



DEPT. OF HEALTH AND HUMAN SERVICES

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

October 2022 PDL

Noted in Red Font that Become Effective October 1, 2022

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

- PDMP Check Requirements- Nebraska Medicaid providers are required to check the
 prescription drug history in the statewide PDMP before prescribing CII controlled substances to
 certain Medicaid beneficiaries. (Exemption to this requirement are for beneficiaries receiving
 cancer treatment, hospice/palliative care, or in long-term care facilities). If not able to check the
 PDMP, then provider is required to document good faith effort, including reasons why unable to
 conduct the check and may be required to submit documentation to the State upon request.
 - PDMP check requirements are under Section 5042 of the SUPPORT for Patients and Communities Act, consistent with section 1944 of the Social Security Act [42 U.S.C. 1396w-3a], beginning October 1, 2021.
- Opioids- The maximum opioid dose covered will decrease from 120 Morphine Milligram
 Equivalents (MME) per day to 90 Morphine Milligram Equivalents (MME) per day. (beginning
 December 1, 2020)

Non-Preferred Drug Coverage

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

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

Specific Class Prior Authorization forms can be found within the PDL class listings and at: https://nebraska.fhsc.com/priorauth/paforms.asp

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

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

Documentation of Medical Necessity PA Form

with Prior Authorization Criteria

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

ACNE AGENTS, TOPICAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benzoyl peroxide (BPO) WASH, LOTION clindamycin/BPO (generic Benzaclin) PUMP clindamycin phosphate PLEDGET clindamycin phosphate SOLUTION DIFFERIN LOTION, CREAM, Rx-GEL (adapalene) DIFFERIN GEL (adapalene) OTC erythromycin GEL erythromycin SOLN erythromycin-BPO (generic for Benzamycin) RETIN-A (tretinoin) AL CREAM, GEL	adapalene (generic differin) adapalene/BPO (generic Epiduo) adapalene/BPO (generic Epiduo Forte) AKLIEF (trifarotene) AL ALTRENO (tretinoin)AL AMZEEQ (minocycline) ARAZLO (tazarotene)AL ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) AZELEX (azelaic acid) BENZACLIN PUMP (clindamycin/BPO) BENZEFOAM (benzoyl peroxide) benzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide GEL OTC benzoyl peroxide GEL OTC benzoyl peroxide GEL Rx benzoyl peroxide TOWELETTE OTC clindamycin FOAM, LOTION clindamycin GEL clindamycin/BPO (generic Acanya) GEL clindamycin/BPO (generic Duac) clindamycin/BPO (generic Duac) clindamycin/tretinoin (generic Veltin, Ziana) dapsone (generic Aczone) EPIDUO FORTE GEL PUMP (adapalene/BPO) erythromycin GEL, PLEDGET erythromycin-BPO (generic for Benzamycin) FVOCLIN (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) OVACE PLUS (sulfacetamide sodium) PLIXDA (adapalene) SWAB RETIN-A ^{AL} GEL, CREAM (tretinoin) sulfacetamide sulfacetamide/sulfur SUMADAN (sulfacetamide/sulfur) tazarotene CREAM, GELNR (generic Tazorac) tazarotene FOAM (generic Fabior) TRETIN-X (tretinoin) tretinoin CREAM, GELAL (generic Avita, Retin-A) tretinoin microspheres (generic for Retin-A Micro) AL TWYNEO (tretinoin/BPO) AL, NR CREAM	Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class It is approved for patients who have failed THREE preferred agents within this drug class.

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

AL_ Age Limit

with Prior Authorization Criteria

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ALZHEIMER'S AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
donepezil (generic Aricept) donepezil ODT (generic Aricept ODT) EXELON Transdermal (rivastigmine)	ASE INHIBITORS ADLARITY (donepezil) ^{NR} PATCH donepezil 23 (generic Aricept 23) galantamine (generic Razadyne) SOLN, TABLET galantamine ER (generic Razadyne ER) rivastigmine (generic for Exelon)	 Non-preferred agents will be approved for patients who have failed a 120-day trial of ONE preferred agent within this drug class within the last 6 months OR Current, stabilized therapy of the non-preferred agent within the previous 45 days
memantine (generic for Namenda)	memantine ER (generic for Namenda XR) memantine SOLN (generic for Namenda) NAMENDA (memantine) NAMZARIC (memantine/donepezil)	 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)

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ANALGESICS, OPIOID LONG-ACTING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) ^{QL} PATCH fentanyl 25, 50, 75, 100 mcg PATCH ^{QL} morphine ER TABLET (generic MS Contin, Oramorph SR) OXYCONTIN ^{CL} (oxycodone ER) tramadol ER (generic Ultram ER) ^{CL}	ARYMO ER (morphine sulfate) QL BELBUCA (buprenorphine) QL BUCCAL buprenorphine BUCCAL (generic for Belbuca) AL, QL buprenorphine PATCH (generic Butrans) QL EMBEDA (morphine sulfate/ naltrexone) DURAGESIC MATRIX (fentanyl) QL fentanyl 37.5, 62.5, 87.5 mcg PATCHQL hydrocodone ER (generic for Hysingla ER) QL hydrocodone bitartrate ER (generic for Zohydro ER) hydromorphone ER (generic for Exalgo) CL HYSINGLA ER (hydrocodone ER) KADIAN (morphine ER) methadone TABLET CL methadone ORAL SYRCL MORPHABOND ER (morphine sulfate) morphine ER (generic for Avinza, Kadian) CAPS NUCYNTA ER (tapentadol) CL oxycodone ER (generic Oxycontin) oxymorphone ER (generic Opana ER) tramadol ER (generic Conzip) CL	The Center for Disease Control (CDC) does not recommend long acting opioids when beginning opioid treatment. • Preferred agents require previous use of a long acting opioid or documentation of a trial on a short acting agent within 90 days • Non-preferred agents will be approved with failure on, or intolerance to TWO preferred agents within this drug class Drug-specific criteria: • Methadone: Will only be approved for use in pain control or end of life care. Trial of preferred agent not required for end of life care • Oxycontin®: Pain contract required for maximum quantity authorization

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ANALGESICS, OPIOID SHORT-ACTINGQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
· ·		 Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the last 12 months Note: for short acting opiate tablets and capsules there is a maximum quantity limit of #150 per 30 days. Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will
oxycodone/APAP Tramadol 50 TABLETAL (generic Ultram) tramadol/APAP (generic Ultracet)	dihydrocodeine/aspirin/caffeine FIORINAL/CODEINE (butalbital/ ASA/codeine/caffeine) hydromorphone LIQUID, SUPPOSITORY (generic Dilaudid) IBUDONE (hydrocodone/ibuprofen) levorphanol meperidine (generic Demerol) morphine SUPPOSITORIES NALOCET (oxycodone/APAP) NUCYNTA (tapentadol) ^{CL} OXAYDO (oxycodone) ^{CL} oxycodone/APAP SOLN oxycodone/APAP SOLN oxycodone/aspirin oxycodone/ibuprofen oxymorphone IR (generic Opana) pentazocine/naloxone ROXICODONE TABLET (oxycodone) ROXYBOND (oxycodone) SEGLENTIS (celecoxib/tramadol) ^{AL} tramadol 100mg TABLET(generic Ultram) ^{AL} tramadol (generic Qdolo) ^{AL,QL} SOLN ZAMICET (hydrocodone/APAP)	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

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ANALGESICS, OPIOID SHORT-ACTINGQL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
N.A.	ASAL	
	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)CL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NDROGEL (testosterone) PUMP CL	ANDRODERM (testosterone) ^{CL} NATESTO (testosterone) ^{CL} testosterone PACKET (generic Androgel) ^{CL} testosterone PUMP (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 last 6 months Drug-specific criteria: Androderm®/Androgel®: Approved for Males only Natesto®: Approved for Males on with diagnosis of: Primary hypogonadism (congenital or acquired) OR Hypogonadotropic hypogonadism (congenital)

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ANGIOTENSIN MODULATORS

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

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ANGIOTENSIN MODULATORS (Continued)

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

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ANTHELMINTICS

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

ANTI-ALLERGENS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ORALAIR (sweet vernal/orchard/rye/timothy/kentucky blue grass mixed pollen allergen extract) PALFORZIA AL,CL (peanut allergen powder-dnfp)	 Drug-specific criteria: ORALAIR

