENTA (linagliptin)	alogliptin (generic for Nesina)	Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within DPP-4

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

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

HYPOGLYCEMICS, MEGLITINIDES

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

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

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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)		

HYPOGLYCEMICS, TZD

Preferred Agents	Non-Preferred Agents	Prior Authorization	n/Class Criteria
THIZAOLIDINEDIONES (TZDs)		Non-preferred agents will be	
pioglitazone (generic for Actos)	AVANDIA (rosiglitazone)		IE preferred agent
TZD COMBINATIONS		within this drug class	
	pioglitazone/glimepiride (generic for Duetact) pioglitazone/metformin (generic for Actoplus Met)	Combination pro clinical reason wh ingredients cannot	ny individual

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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

IMMUNOMODULATORS, ATOPIC DERMATITISAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIDEL (pimecrolimus)	DUPIXENT (dupilumab) ^{CL} EUCRISA (crisaborole) 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: For atopic dermatitis, must have trial of Eucrisa; For moderate to severe asthma, must have eosinophilic phenotype or oral corticosteroid dependent asthma uncontrolled with maintenance controller medication; For adults with chronic rhinosinusitis with nasal polyposis, must document inadequate control on current treatment regimen and be used as add-on maintenance treatment with intranasal steroid

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

IMMUNOSUPPRESSIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathiaprine (generic Imuran) cyclosporine, modified CAPSULE (generic Neoral) mycophenolate CAPSULE, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLUTION 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
ANTICHO	LINERGICS	Non-preferred agents will be approved
pratropium (generic for Atrovent)		for patients who have failed a 30-day trial of ONE preferred agent within this
ANTIHIS	TAMINES	drug class
azelastine 0.1% (generic for Astelin)	azelastine 0.15% (generic for Astepro) azelastine/fluticasone (generic for Dymista) olopatadine (generic for Patanase)	 Drug-specific criteria: mometasone: Prior authorization NOT required for children ≤ 12 years budesonide: Approved for use in Pregnancy (Pregnancy Category
CORTICO	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)	 Veramyst®: Prior authorization 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. CL – Prior Authorization / Class Criteria apply QL – Quantity/Duration Limit AL – Age Limit

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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

incogainides / Oxazolidinones / STREFTOGRAMINS		
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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

LIPOTROPICS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BILE ACID SE	QUESTRANTS	 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
	JUXTAPID (lomitapide) ^{CL}	inadequate
	KYNAMRO (mipomersen) ^{CL}	Juxtapid®/ Kynamro®:
FIBRIC ACID DERIVATIVES		 Approved for diagnosis of homozygous 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,
niacin ER (generic for Niaspan)	NIACOR (niacin IR) NIASPAN (niacin ER)	bile acid sequestrantsRequire faxed copy of REMS PA form
OMEGA-3 F	ATTY ACIDS	_ Lovaza ®: Approved for TG ≥ 500
CHOLESTEROL ABSO	omega-3 fatty acids (generic for Lovaza) ^{CL} VASCEPA (icosapent) ^{CL} DRPTION INHIBITORS	 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
ezetimibe (generic for Zetia)	NEXLIZET (bempedoic acid/ezetimibe) ^{NR,QL}	

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

LIPOTROPICS, OTHER (continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BTILISIN/KEXIN TYPE 9 (PCSK9) BITORS 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)	ATINS ALTOPREV (lovastatin ER) ^{CL} EZALLOR SPRINKLE (rosuvastatin) ^{QL} fluvastatin IR/ER (generic Lescol/ Lescol XL) LIVALO (pitavastatin) ZYPITAMAG (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: Altoprev®: One of the TWO trials must be IR lovastatin Combination products: Require clinical
STATIN CO	MBINATIONS	reason why individual ingredients cannot be
	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

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

MACROLIDES AND KETOLIDES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MACRO	MACROLIDES	
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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