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ANTIBIOTICS, GASTROINTESTINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FIRVANQ (vancomycin) SOLN netronidazole TABLET neomycin inidazole (generic Tindamax) ^{CL}	DIFICID (fidaxomicin) CL TABLET, SUSP FLAGYL ER (metronidazole)CL metronidazoleCL CAPS nitazoxanide (generic Alinia) TABLETAL, CL, QL paromomycin SOLOSEC (secnidazole) vancomycin CAPS (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®: For diagnosis of C. difficile diarrhea (pseudomembranous colitis), trial and failure or intolerance to oral vancomycin is required. For diagnosis of relapsed or recurrent C. difficile, an appropriate ICD-10 diagnosis code must be submitted for coverage. Flagyl ER®: Trial and failure with metronidazole is required Flagyl "Metronidazole 375mg capsules and Flagyl ER®/ Metronidazole 750mg E tabs: Clinical reason why the generic regular-release cannot be used tinidazole: 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's 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®

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

ANTIBIOTICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
bacitracin OINT bacitracin/polymyxin (generic Polysporin) mupirocin OINT (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/ pramoxine	CENTANY (mupirocin) gentamicin OINT, CREAM mupirocin CREAM (generic Bactroban) ^{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

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ANTIBIOTICS. VAGINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CLEOCIN OVULES (clindamycin) clindamycin CREAM (generic Cleocin) CLINDESSE (clindamycin) metronidazole, vaginal NUVESSA (metronidazole)	CLEOCIN CREAM (clindamycin) METROGEL (metronidazole) VANDAZOLE (metronidazole) XACIATO (clindamycin phosphate) vaginal gel ^{AL,NR}	Non-preferred agents will be approved for patients who have failed a therapeutic trial (duration = 3 days) with ONE preferred agent within this drug class within the las 6 months

ANTICOAGULANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ELIQUIS (apixaban) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) KARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg KARELTO (rivaroxaban) 2.5 mg ^{CL,QL} KARELTO DOSE PACK (rivaroxaban)	BEVYXXA (betrixaban) ^{QL} dabigatran etexilate ^{NR} (generic Pradaxa) fondaparinux (generic Arixtra) FRAGMIN (dalteparin) SAVAYSA (edoxaban) ^{QL} XARELTO (rivaroxaban) ^{CL} SUSP	 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:

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ANTIEMETICS/ANTIVERTIGO AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BINOIDS	Non-preferred agents will be approved for patients who have
dronabinol (generic Marinol) ^{AL}	CESAMET (nabilone)	failed ONE preferred agent within this drug class within the same
5HT3 RECEPTO	OR BLOCKERS	group
ondansetron (generic Zofran/Zofran ODT) ^{QL}	ANZEMET (dolasetron) granisetron (generic Kytril) SANCUSO (granisetron) ^{CL} ZUPLENZ (ondansetron)	Drug-specific criteria: • Akynzeo®/Varubi®: Approved for Moderately/Highly emetogenic chemotherapy with dexamethasone and a 5-HT3 antagonist
NK-1 RECEPTO	R ANTAGONIST	Regimens include: AC combination (Doxorubicin or Epirubicin with
EMEND (aprepitant) CAPS, CAPS PACK ^{QL}	aprepitant (generic Emend) QL,CL AKYNZEO (netupitant/palonosetron)CL VARUBI (rolapitant) TABLET CL	Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine, Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine, Dacarbazine, Dactinomycin, Daunorubicin,
TRADITIONAL	ANTIEMETICS	Epirubicin, Etoposide, Hexamethylmelamine, Idarubicin,
DICLEGIS (doxylamine/pyridoxine) ^{CL,QL} dimenhydrinate (generic Dramamine) OTC meclizine (generic Antivert) metoclopramide (generic Reglan) phosphoric acid/dextrose/fructose SOLN (generic Emetrol) prochlorperazine, oral (generic Compazine) promethazine SYRUP, TABLET (generic Phenergan) promethazine SUPPOSITORY 12.5mg, 25mg TRANSDERM-SCOP (scopolamine)	BONJESTA (doxylamine/pyridoxine),CL,QL COMPRO (prochlorperazine) doxylamine/pyridoxine (generic Diclegis)CL,QL metoclopramide ODT (generic Metozolv ODT) prochlorperazine SUPPOSITORY (generic Compazine) promethazine SUPPOSITORY 50mg scopolamine TRANSDERMAL trimethobenzamide TABLET (generic Tigan)	 Hexametry interamine, toardolcin, 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

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche) fluconazole SUSP, TABLET (generic Diflucan) griseofulvin SUSP griseofulvin microsized TABLET nystatin SUSP, TABLET terbinafine (generic Lamisil)	BREXAFEMME (ibrexafungerp) ^{QL} CRESEMBA (isavuconazonium) ^{CL} flucytosine (generic Ancobon) ^{CL} griseofulvin ultramicrosize (generic GRIS-PEG) itraconazole (generic Sporanox) ^{CL} ketoconazole (generic Nizoral) nystatin POWDER ONMEL (itraconazole) posaconazole (generic Noxafil) ^{AL,CL} TOLSURA (itraconazole) ^{CL} VIVJOA (oteseconazole) CAPS ^{NR} voriconazole (generic VFEND) ^{CL}	 Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class Drug-specific criteria: Cresemba®: Approved for diagnosis of invasive aspergillosis or invasive mucomycosis Flucytosine: Approved for diagnosis of: Candida: Septicemia, endocarditis, UTIs Cryptococcus: Meningitis, pulmonary infections Noxafil®: No trial for diagnosis of Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropenic hematologic malignancies, Graft vs. Host disease(GVHD), Immunosuppression secondary to hematopoietic stem cell transplant Noxafil® Suspension:

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ANTIFUNGALS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Clotrimazole CREAM (generic Lotrimin) RX, OTC clotrimazole SOLN OTC ketoconazole CREAM, SHAMPOO (generic Nizoral) LAMISIL (terbinafine) SPRAY OTC LAMISIL AT CREAM (terbinafine) OTC miconazole CREAM, POWDER OTC nystatin terbinafine OTC (generic Lamisil AT) tolnaftate POWDER, CREAM, POWDER OTC (generic Tinactin)	EUNGAL ALEVAZOL (clotrimazole) OTC ciclopirox CREAM, GEL, SUSP (generic Ciclodan, Loprox) ciclopirox NAIL LACQUERCL (generic Penlac) ciclopirox SHAMPOO (generic Loprox) clotrimazole SOLN RX (generic Lotrimin) DESENEX POWDER OTC (miconazole) econazole (generic Spectazole) ERTACZO (sertaconazole) EXELDERM (sulconazole) FUNGOID OTC JUBLIA (efinaconazole) ^{CL} ketoconazole FOAMCL (generic Extina, Ketodan) LAMISIL AT GEL, SPRAY (terbinafine) OTC LOPROX (ciclopirox) SUSP, SHAMPOO, CREAM LOTRIMIN AF CREAM OTC (clotrimazole) LOTRIMIN ULTRA (butenafine) luliconazole (generic Luzu) MENTAX (butenafine) miconazole OTC OINTMENT, SPRAY miconazole/zinc oxide/petrolatum (generic Vusion) naftifine CREAM, GEL (generic Naftin) oxiconazole (generic Oxistat) salicylic acid (generic Bensal HP) tavaborole SOLUTIONCL (generic Kerydin) 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 and tavaborole:
ANTIFUNGAL/STEF	ROID COMBINATIONS	
	clotrimazole/betamethasone LOTION (generic Lotrisone)	

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ANTIHISTAMINES, MINIMALLY SEDATING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
cetirizine TABLET, SOLN (Rx only) (generic for Zyrtec) loratadine TABLET, SOLN (generic for Claritin) levocetirizine TABLET (generic for Xyzal)	cetirizine CHEWABLE (generic for Zyrtec) cetirizine SOLN (OTC) 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) SOLN loratadine CAPS, CHEWABLE, ODT (generic fClaritin 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

with Prior Authorization Criteria

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ANTIHYPERURICEMICS

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

with Prior Authorization Criteria

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ANTIMIGRAINE AGENTS, OTHER

AJOVY (fremanezumab-vfrm) Autoinjector 3-pack ^{CL,QL} EMGALITY 120 mg/mL (galcanezumab- ELYXYB (celecoxib) ^{AL,QL} SOLN triptan. In addition, all non-preferred agents will require a failed trial or	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NURTEC ODT (rimegepant) ^{AL,CL,QL} UBRELVY (ubrogepant) ^{AL,CL,QL} TABLET ERGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL QULIPTA (atogepant) ^{AL,QL} REYVOW (lasmiditan) ^{AL,CL,QL} REYVOW (lasmiditan) ^{AL,CL,QL} TABLET TRUDHESA (dihydroergotamine mesylate) ^{AL,QL} NASAL agent of the same indication Drug-specific criteria: Cambia®: Requires diagnosis of migraine and documentation of why solid dosing forms not appropriate Emgality 120mg is recommended for preventative treatment of Migraine, Emgality 100mg is recommended for treatment of Episodic Cluster Headache Aimovig, Ajovy, Emgality 120mg, Nurtec ODT (prophylaxis), and Qulipta:: 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	PEN, Autoinjector AJOVY (fremanezumab-vfrm) Autoinjector 3-pack ^{CL,QL} EMGALITY 120 mg/mL (galcanezumab-gnlm) CL, QL PEN, SYRINGE NURTEC ODT (rimegepant) AL, CL, QL UBRELVY (ubrogepant) AL, CL, QL	CAFERGOT (ergotamine/caffeine) CAMBIA (diclofenac potassium) dihydroergotamine mesylate NASAL ELYXYB (celecoxib)AL,QL SOLN EMGALITY 100 mg (galcanezumabgnlm) CL,QL SYRINGE ERGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL QULIPTA (atogepant)ALQL REYVOW (lasmiditan)AL, CL,QL TABLET TRUDHESA (dihydroergotamine)	 approved for patients who have a failed trial or a contraindication to a triptan. In addition, all non-preferred agents will require a failed trial or contraindication of a preferred agent of the same indication Drug-specific criteria: Cambia®: Requires diagnosis of migraine and documentation of why solid dosing forms not appropriate Emgality 120mg is recommended for preventative treatment of Migraine, Emgaility 100mg is recommended for treatment of Episodic Cluster Headache Aimovig, Ajovy, Emgality 120mg, Nurtec ODT (prophylaxis), and Qulipta:: 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,

with Prior Authorization Criteria

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ANTIMIGRAINE AGENTS, TRIPTANSQL

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

with Prior Authorization Criteria

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ANTIPARASITICS, TOPICAL

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

with Prior Authorization Criteria

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ANTIPARKINSON'S AGENTS, ORAL

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

Nebraska Medicaid **Preferred Drug List** with Prior Authorization Criteria

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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
alcipotriene CREAM, OINT, SOLN	calcitriol (generic for Vectical) calcipotriene/betamethasone OINT (generic for Taclonex) calcipotriene/betamethasone SUSP (generic for Taclonex Scalp) CALCITRENE (calcipotriene) DOVONEX CREAM (calcipotriene) DUOBRII (halobetasol prop/tazarotene ENSTILAR (calcipotriene/betamethasone) SORILUX (calcipotriene) VTAMA CREAM (tapinarof) ^{AL,NR} ZORYVE CREAM (roflumilast) ^{AL,NR}	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

with Prior Authorization Criteria

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ANTIVIRALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTI-HERPETIC DRUGS		Non-preferred agents will be
acyclovir (generic Zovirax) famciclovir (generic Famvir) valacyclovir (generic Valtrex)	acyclovir (generic for Zovirax) ^{CL} SUSP SITAVIG (acyclovir buccal) ^{CL}	approved for patients who have failed a 10-day trial of ONE preferred agent within the same group
ANTI-INFLUENZA DRUGS		Drug-specific criteria:
oseltamivir (generic Tamiflu) ^{QL}	rimantadine (generic Flumadine) RELENZA (zanamivir) ^{QL} TAMIFLU (oseltamivir) ^{QL} XOFLUZA (baloxavir marboxil) ^{AL,CL,QL}	 Acyclovir Susp: Prior authorization NOT required for children ≤ 12 years old Sitavig®: Approved for recurrent herpes labialis (cold sores) in immunocompetent adults Xofluza: Requires clinical, patient specific reason that a preferred agent cannot be used

ANTIVIRALS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acyclovir OINT	acyclovir CREAM, (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

ANVIOLVTICE

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
alprazolam TABLET (generic for Xanax) buspirone (generic for Buspar) chlordiazepoxide diazepam TABLET , SOLN (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} lorazepam ORAL SYRINGE ^{NR} LOREEV XR (lorazepam) ^{AL.NR} meprobamate oxazepam	 Non-preferred agents will be approved for patients who have failed a trial with TWO preferred agents within this drug class Drug-specific criteria: Diazepam Intensol®: Requires clinical reason why diazepam solution cannot be used Alprazolam Intensol®: Requires trial of diazepam solution OR lorazepam Intensol®

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BETA BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atenolol (generic Tenormin) atenolol/chlorthalidone (generic Tenoretic) bisoprolol (generic Zebeta) bisoprolol/HCTZ (generic Ziac) metoprolol (generic Lopressor) metoprolol ER (generic Toprol XL) propranolol (generic Inderal) propranolol ER (generic Inderal LA)	acebutolol (generic Sectral) betaxolol (generic Kerlone) BYSTOLIC (nebivolol) HEMANGEOL (propranolol) SOLN INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) LEVATOL (penbutolol) metoprolol/HCTZ (generic Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide nebivolol (generic Bystolic) pindolol (generic Viskin) propranolol/HCTZ (generic Inderide) timolol (generic Blocadren) TOPROL XL (metoprolol ER)	 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 ^{CL} (generic Coreg CR)	
ANTIARR	HYTHMIC	
sotalol (generic Betapace)	SOTYLIZE (sotalol)	

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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ursodiol CAPSULE 300mg (generic Actigall) ursodiol 250mg TABLET (generic URSO) ursodiol 500mg TABLET (generic URSO FORTE)	BYLVAY (odevixibat) ^{NR} CAP, PELLET CHENODAL (chenodiol) CHOLBAM (cholic acid) LIVMARLI (maralixibat) SOLN ^{AL,NR} OCALIVA (obeticholic acid) RELTONE (ursodiol 200mg,400mg) CAP ^{NR}	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

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

with Prior Authorization Criteria

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BLADDER RELAXANT PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Oxybutynin IR, ER (generic Ditropan/Ditropan XL) solifenacin (generic Vesicare) TOVIAZ (fesoterodine ER)	darifenacin ER (generic Enablex) fesoterodine ^{NR} (generic Toviaz) flavoxate GELNIQUE (oxybutynin) GEMTESA (vibegron) ^{AL,QL} MYRBETRIQ TABLET , SUSP ^{AL,CL,QL} (mirabegron) OXYTROL (oxybutynin) tolterodine IR, ER (generic Detrol/ Detrol LA) trospium IR, ER (generic Sanctura/ Sanctura XR) VESICARE (solifenacin) VESICARE LS SUSP (solifenacin) AL	 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 Myrbetriq suspension: Covered for pediatric patients ≥ 3 years old with a diagnosis of Neurogenic Detrusor Overactivity (NDO)

with Prior Authorization Criteria

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BONE RESORPTION SUPRESSION AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BISPHOSPHONATES		Non-preferred agents will be
alendronate (generic Fosamax) TABLET ibandronate (generic Boniva) QL	alendronate SOLN (generic Fosamax) ^{QL} ATELVIA DR (risedronate)	approved for patients who have failed a trial of ONE preferred agent within the same group
	BINOSTO (alendronate)	Drug-specific criteria:
	etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL}	 Actonel® Combinations: Covered as individual agents without prior authorization Atelvia DR®: Requires clinical reason
	risedronate (generic Actonel) ^{QL}	alendronate cannot be taken on an empty stomach
	PRESSION AND RELATED DRUGS	Binosto®: Requires clinical reason why alendronate tablets OR Fosamax® solution cannot be used
calcitonin-salmon NASAL FORTEO (teriparatide) ^{CL,QL}	EVISTA (raloxifene) teriparatide (generic Forteo) ^{CL,QL}	Etidronate disodium: Trial not required for diagnosis of hetertrophic ossification
raloxifene (generic Evista)	TYMLOS (abaloparatide)	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 therapy
		Trial of calcitonin-salmon not required
		 Maximum of 24 months treatment per lifetime

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BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS

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

with Prior Authorization Criteria

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BRONCHODILATORS, BETA AGONIST