MOVEMENT DISORDERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) ^{CL} tetrabenazine (generic for Xenazine) ^{CL}	INGREZZA (valbenazine) ^{CL} CAP, INITIATION PACK	Non-preferred agent requires trial of Austedo
		All drugs require an FDA approved indication – ICD-10 diagnosis code required.
		Drug-specific criteria:
		 Austedo: Diagnosis of Tardive Dyskinesia or chorea associated with Huntington's Disease; Requires a Step through tetrabenazine with the diagnosis of chorea associated with Huntington's Disease
		 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) dalfampridine (generic Ampyra) ^{QL} EXTAVIA (interferon beta-1b) ^{QL} glatiramer (generic Copaxone) ^{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

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

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

NSAIDs, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Ü	diclofenac potassium (generic for Cataflam, Zipsor) diclofenac SR (generic for Voltaren-XR) diflunisal (generic for Dolobid)	Prior Authorization/Class Criteria Non-preferred agents within COX-1 SELECTIVE group will be approved for patients who have failed no less than 30-day trial of TWO preferred agents within this drug class Drug-specific criteria:
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)	etodolac & SR (generic for Lodine/XL) fenoprofen (generic for Nalfon) flurbiprofen (generic for Ansaid) ibuprofen OTC (generic for Advil,	 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 meloxicam suspension: Approved for age≤ 11 years

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

NSAIDs, ORAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COX-I SELECT	ALL BRAND NAME NSAIDs including: CAMBIA (diclofenac oral solution)	Drug-specific criteria: Sprix®: Approved for patients unable to tolerate, swallow OR absorb oral NSAIDs OR
	DUEXIS (ibuprofen/famotidine) SPRIX (ketorolac nasal spray) NASALQL, CL TIVORBEX (indomethacin) VIVLODEX (meloxicam submicronized) ZIPSOR (diclofenac) ZORVOLEX (diclofenac)	 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
NSAID/GI PROTECTA	ANT COMBINATIONS	
	diclofenac/misoprostol (generic for Arthrotec)	
COX-II SE	ELECTIVE	
celecoxib (generic for Celebrex)		

NSAIDs, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	diclofenac (generic for Pennsaid Solution) ^{CL} FLECTOR PATCH (diclofenac) ^{CL} <i>LICART PATCH</i> (diclofenac) PENNSAID PACKET , PUMP (diclofenac) ^{CL} VOLTAREN GEL (diclofenac) ^{CL}	 Flector®: Approved for diagnosis of acute pain due to sprain/strain/contusion AND trial of oral diclofenac OR clinical reason patient cannot use oral dosage form 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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
CDK 4/6 INHIBITOR		Non-preferred agents DO NOT
IBRANCE (palbociclib)	KISQALI (ribociclib) KISQALI FEMARA CO-PACK VERZENIO (abemaciclib)	require a trial of a preferred agent, but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
CHEMO	THERAPY	- - Drug-specific critera
cyclophosphamide XELODA (capecitabine)	capecitabine (generic for Xeloda) ^{CL}	 anastrozole: May be approved for malignant neoplasm of male breast (male breast cancer)
HORMONE BLOCKADE		capecitabine: Requires trial of Xeloda or clinical reason Xeloda
anastrozole (generic for Arimidex) exemestane (generic for Aromasin) letrozole (generic for Femara) tamoxifen citrate (generic for Nolvadex)	SOLTAMOX SOLN (tamoxifen) ^{CL} toremifene (generic for Fareston) ^{CL}	 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
ОТ	HER	for short term use
	NERLYNX (neratinib) PIQRAY (alpelisib) TYKERB (lapatinib) TALZENNA (talazoparib tosylate) QL TUKYSA(tucatinib)NR,QL	 Soltamox: May be approved with documented swallowing difficulty

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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	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) LEUKERAN (chlorambucil) VENCLEXTA (venetoclax)	DAURISMO (glasdegib maleate) ^{QL} IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) ^{QL} XOSPATA (gilteritinib) ^{QL} LL COPIKTRA (duvelisib) ^{QL} ZYDELIG (idelalisib)	 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 Tasigna: Patients receiving Tasigna, which changed from
	ML	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
М	PN	-
JAKAFI (ruxolitinib)		-
MYE	LOMA	
ALKERAN (melphalan) REVLIMID (lenalidomide)	FARYDAK (panobinostat) melphalan (generic for Alkeran) NINLARO (ixazomib) POMALYST (pomalidomide) THALOMID (thalidomide) XPOVIO (selinexor) CL	
ОТ	HER	
MATULANE (procarbazine) TABLOID (thioguanine) tretinoin (generic for Vesanoid)	BRUKINSA (zanubrutinib) ^{NR,QL} CALQUENCE (acalabrutinib) ^{QL} INREBIC (fedratinib dihydrochloride) ^{QL} ZOLINZA (vorinostat)	