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
INHALERS – Short Acting		 Non-preferred agents will
albuterol HFA (generic for ProAir HFA)	albuterol HFA (Proventil HFA, Ventolin HFA) levalbuterol HFA (generic for Xopenex HFA) PROAIR DIGIHALER (albuterol) PROAIR RESPICLICK (albuterol) PROVENTIL HFA (albuterol) VENTOLIN HFA (albuterol)	be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: • Xopenex®: Covered for cardiac diagnoses or side effect of tachycardia with albuterol product
INH	ALERS – Long Acting	Ventolin HFA is temporarily authorized
SEREVENT (salmeterol)	ARCAPTA NEOHALER (indacaterol) STRIVERDI RESPIMAT (olodaterol)	due to ProAir HFA discontinuation
INI	HALATION SOLUTION	
albuterol (2.5mg/3ml premix or 2.5mg/0.5ml) albuterol 100 mg/20 mL albuterol low dose (0.63mg/3ml & 1.25mg/3ml)	arformoterol tartrate (generic Brovana) BROVANA (arformoterol) formoterol fumarate (generic Perforomist) levalbuterol (generic for Xopenex) PERFOROMIST (formoterol)	
	ORAL	
albuterol SYRUP	albuterol TABLET albuterol ER (generic for Vospire ER) metaproterenol (formerly generic for Alupent) terbutaline (generic for Brethine)	

with Prior Authorization Criteria

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CALCIUM CHANNEL BLOCKERS, ORAL

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

with Prior Authorization Criteria

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

COLONY STIMULATING FACTORS

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

with Prior Authorization Criteria

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CONTRACEPTIVES, ORAL

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

with Prior Authorization Criteria

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COPD (CHRONIC OBSTRUCTIVE PULMONARY DISEASE) AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANORO ELLIPTA (umeclidinium/vilanterol) ATROVENT HFA (ipratropium) COMBIVENT RESPIMAT (albuterol/ipratropium) SPIRIVA (tiotropium) STIOLTO RESPIMAT (tiotropium/olodaterol)	BEVESPI AEROSPHERE (glycopyrolate/formoterol) DUAKLIR PRESSAIR (aclidinium br and formoterol fum) INCRUSE ELIPTA (umeclidnium) 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 SOLN (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

with Prior Authorization Criteria

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CYSTIC FIBROSIS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BRONCHITOL (mannitol) ^{AL,CL,QL} KALYDECO PACKET, TABLET (ivacaftor) ^{QL, AL} ORKAMBI (lumacaftor/ivacaftor) PACKET, TABLET ^{QL, AL} SYMDEKO (tezacaftor/ivacaftor) ^{QL, AL} TRIKAFTA (elexacaftor, tezacaftor, ivacaftor) ^{AL, CL}	 Drug-specific criteria: Bronchitol: Approved for diagnosis of CF and documentation that the patient has passed the BRONCHITOL Tolerance Test 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

with Prior Authorization Criteria

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CYTOKINE & CAM ANTAGONISTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ENBREL (etanercept) KIT, MINI CART, PEN, SYR, VIAL ^{QL} HUMIRA (adalimumab) ^{QL} OTEZLA (apremilast) ORAL ^{CL,QL}	ACTEMRA (tocilizumab) SUB-Q ARCALYST (nilonacept) CIBINQO (abrocitinib) ^{AL,NR,QL} CIMZIA (certolizumab pegol) ^{QL} COSENTYX (secukinumab) ENSPRYNG (satralizumab-mwge) SUB-Q ILUMYA (tildrakizumab) SUB-Q KEVZARA (sarilumab) SUB-Q, PEN, SYRINGE KINERET (anakinra) OLUMIANT (baricitinib) TABLET ^{CL,QL} ORENCIA (abatacept) SUB-Q RINVOQ ER (upadacitinib) ^{CL,QL} SILIQ (brodalumab) SIMPONI (golimumab) SKYRIZI (risankizamab-rzaa) SYRINGE SKYRIZI ON-BODY (risankizamab-rzaa) ^{NR,QL} SKYRIZI PEN (risankizamab-rzaa) ^{QL} SOTYKTU (deucravacitinib) ^{NR} TABLET STELARA (ustekinumab) SUB-Q TALTZ (ixekizumab) ^{AL} TREMFYA (guselkumab) ^{QL} XELJANZ (tofacitinib) TABLET SCL,QL XELJANZ XR (tofacitinib) TABLET	 Preferred agents will be approved with FDA-approved indication – ICD-10 diagnosis code is required. Non-preferred agents will be approved for FDA-approved indications in patients who have failed a trial of ONE preferred agent within this drug class, or upon diagnosis for non-preferred agent with FDA-approved indication if no preferred agent has FDA approval for diagnosis. Drug-specific criteria: Otezla: Requires a trial of Humira Olumiant: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira) Rinvoq: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira) Xeljanz, Xeljanz XR: Requires documentation of inadequate response or intolerance to a Tumor Necrosis Factor (TNF) blocker (ex., Enbrel, Humira).

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DIURETICS

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

AL – Age Limit

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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 SUSP (generic Cipro) levofloxacin SOLN 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)

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

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GI MOTILITY, CHRONIC

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AMITIZA (lubiprostone) ^{AL, QL} LINZESS (linaclotide) ^{QL} MOVANTIK (naloxegol oxalate) ^{QL}	alosetron (generic Lotronex) IBSRELA (tenapanor) ^{AL,NR,QL} lubiprostone (generic Amitiza) ^{AL,QL} MOTEGRITY (prucalopride succinate) RELISTOR (methylnaltrexone) TABLET ^{QL} SYMPROIC (naldemedine) TRULANCE (plecanatide) ^{QL} VIBERZI (eluxodoline)	 Non-preferred agents will be approved for patients who have failed a 30-day trial of ONE preferred agent within this drug class 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 failure of loperamide AND diphenoxylate

GLUCAGON AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) ^{AL,QL} NASAL GLUCAGON EMERGENCY (glucagon) ^{QL} INJ KIT (Lilly) glucagon ^{QL} INJECTION PROGLYCEM (diazoxide) SUSP	diazoxide SUSP (generic Proglycem) GLUCAGON EMERGENCY (glucagon) ^{QL} INJ KIT (Fresenius) GVOKE (glucagon) ^{AL,QL} KIT, PEN, SYRINGE, VIAL ZEGALOGUE (dasiglucagon) ^{AL,QL} AUTO-INJECTOR, SYRINGE	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

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GLUCOCORTICOIDS, INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GLUCOCO	RTICOIDS	Non-preferred agents within the
ASMANEX (mometasone) QL,AL FLOVENT HFA (fluticasone) PULMICORT FLEXHALER (budesonide) GLUCOCORTICOID/BRONCH ADVAIR DISKUS (fluticasone/ salmeterol) QL	AEROSPAN (flunisolide) ALVESCO (ciclesonide) ^{AL,CL} ARMONAIR DIGIHALER (fluticasone) ^{AL,QL} ARMONAIR RESPICLICK (fluticasone) ^{AL} ARNUITY ELLIPTA (fluticasone) ASMANEX HFA (mometasone) ^{CL,AL,QL} FLOVENT DISKUS (fluticasone) fluticasone HFA (generic Flovent HFA) ^{NR} QVAR (beclomethasone) QVAR Redihaler (beclomethasone)	 Non-preferred agents within the Glucocorticoids and Glucocorticoid/Bronchodilator Combo groups will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months 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.
ADVAIR HFA (fluticasone/salmeterol) ^{QL} DULERA (mometasone/formoterol) SYMBICORT (budesonide/ formoterol)	BREO ELLIPTA (fluticasone/vilanterol) BREZTRI (budesonide/formoterol/glycopyrrolate)QL Budesonide/formoterol (generic for Symbicort) fluticasone/salmeterol (generic for Advair Diskus)QL fluticasone/salmeterol (generic for Airduo Respiclick) fluticasone/vilanterolNR (Breo Ellipta) TRELEGY ELLIPTA (fluticasone/umeclidinium/vilanterol) WIXELA INHUB (generic for Advair Diskus)QL	
	budesonide RESPULES (generic for Pulmicort)	

with Prior Authorization Criteria

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GLUCOCORTICOIDS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	ALKINDI (hydrocortisone) GRANULES ^{AL} CORTEF (hydrocortisone) cortisone TABLET dexamethasone INTENSOL DEXPAK (dexamethasone) DXEVO (dexamethasone) EMFLAZA (deflazacort) SUSP, TABLET ^{CL} ENTOCORT EC (budesonide) methylprednisolone 8mg, 16mg, 32mg ORTIKOS ER (budesonide) ^{AL,QL} PEDIAPRED (prednisolone sodium phosphate) prednisolone sodium phosphate	 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: 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)	HUMATROPE (somatropin)	Growth Hormone PA Form
NUTROPIN AQ (somatropin)	OMNITROPE (somatropin)	Growth Hormone Criteria
NORDITROPIN (somatropin)	SAIZEN (somatropin)	
	SEROSTIM (somatropin)	
	SKYTROFA (lonapegsomatropin-tcgd)	
	ZOMACTON (somatropin)	
	ZORBTIVE (somatropin)	

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

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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 HAEGARDA (C1 esterase inhibitor, human) AL,CL SUB-Q icatibant acetate (generic for FIRAZYR) BUB-Q	CINRYZE (C1 esterase inhibitor, human) ^{AL,CL} INTRAVENOUS FIRAZYR (icatibant acetate) ^{AL} SUB-Q ORLADEYO (berotralstat) CAP ^{AL,QL} RUCONEST (recombinant human	 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
	C1 inhibitor) ^{AL} INTRAVENOUS TAKHZYRO (lanadelumab-flyo) ^{AL,CL} VIAL, SYRINGE ^{NR}	 inhibitors, NSAIDs, or estrogen-containing products is contraindicated Non-preferred agents will be approved for patients who have a failed trial or a contraindication to ONE preferred agent within this drug class with the same indication
		Cinryze, Haegarda, Orladeyo, and Takhzyro, require a history of two or more HAE attacks monthly, and trial and failure or contraindication to oral danazol

with Prior Authorization Criteria

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HEMOPHILIA TREATMENTS

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

with Prior Authorization Criteria

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HEPATITIS B TREATMENTS

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

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HEPATITIS C TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIRECT ACTIN	IG ANTI-VIRAL	Hepatitis C Treatments PA Form
sofosbuvir/velpatasvir (generic Epclusa) ^{CL} MAVYRET (glecaprevir/pibrentasvir) TABLET^{CL}, PELLET^{AL,CL,NR} VOSEVI (sofosbuvir/velpatasvir/ voxilaprevir) ^{CL}	HARVONI 200/45MG, TABLET (sofosbuvir/ledipasvir) ^{CL} HARVONI (ledipasvir/sofosbuvir) ^{CL} PELLET sofosbuvir/ledipasvir (generic Harvoni) ^{CL} SOVALDI (sofosbuvir) ^{CL} 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/or will only be considered with documentation of why the preferred product within this drug class is not appropriate for patient Patients newly eligible for Medicaid will be allowed to complete treatment with the original that treatment was initially authorized by another payor Drug-specific criteria: Trial with with a preferred agent not required in the following: Harvoni: Post liver transplant for genotype
DIDA	VIRIN	1 or 4
	REBETOL (ribavirin)	Vosevi: Requires documentation of non- response after previous treatment course of Direct Acting Anti-viral agent (DAA) for
INTER	FERON	genotype 1-6 without cirrhosis or with compensated cirrhosis
PEGASYS (pegylated interferon alfa- 2a) CL PEG-INTRON (pegylated interferon alfa-2b) CL		— Compensated cirriosis

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HISTAMINE II RECEPTOR BLOCKERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
famotidine TABLET (generic for Pepcid) nizatidine SOLN (generic for Axid)	cimetidine TABLET , SOLN ^{CL} (generic for Tagamet) famotidine SUSP nizatidine CAPS (generic for Axid) ranitidine CAPS ,	 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 cimetidine solution/ famotidine suspension/ranitidine syrup: Requires clinical reason why nizatidine syrup cannot be used ***famotidine suspension is authorized during shortage of nizatidine syrup.***

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HIV / AIDSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CCR5 ANT	AGONISTS	 Non-preferred agents will be
SELZENTRY SOLN, TAB (maraviroc)	maraviroc (generic Selzentry)	approved for patients who have a diagnosis of HIV/AIDS and patien
FUSION IN	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 to, drug resistance or concomitan conditions not recommended with
HIV-1 ATTACH	MENT INHIBITOR	preferred agents
	RUKOBIA ER (fostemsavir)AL,QL	 Patients undergoing treatment at the time of any preferred status change will be allowed to continue
INTEGRASE STRAND TRAI	NSFER INHIBITORS (INSTIS)	therapy
ISENTRESS (raltegravir) ^{QL} ISENTRESS HD (raltegravir) TIVICAY (dolutegravir)	TIVICAY PD (dolutegravir)	 Diagnosis of HIV/AIDS required OR Pre and Post Exposure Prophylaxis
NON-NUCLEOSIDE REVERSE TRA	NSCRIPTASE INHIBITORS (NNRTIs)	
efavirenz CAPS, TABLET (generic Sustiva) INTELENCE (etravirine) ^{QL} PIFELTRO (doravirine) ^{QL}	EDURANT (rilpivirine) etravirine (generic Intelence) ^{QL} nevirapine IR, ER (generic Viramune/Viramune XR) RESCRIPTOR (delavirdine) SUSTIVA CAPS, TABLET (efavirenz) VIRAMUNE (nevirapine) SUSP	
NUCLEOSIDE REVERSE TRANS	SCRIPTASE INHIBITORS (NRTIs)	
abacavir SOLN, TABLET (generic Ziagen) EMTRIVA CAPS, SOLN (emtricitabine) lamivudine SOLN, TABLET (generic Epivir) zidovudine CAPS, SYRUP, TABLET (generic Retrovir)	didanosine DR (generic Videx EC) emtricitabine CAPS (generic for Emtriva) EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPS (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
NUCLEOTIDE REVERSE TRAN	SCRIPTASE INHIBITORS (NRTIs)	
tenofovir TABLET (generic Viread)	VIREAD (tenofovir) POWDER	
PHARMACOKIN	ETIC ENHANCER	
	TYBOST (cobicistat)QL	

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

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HIV / AIDS^{CL} (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
•	APTIVUS CAPS, SOLN (tipranavir) CRIXIVAN (indinavir) fosamprenavir TAB (generic Lexiva) INVIRASE (saquinavir) LEXIVA SUSP (fosamprenavir) LEXIVA TABLET (fosamprenavir) NORVIR POWDER, SOLN (ritonavir)	 Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with
	NORVIR (ritonavir) TAB PREZISTA (darunavir) SUSP, TABLET REYATAZ POWDER (atazanavir) VIRACEPT (nelfinavir)	preferred agents Patients undergoing treatment at

with Prior Authorization Criteria

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HIV / AIDSCL (Continued)

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

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HIV / AIDS^{CL} (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COMBINATION PRODU	CTS – MULTIPLE CLASSES	
BIKTARVY (bictegravir/emtricitabine/ tenofovir)QL COMPLERA (rilpivirine/emtricitabine/tenofovir) DELSTRIGO (doravirine/lamivudine/tenofovir)QL DOVATO (dolutegravir/lamivudine)QL efavirenz/emtricitabine/tenofovir (generic Atripla)CL GENVOYA (elvitegravier/cobicistat/emtricitabine/tenofovir)QL, AL ODEFSEY (emtricitabine/rilpivirine/tenofovir)QL STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir)QL SYMFI (efavirenz/lamivudine/tenofovir)QL SYMFI LO (efavirenz/lamivudine/tenofovir)QL SYMTUZA (darunavir/cobicistat/emtricitabine/tenofovir)QL SYMTUZA (dolutegravir/abacavir/lamivudine)	ATRIPLA (efavirenz/emtricitabine/tenofovir) efavirenz/lamivudine/tenofovir (generic for Symfi) ^{QL} efavirenz/lamivudine/tenofovir (generic for Symfi Lo) ^{QL} JULUCA (dolutegravir/rilpivirine) ^{QL} TRIUMEQ PD (abacavir, dolutegravir, and lamivudine) SUSP ^{NR}	 Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy Diagnosis of HIV/AIDS required OR Pre and Post Exposure Prophylaxis

with Prior Authorization Criteria

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HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

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

with Prior Authorization Criteria

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HYPOGLYCEMICS. INCRETIN MIMETICS/ENHANCERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
·	CEPTOR AGONIST (GLP-1 RA) ^{CL}	GLP-1 RA Criteria
OZEMPIC (semaglutide) TRULICITY (dulaglutide) VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) BYDUREON BCISE PEN (exenatide) ^{QL} BYDUREON (exenatide ER) BYDUREON PEN (exenatide ER) subcutaneous BYETTA (exenatide) subcutaneous MOUNJARO (tirazepatide) ^{NR} PEN RYBELSUS (semaglutide)	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR A diagnosis of ASCVD associated with a diagnosis of Type II diabetes (no metformin trial required) Non-preferred agents will be approved for patients who have: Failed a trial of TWO preferred agents within GLP-1 RA
INSULIN/GLP-1 RA	A COMBINATIONS	AND Diagnosis of disperses with HhA1C
	SOLIQUA (insulin glargine/lixisenatide) XULTOPHY (insulin degludec/liraglutide)	 Diagnosis of diabetes with HbA1C ≥ 7 AND Trial of metformin, or contraindication or intolerance to metformin
AMYLIN	ANALOG	Amylin Analog Criteria
	SYMLIN (pramlintide) subcutaneous	 ALL criteria must be met Concurrent use of short-acting mealtime insulin Current therapy compliance No diagnosis of gastroparesis HbA1C ≤ 9% within last 90 days Monitoring of glucose during initiation of therapy
DIPEPTIDYL PEPTIDASI	E-4 (DPP-4) INHIBITOR ^{QL}	- Illitiation of therapy
JANUMET (sitagliptin/metformin) JANUMET XR (sitagliptin/metformin) JANUVIA (sitagliptin) JENTADUETO (linagliptin/metformin) TRADJENTA (linagliptin)	alogliptin (generic for Nesina) alogliptin/metformin (generic for	DPP-4 Inhibitor Criteria Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin. Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within the DPP-4 inhibitor class