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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
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
ALK/F	ROS1 / NTRK	
XALKORI (crizotinib)	ROZLYTREK (entrectinib) AL,QL	
	EGFR	
IRESSA (gefitinib) TAGRISSO (osimertinib)	erlotinib (generic for Tarceva) GILOTRIF (afatinib) TARCEVA (erlotinib) VIZIMPRO (dacomitinib) ^{QL}	
	OTHER	
	HYCAMTIN (topotecan) RETEVMO (selpercatinib) ^{NR,AL} TABRECTA (capmatinib) ^{NR,QL}	

ONCOLOGY AGENTS, ORAL, OTHER

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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

ONCOLOGY AGENTS, ORAL, RENAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
LENVIMA (lenvatinib) SUTENT (sunitinib) VOTRIENT (pazopanib)	AFINITOR DISPERZ (everolimus) ^{cL} CABOMETYX (cabozantinib) everolimus (generic for Afinitor) INLYTA (axitinib) 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
BASAL (RIVEDGE (vismodegib)	DOMZO (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
FINLAR (dabrafenib)	RAFTOVI (encorafenib)	rug-specific critera Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17-
EKINIST (trametinib) FINLAR (dabrafenib) N	RAFTOVI (encorafenib) DTELLIC (cobimetinib) EKTOVI (binimetinib)	rug-specific critera Odomzo: Patients Odomzo, which ch

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

Nebraska Medicaid **Preferred Drug List**

with Prior Authorization Criteria

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

OPHTHALMICS, ALLERGIC CONJUNCTIVITIS

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

OPHTHALMICS, ANTIBIOTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FLUOROQUINOLONES		 Non-preferred agents will be
ciprofloxacin SOLUTION (generic for Ciloxan) MOXEZA (moxifloxacin) ofloxacin (generic for Ocuflox)	BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) gatifloxacin 0.5% (generic for Zymaxid) levofloxacin moxifloxacin (generic for Vigamox) moxifloxacin (generic for Moxeza) VIGAMOX (moxifloxacin)	 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
MACRO	DLIDES	documented fungal infection
erythromycin	AZASITE (azithromycin) ^{CL}	
AMINOGL	YCOSIDES	
gentamicin SOLUTION tobramycin (generic for Tobrex drops) TOBREX OINTMENT (tobramycin)	gentamicin OINTMENT	
OTHER OPHTH	ALMIC AGENTS	
bacitracin/polymyxin B (generic Polysporin) polymyxin B/trimethoprim (generic for Polytrim)	bacitracin NATACYN (natamycin) ^{CL} neomycin/bacitracin/polymyxin B OINTMENT neomycin/polymyxin B/gramicidin NEOSPORIN (neomycin/polymyxin B/gramcidin) sulfacetamide SOLUTION (generic for Bleph-10) sulfacetamide OINTMENT	

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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

OPHTHALMICS, ANTI-INFLAMMATORIES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CORTICOSTEROIDS		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%) 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	<u> </u>
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%)	

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 **60** of **77**

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
RESTASIS (cyclosporine) RESTASIS MULTIDOSE (cyclosporine)	CEQUA (cyclosporine) QL XIIDRA (lifitegrast)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

OPHTHALMICS, GLAUCOMA

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIOTICS		Non-preferred agents will be
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide)	approved for patients who have failed a trial of ONE preferred agent within this drug class
SYMPATHO	MIMETICS	
brimonidine 0.2% (generic for Alphagan)	Alphagan P (brimonidine 0.1%) Alphagan P (brimonidine 0.15%) apraclonidine (generic for lopidine)	
BETA BL	OCKERS	_
levobunolol (generic for Betagan) timolol (generic for Timoptic)	betaxolol (generic for Betoptic) BETIMOL (timolol) BETOPTIC S (betaxolol) carteolol (generic for Ocupress) timolol (generic for Istalol) TIMOPTIC OCUDOSE TIMOPTIC XE (timolol gel forming solution)	
CARBONIC ANHYD	RASE INHIBITORS	
dorzolamide (generic for Trusopt)	AZOPT (brinzolamide)	
PROSTAGLAN	DIN ANALOGS	
latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATI	ON DRUGS	
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine	
ОТ	HER	•
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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

OPIOID DEPENDENCE TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SUBOXONE 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 per label. Trial of preferred product not required.