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

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HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HUMALOG (insulin lispro) U-100 CARTRIDGE, PEN, VIAL HUMALOG JR. (insulin lispro) U-100 KWIKPEN HUMALOG MIX VIAL (insulin lispro/lispro protamine) HUMALOG MIX KWIKPEN (insulin lispro/lispro protamine) HUMULIN (insulin) VIAL HUMULIN 70/30 VIAL HUMULIN U-500 VIAL HUMULIN OTC PEN HUMULIN OTC PEN HUMULIN 70/30 OTC PEN insulin aspart (generic for Novolog) insulin aspart/insulin aspart protamine PEN, VIAL (generic for Novolog Mix) insulin lispro (generic for Humalog) PEN, VIAL, JR KWIKPEN insulin lispro/lispro protamine KWIKPEN (Humalog Mix Kwikpen) LANTUS SOLOSTAR PEN (insulin glargine) LANTUS (insulin glargine) VIAL LEVEMIR (insulin detemir) PEN, VIAL NOVOLIN (insulin) PEN NOVOLOG (insulin aspart) CARTRIDGE, FLEXPEN, VIAL (insulin aspart/aspart protamine)	ADMELOG (insulin lispro) PEN, VIAL AFREZZA (regular insulin) INHALATION APIDRA (insulin glulisine) BASAGLAR (insulin glargine, rec) PEN FIASP (insulin aspart) CARTRIDGE, PEN, VIAL HUMALOG (insulin lispro) U-200 KWIKPEN insulin degludec (generic Tresiba) ^{NR} 100U/mL PEN, 200U/mL PEN, VIAL insulin glargine PEN, VIAL insulin Glargine-YFGN PEN, VIAL (generic for Semglee-YFGN) LYUMJEV KWIKPEN, VIAL(insulin lispro-aabc) NOVOLIN (insulin) NOVOLIN 70/30 VIAL(insulin) NOVOLOG MIX (insulin aspart/aspart protamine) VIAL TOUJEO SOLOSTAR (insulin glargine) SEMGLEE (insulin glargine) PEN, VIAL SEMGLEE YFGN (insulin glargine) PEN, VIAL TRESIBA (insulin degludec)	 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:

HYPOGLYCEMICS, MEGLITINIDES

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

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AL – Age Limit

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HYPOGLYCEMICS, METFORMINS

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

HYPOGLYCEMICS, SGLT2

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) ^{QL,CL} INVOKAMET (canagliflozin/metformin) ^{QL,CL} INVOKANA (canagliflozin) ^{CL} JARDIANCE (empagliflozin) ^{QL,CL} SYNJARDY (empagliflozin/metformin) ^{AL,CL,QL} XIGDUO XR (dapagliflozin/metformin) ^{QL,CL}	INVOKAMET XR (canagliflozin/metformin) ^{QL} SEGLUROMET (ertugliflozin/metformin) ^{QL} STEGLATRO (ertugliflozin) ^{QL} SYNJARDY XR (empagliflozin/metformin) ^{AL,QL}	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR A diagnosis of ASCVD or Heart Failure, or Chronic Kidney Disease associated with a diagnosis of Type II diabetes (no metformin trial required) Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class Drug Specific Criteria: Farxiga: May be approved for a diagnosis of heart failure with reduced ejection fraction (NYHA class II-IV) without a diagnosis of diabetes May be approved for a diagnosis of chronic kidney disease at risk of progression without a diagnosis of diabetes Jardiance: May be approved for a diagnosis of diabetes

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HYPOGLYCEMICS, SULFONYLUREAS

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

IDIOPATHIC PULMONARY FIBROSIS

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

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IMMUNOMODUI ATORS ASTHMACL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ASENRA (benralizumab) ^{AL} PEN (OLAIR (omalizumab) SYR ^{AL,QL}	NUCALA (mepolizumab) ^{AL} AUTO-INJ, SYR	Asthma Immunomodulator PA Form Non-preferred agents require a tria of a preferred agent within this drug class with the same indication Drug Specific Criteria: Dupixent: is indicated for Patients 6 years and older as an addomaintenance treatment in patients with moderate-to-severe asthma with an eosinophilic phenotype or with oral corticosteroid dependent asthma For other indications, see Immunomodulators, Atopic Dermatitis Fasenra: is indicated for Patient 12 years and older for addon maintenance treatment of severe asthma, and with an eosinophilic phenotype Nucala: is indicated for Patients 6 years and older for addon maintenance treatment of severe asthma, and with an eosinophilic phenotype Patients 12 years and older with hypereosinophilic syndrome (HES) for ≥6 months without identifiable non-hematologic secondary cause Patients 18 years and older for addomaintenance treatment of chronic rhinosinusitis with nasal polyps (CRWSwNP) with inadequate response to nasal corticosteroids. Adult patients with eosinophilic granulomatosis with polyangiitic syndrome in vitro reactivity to a perennial aeroallergen and symptoms that are inadequately controlled with inhaled corticosteroids Patients 12 years and older with Chronic spontaneous urticaria (CSU) who remain symptomatic despite H1 antihistamine treatmen Patients 18 years and older with Nasa Polyps with inadequate response to nasal corticosteroids. As add-on maintenance treatment

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IMMUNOMODULATORS, ATOPIC DERMATITISAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
EUCRISA (crisaborole) ^{CL,QL}	ADBRY (tralokinumab-ldrm) SUB-Q ^{AL,NR,QL} DUPIXENT (dupilumab) ^{AL,CL} DUPIXENT PEN ^{AL} OPZELURA (ruxolitinib phosphate) CREAM ^{AL,NR,QL} pimecrolimus (generic for Elidel) tacrolimus (generic for Protopic) ^{CL}	Non-preferred agents require: Trial of a topical steroid AND Trial of one preferred product within this drug class Drug-specific criteria: Dupixent: Indicated for the treatment of patients aged 6 months and older with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. DUPIXENT can be used with or without topical corticosteroids. -as an add-on maintenance treatment of patients aged 6 years and older with moderate-to-severe asthma characterized by an eosinophilic phenotype or with oral corticosteroid dependent asthma. - as an add-on maintenance treatment in adult patients with inadequately controlled chronic rhinosinusitis with nasal polyposis (CRSwNP) - for treatment of eosinophilic esophagitis in adult and pediatric patients aged 12 years and older, weighing at least 40 kg • Eucrisa: Requires use and failure of 1 topical steroid or Elidel.

IMMUNOMODULATORS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
imiquimod (generic for Aldara)	ALDARA (imiquimod) HYFTOR (sirolimus) ^{AL,NR} GEL 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

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IMMUNOSUPPRESSIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathioprine (generic Imuran) azathioprine (generic Azasan) ^{NR} cyclosporine, modified CAPS (generic Neoral) everolimus (generic for Zortress) ^{AL} mycophenolate CAPS, TABLET (generic Cellcept) RAPAMUNE (sirolimus) SOLN RAPAMUNE (sirolimus) TABLET tacrolimus	ASTAGRAF XL (tacrolimus) AZASAN (azathioprine) cyclosporine CAP, SOFTGEL cyclosporine, modified SOLN (generic Neoral) ENVARSUS XR (tacrolimus) GENGRAF (cyclosporine, modified) CAP, SOLN mycophenolate SUSP (generic Cellcept) mycophenolic acid MYFORTIC (mycophenolate sodium) PROGRAF (tacrolimus) CAPS, PACKET REZUROCK (belumosudil) ^{AL,QL} TAB SANDIMMUNE (cyclosporine) CAPS, SOLN sirolimus SOLN, TABLET (generic Rapamune) TAVNEOS (avacopan) ^{QL} CAPS ZORTRESS (everolimus) AL	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Patients established on existing therapy will be allowed to continue

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INTRANASAL RHINITIS DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTICHOLINERGICS		Non-preferred agents will be approved
ipratropium (generic for Atrovent)		for patients who have failed a 30-day trial of ONE preferred agent within this
ANTIHIS	TAMINES	drug class
azelastine 0.1% (generic for Astelin)	azelastine 0.15% (generic for Astepro) azelastine/fluticasone (generic for Dymista) olopatadine (generic for Patanase) RYALTRIS (olopatadine/mometasone)AL,NR	 Drug-specific criteria: mometasone: Prior authorization NOT required for children ≤ 12 years budesonide: Approved for use in Pregnancy (Pregnancy Category
CORTICOSTEROIDS		 B) Xhance: Indicated for treatment of
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) XHANCE (fluticasone) ZETONNA (ciclesonide)	nasal polyps in ≥ 18 years only

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LEUKOTRIENE MODIFIERS

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

LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

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

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LIPOTROPICS, OTHER

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

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LIPOTROPICS, OTHER (continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROPROTEIN CONVERTASE SU	BTILISIN/KEXIN TYPE 9 (PCSK9) BITORS PRALUENT (alorocumab) ^{CL} REPATHA (evolocumab) ^{CL}	 Prior Authorization/Class Criteria Praluent®: Approved for diagnoses of: atherosclerotic cardiovascular disease (ASCVD) heterozygous familial hypercholesterolemia (HeFH) Homozygous familial hypercholesterolemia (HoFH) as an adjunct to other LDL-C lowering therapies AND Maximized high-intensity statin WITH ezetimibe for at 3 continuous months Failure to reach target LDL-C levels: ASCVD - < 70 mg/dL, HeFH - < 100 mg/dL Repatha®: Approved for: adult diagnoses of atherosclerotic cardiovascular disease (ASCVD) heterozygous familial hypercholesterolemia (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:
		 Maximized high-intensity statin WITH ezetimibe for 3+ continuous months

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LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atorvastatin (generic Lipitor) ^{QL} lovastatin (generic Mevacor) pravastatin (generic Pravachol) rosuvastatin (generic Crestor) simvastatin (generic Zocor)	ALTOPREV (lovastatin ER) ^{CL} EZALLOR SPRINKLE (rosuvastatin) ^{QL} fluvastatin IR/ER (generic Lescol/ Lescol XL) LIVALO (pitavastatin) 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
STATIN COI	MBINATIONS	IR lovastatin Combination products: Require clinical reason why individual ingredients cannot be
	atorvastatin/amlodipine (generic Caduet) simvastatin/ezetimibe (generic Vytorin)	 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

MACROLIDES AND KETOLIDES, ORAL

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

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METHOTREXATE

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
methotrexate PF VIAL, TABLET, VIAL	OTREXUP (methotrexate) SUB-Q RASUVO (methotrexate) SUB-Q REDITREX	Non-preferred agents require a trial of the preferred agent AND will be approved for an FDA-approved indication Drug-specific criteria: ■ Xatmep™:Indicated for pediatric patients only

MOVEMENT DISORDERS

FPreferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AUSTEDO (deutetrabenazine) ^{CL} INGREZZA (valbenazine) ^{AL,CLQL} CAPS	INGREZZA (valbenazine) ^{CL} INITIATION PACK XENAZINE (tetrabenazine) ^{CL}	All drugs require an FDA approved indication – ICD-10 diagnosis code required.
tetrabenazine (generic for Xenazine) ^{CL}		Non-preferred agents require a trial and failure of a preferred agent with the same indication or a clinical reason why a preferred agent in this class cannot be used. 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 • tetrabenazine: Diagnosis of chorea with Huntington's Disease

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MULTIPLE SCLEROSIS DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AVONEX (interferon beta-1a) ^{QL} BETASERON (interferon beta-1b) ^{QL} COPAXONE 20mg (glatiramer) ^{QL} dimethyl fumarate (generic for Tecfidera) KESIMPTA (Ofatumumab) ^{CL,QL}	AUBAGIO (teriflunomide) BAFIERTAM (monomethyl fumarate) ^{QL} dalfampridine (generic Ampyra) ^{QL} EXTAVIA (interferon beta-1b) ^{QL} GILENYA (fingolimod) ^{QL} glatiramer (generic Copaxone) ^{QL} MAVENCLAD (cladribine) MAYZENT (siponimod) ^{QL} PLEGRIDY (peginterferon beta-1a) ^{QL} PONVORY (ponesimod) REBIF (interferon beta-1a) ^{QL} TASCENSO ODT (fingolimod) TABLET ^{AL,NR} TECFIDERA (dimethyl fumarate) VUMERITY (diroximel) ^{QL} ZEPOSIA (ozanimod) ^{AL,CL,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 Kesimpta: Approved for patients who have failed a trial of a preferred injectable agent within this class Zeposia: Approved for a diagnosis of relapsing forms of multiple sclerosis (MS) with trial of ONE preferred agent OR a diagnosis of moderately to severely active ulcerative colitis and treatment failure of Humira.

NITROFURAN DERIVATIVES

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

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NSAIDs, ORAL

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

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NSAIDs, ORAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COX-I SELECT	IVE (continued)	
NSAID/GI PROTECT	ALL BRAND NAME NSAIDs including: CAMBIA (diclofenac oral solution) DUEXIS (ibuprofen/famotidine) ^{CL} ibuprofen/famotidine (generic Duexis) ^{CL} SPRIX (ketorolac nasal spray) NASAL ^{QL, CL} TIVORBEX (indomethacin) VIVLODEX (meloxicam submicronized) ZIPSOR (diclofenac) ZORVOLEX (diclofenac) ANT COMBINATIONS diclofenac/misoprostol (generic for	Drug-specific criteria: Sprix®: Approved for patients unable to tolerate, swallow OR absorb oral NSAIDs OR contraindication OR trial of TWO preferred oral NSAIDs Tivorbex®: Requires clinical reason why indomethacin capsules cannot be used Zorvolex®: Requires trial of oral diclofenac OR clinical reason why diclofenac potassium/sodium cannot be used
	Arthrotec)	
COX-II SI	ELECTIVE	
celecoxib (generic for Celebrex)		

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NSAIDs, TOPICAL

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

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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 require a trial of a preferred agent,
IBRANCE (palbociclib)	KISQALI (ribociclib) KISQALI FEMARA CO-PACK VERZENIO (abemaciclib)	but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
CHEMOT	HERAPY	- 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
ОТІ	HER	for short term use
	NERLYNX (neratinib) PIQRAY (alpelisib) lapatinib (generic Tykerb) ^{CL} TALZENNA (talazoparib tosylate) QL TUKYSA(tucatinib) ^{QL}	 Soltamox: May be approved with documented swallowing difficulty

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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	PURIXAN (mercaptopurine) ^{AL}	 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
	AML	from current treatment guidelines
IMBRUVICA (ibrutinib) LEUKERAN (chlorambucil) VENCLEXTA (venetoclax)	DAURISMO (glasdegib maleate) ^{QL} IDHIFA (enasidenib) RYDAPT (midostaurin) TIBSOVO (ivosidenib) ^{QL} XOSPATA (gilteritinib) ^{QL} CLL COPIKTRA (duvelisib) ^{QL} IMBRUVICA (ibrutinib) ^{NR} SUSP 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 Purixan: Prior authorization not required for age ≤12 or for documented swallowing disorder Tabloid: Prior authorization not required for age ≤10
	CML	required for age <19 Tasigna: Patients receiving
hydroxyurea (generic for Hydrea) imatinib (generic for Gleevec) MYLERAN (busulfan) SPRYCEL (dasatinib)	BOSULIF (bosutinib) GLEEVEC (imatinib) HYDREA (hydroxyurea) ICLUSIG (ponatinib) SCEMBLIX (asciminib) ^{NR} TASIGNA (nilotinib) ^{CL}	 Tasigna, which changed from preferred to non-preferred on 1-17-19 will be allowed to continue therapy Xpovio: Indicated for relapsed or refractory multiple myeloma. Requires concomitant therapy with
MPN		dexamethasone
JAKAFI (ruxolitinib)		
M	YELOMA	
ALKERAN (melphalan) REVLIMID ^{QL} (lenalidomide)	FARYDAK (panobinostat) lenalidomide ^{NR,QL} (generic for Revlimid) melphalan (generic for Alkeran) NINLARO (ixazomib) POMALYST (pomalidomide) THALOMID (thalidomide) XPOVIO (selinexor) CL	
	OTHER	
MATULANE (procarbazine) TABLOID (thioguanine) tretinoin (generic for Vesanoid) ^{AL}	BRUKINSA (zanubrutinib ^{QL} CALQUENCE (acalabrutinib) ^{QL} INREBIC (fedratinib dihydrochloride) ^{QL} INQOVI (decitabine/cedazuridine) VONJO (pacritinib) ^{NR,QL} ZOLINZA (vorinostat)	