OPIOID-REVERSAL TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naloxone SYRINGE, VIAL naltrexone TABLET NARCAN (naloxone) SPRAY		 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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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 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

DALI (DI II MONADY ADTEDIAI LIVDEDTENSION ACENTS). ODAL AND INLIAI ED

PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED			
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria	
ADCIRCA (tadalafil) ^{CL} ambrisentan (generic Letairis) sildenafil TABLET (generic Revatio) ^{CL} TRACLEER TABLET (bosentan) TYVASO INHALATION (treprostinil) VENTAVIS INHALATION (iloprost)	ADEMPAS (riociguat) ^{CL} bosentan TABLET (generic Tracleer) <i>LETAIRIS</i> (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) sildenafil SUSPENSION (generic Revatio) ^{CL} tadalafil (genericAdcirca) ^{CL} 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®:	

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

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD LITTLE ANIMALS VITAMINS CHEW OTC (pedi multivit 91/iron fum) CHEW child multivitamins chew otc (pedi multivit 19/folic acid) CHEW CHILDREN'S CHEW MULTIVIT-IRON OTC (pedi multivit 91/iron fum) CHEW	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)	 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
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	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 A,C,D3, 21/fluoride) DROPS infant-toddler multivit drop OTC (pediatric multivit no. 165 drops)	multivit 153/D3/K) POLY-VI-FLOR (pedi multivit 33/fluoride) CHEW POLY-VI-FLOR (pedi multivit 37/fluoride) DROPS	
infant-toddler multivit-iron OTC (pedi mv no.164/ferrous sulfate drops) infant-toddler tri-vit drop (vit a	POLY-VI-FLOR w/IRON (pedi multivit 33/fluoride/iron) CHEW POLY-VI-FLOR w/IRON (pedi multivit 37/fluoride/iron) DROPS	
palmitate/vit c/vit d3 drops) multivitamins with fluoride (pedi multivit 2/fluoride) DROPS multivits with iron and fluoride (pedi	QUFLORA OTC and Rx (pedi multivit 84/fluoride) QUFLORA FE (pedi multivit 142/iron/fluoride)	
multivit 45/fluoride/iron) DROPS MVC-FLUORIDE (pedi multivit 12/fluoride) CHEW TAB ped mvi A,C,D3,No 21/fluoride DROPS	TRI-VI-FLORO (ped multivit A, C, D3, 38/fluoride)	
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		

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

PENICILLINS

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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 voltamin 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/omega3) trust natal dha (pnv2/iron b-g suc-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 dss/dha) virt-select CAPSULE (pnv80/iron fum/fa/dss/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#16/iron fum &ps/fa/om-3) virt-pm dha SOFTGEL (pnv combo#47/iron/fa #1/dha) zatean-pn dha CAPSULE (pnv #47/iron/fa #1/dha)	Non-preferred agents will be approved for patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class Non-preferred agents within trial of the patients of the patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class.

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

PROGESTERONE (hydroxyprogesterone caproate)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MAKENA AUTO INJECTOR (hydroxyprogesterone caproate) MAKENA MDV, SDV (hydroxyprogesterone caproate)	hydroxyprogesterone caproate (generic Makena)	 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)	DEXILANT (dexlansoprazole) esomeprazole magnesium (generic Nexium) esomeprazole strontium lansoprazole (generic Prevacid) NEXIUM SUSPENSION (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) 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:

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

SEDATIVE HYPNOTICS

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

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

SKELETAL MUSCLE RELAXANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
paclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) ^{QL} nethocarbamol (generic Robaxin) izanidine 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 cannobe 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 day 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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

STEROIDS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
LOW POTENCY		Low Potency Non-preferred agents
hydrocortisone OTC & RX CREAM, LOTION, OINTMENT hydrocortisone/aloe OINTMENT, CREAM 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) MICORT-HC (hydrocortisone) TEXACORT (hydrocortisone)	will be approved for patients who have failed a trial of ONE preferred agent within this drug class
MEDILIM	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)	Medium Potency Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

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 drug class
triamcinolone LOTION	betamethasone dipropionate betamethasone / propylene glycol	
	betamethasone valerate	
	desoximetasone	
	diflorasone diacetate fluocinonide SOLUTION	
	fluocinonide CREAM, GEL, OINTMENT	
	fluocinonide emollient	
	halcinonide CREAM (generic for Halog)	
	HALOG (halcinonide) CREAM , SOLN ^{NR}	
	KENALOG AEROSOL (triamcinolone)	
	SERNIVO (betamethasone dipropionate)	
	triamcinolone SPRAY (generic for	
	Kenalog spray) TRIANEX OINTMENT (triamcinolone)	
	VANOS (fluocinonide)	
VERY HIG	H POTENCY	very riight oterioy rion preferred
clobetasol emollient (generic for Temovate-E)	APEXICON-E (diflorasone)	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class
clobetasol propionate CREAM, GEL,	BRYHALI (halobetasol prop) LOTION clobetasol SHAMPOO, LOTION	
OINTMENT, SOLUTION	clobetasol propionate FOAM, SPRAY	S
halobetasol propionate (generic for	CLOBEX (clobetasol)	
Ultravate)	halobetasol propionate FOAM (generic	
	for Lexette) AL,QL	
	LEXETTE(halobetasol propionate) AL,QL	
	OLUX-E /OLUX/OLUX-E CP (clobetasol)	
	,	

Nebraska Medicaid **Preferred Drug List**

with Prior Authorization Criteria

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020 **STIMULANTS AND RELATED AGENTS**^{AL}

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

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Methylphe APTENSIO XR (methylphenidate) dexmethylphenidate (generic for Focalin IR)	ADHANSIA XR (methylphenidate) QL CONCERTA (methylphenidate ER)QL 18mg, 27mg, 36mg, 54mg	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class Maximum accumulated dose of 108mg per day for ages < 18
FOCALIN XR (dexmethylphenidate) METHYLIN SOLUTION (methylphenidate) methylphenidate (generic for Ritalin) methylphenidate 30/70 (generic for Metadate CD) methylphenidate SOLUTION (generic for Methylin) methylphenidate ER 10mg, 20mg (generic for Ritalin SR, Metadate ER) methylphenidate ER 18mg, 27mg, 36mg, 54mg (generic Concerta) ^{QL} QUILLICHEW ER CHEWTAB (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 LA) methylphenidate ER (generic for Ritalin SR) methylphenidate ER CAP (generic for Aptensio XR) ^{NR,QL} methylphenidate ER 72mg (generic for RELEXXI) ^{QL} QUILLIVANT XR SUSP (methylphenidate) RITALIN (methylphenidate)	 Maximum accumulated dose of 72mg per day for ages > 19

May 2020 P&T Proposed Changes *Highlighted in Red become* effective July 16, 2020

STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

TETRACYCLINES

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

THYROID HORMONES

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) THYROLAR TABLET (liotrix) TIROSINT TABLET (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

May 2020 P&T Proposed Changes Highlighted in Red become effective July 16, 2020

ULCERATIVE COLITIS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
OR	AL	Non-preferred agents will be
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine)	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
REC	TAL	used NOT covered in females
CANASA (mesalamine)	mesalamine ENEMA (generic Rowasa) mesalamine SUPPOSITORY (generic Canasa) UCERIS (budesonide)	NOT covered in ternales

UTERINE DISORDER TREATMENT - ENDOMETRIOSIS

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

VASODILATORS, CORONARY

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate TABLET isosorbide dinitrate ER, SA TABLET (generic Dilatrate-SR/Isordil) isosorbide mono IR/SR TABLET nitroglycerin SUBLINGUAL, TRANSDERMAL nitroglycerin ER TABLET	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