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

with Prior Authorization Criteria

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ONCOLOGY AGENTS, ORAL, LUNG

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALECENSA (alectinib)	ALK ALUNBRIG (brigatinib) ^{QL} LORBRENA (lorlatinib) ^{QL} ZYKADIA (ceritinib) CAPS , TABLET	 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 Criteria Iressa/ Xalkori: Patients receiving Iressa or Xalkori prior to 1/21/21 (which changed from preferred to non-preferred) will be allowed to continue current treatment
ALK / ROS1 / NTRK		
	ROZLYTREK (entrectinib) ^{AL,QL} XALKORI (crizotinib)	
EGFR		
TAGRISSO (osimertinib)	erlotinib (generic for Tarceva) EXKIVITY (mobocertinib) ^{NR,QL} GILOTRIF (afatinib) IRESSA (gefitinib) TARCEVA (erlotinib) VIZIMPRO (dacomitinib) ^{QL}	
C	THER	
	GAVRETO (pralsetinib) ^{QL} HYCAMTIN (topotecan) LUMAKRAS (sotrasib) ^{QL} RETEVMO (selpercatinib) ^{AL} TABRECTA (capmatinib) ^{QL} TEPMETKO (tepotinib) ^{QL}	

with Prior Authorization Criteria

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ONCOLOGY AGENTS, ORAL, OTHER

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

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ONCOLOGY AGENTS, ORAL, PROSTATE

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
abiraterone (generic for Zytiga) ^{AL,QL} bicalutamide (generic for Casodex) flutamide XTANDI (enzalutamide) ^{AL,QL} ZYTIGA (abiraterone) ^{AL,QL}	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
INLYTA (axitinib) LENVIMA (lenvatinib) SUTENT (sunitinib) VOTRIENT (pazopanib	AFINITOR DISPERZ (everolimus) ^{CL} CABOMETYX (cabozantinib) everolimus (generic for Afinitor) everolimus SUSP (generic for Afinitor Disperz) ^{NR} FOTIVDA (tivozanib) ^{NR} NEXAVAR (sorafenib) sorafenib (generic Nexavar) ^{NR} sunitinib malate (generic for Sutent) WELIREG (belzutifan) ^{NR,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 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 CELL ODOMZO (coniderib)		 Non-preferred agents DO NOT require a trial of a preferred agent,
ERIVEDGE (vismodegib)	ODOMZO (sonidegib) ^{CL}	but DO require an FDA-approved indication OR documentation submitted supporting off-label use from current treatment guidelines
BRAF MUTATION		
MEKINIST (trametinib)	BRAFTOVI (encorafenib)	Drug-specific critera
TAFINLAR (dabrafenib)	COTELLIC (cobimetinib) MEKTOVI (binimetinib) ZELBORAF (vemurafenib)	 Odomzo: Patients receiving Odomzo, which changed from preferred to non-preferred on 1-17- 19 will be allowed to continue therapy

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

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OPHTHALMICS, ALLERGIC CONJUNCTIVITIS

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

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OPHTHALMICS, ANTIBIOTICS

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

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OPHTHALMICS, ANTIBIOTIC-STEROID COMBINATIONS

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

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OPHTHALMICS, ANTI-INFLAMMATORIES

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

OPHTHALMICS, ANTI-INFLAMMATORY / IMMUNOMODULATORS

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

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AL – Age Limit

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OPHTHALMICS, GLAUCOMA

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MIO	rics	 Non-preferred agents will be
pilocarpine	PHOSPHOLINE IODIDE (echothiophate iodide) VUITY (pilocarpine) ^{NR}	approved for patients who have failed a trial of ONE preferred agent within this drug class
SYMPATHO	MIMETICS	<u> </u>
Alphagan P (brimonidine 0.15%) brimonidine 0.2% (generic for Alphagan)	Alphagan P (brimonidine 0.1%) apraclonidine (generic for lopidine) brimonidine P 0.15%	
BETA BLO	OCKERS	<u>[</u>
levobunolol (generic for Betagan) timolol (generic for Timoptic)	betaxolol (generic for Betoptic) BETIMOL (timolol) BETOPTIC S (betaxolol) carteolol (generic for Ocupress) timolol (generic for Istalol) timolol (generic for Timoptic Ocudose) TIMOPTIC OCUDOSE TIMOPTIC XE (timolol gel forming solution)	
CARBONIC ANHYDR	RASE INHIBITORS	
. ,	AZOPT (brinzolamide) brinzolamide (generic for Azopt)	
PROSTAGLAND	IN ANALOGS	
latanoprost (generic for Xalatan) TRAVATAN Z (travoprost)	bimatoprost (generic for Lumigan) travoprost (generic for Travatan Z) VYZULTA (latanoprostene) XALATAN (latanoprost) ZIOPTAN (tafluprost)	
COMBINATIO	ON DRUGS	
COMBIGAN (brimonidine/timolol) dorzolamide/timolol (generic for Cosopt)	brimonidine/timolol (generic Combigan) ^{NR} dorzolamide/timolol PF (generic for Cosopt PF) SIMBRINZA (brinzolamide/brimonidine)	
ОТІ	IER	
RHOPRESSA (netarsudil) ^{CL} ROCKLATAN (netarsudil and latanoprost) ^{CL}		Drug-specific criteria: Rhopressa and Rocklatan: Electronically approved for patients who have a trial of ONE generic agent, within ophthalmics - glaucoma within 60 days

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OPIOID DEPENDENCE TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
buprenorphine SL buprenorphine/naloxone TAB (SL) SUBOXONE FILM (buprenorphine/naloxone)	buprenorphine/naloxone FILM LUCEMYRA (lofexidine) ^{CL,QL} ZUBSOLV (buprenorphine/naloxone)	Buprenorphine PA Form Buprenorphine Informed Consent Non-preferred agents require a treatment failure of a preferred drug or patient-specific documentation of why a preferred product is not appropriate for the patient. Drug-specific criteria: Lucemyra: Approved for FDA approved indication and dosing per label. Trial of preferred product not required.

OPIOID-REVERSAL TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naloxone SYRINGE, VIAL naltrexone TABLET NARCAN (naloxone) SPRAY	KLOXXADO (naloxone) NASAL naloxone SPRAY (generic for Narcan) ZIMHI (naloxone) ^{AL} SYRINGE	 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

Nebraska Medicaid Preferred Drug List with Prior Authorization Criteria

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OTIC ANTIBIOTICS

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

PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED

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

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CL – Prior Authorization / Class Criteria apply

QL – Quantity/Duration Limit

AL – Age Limit

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

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PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD CHEW + IRON (MULTIVITAMIN WITH IRON) CHEW CHILDREN'S CHEWABLES (PEDI MULTIVIT NO.31/IRON/FOLIC, PEDI MULTIVIT NO.25/FOLIC ACID, PEDI MULTIVIT NO.23/FOLIC ACID) MULTIVIT-FLUOR (PEDI MULTIVIT NO.17 W-FLUORIDE, PEDI MULTIVIT NO.16 W-FLUORIDE) CHEW MULTIVIT-FLUOR (PEDI MULTIVIT NO.2 W-FLUORIDE) DROP MULTIVIT-IRON-FLUOR (PEDI MULTIVIT NO.2 W-FLUORIDE) DROP	Non-Preferred Agents DEKAS PLUS LIQUID ^{AL} FLORIVA (PEDI MULTIVIT NO.85/FLUORIDE) CHEW FLORIVA PLUS (PEDI MULTIVIT NO.161/FLUORIDE DROP MULTI-VIT-FLOR (PEDI MULTIVIT NO.205/FLUORIDE) CHEW POLY-VI-FLOR (PEDI MULTIVIT NO.33/FLUORIDE) CHEW POLY-VI-FLOR (PEDI MULTIVIT NO.37 W-FLUORIDE) DROPS POLY-VI-FLOR /0.25mg IRON (PEDI MULTIVIT 37/FLUORIDE/IRON) POLY-VI-FLOR /0.5mg IRON (PEDI MULTIVIT 33/FLUORIDE/IRON) POLY-VI-SOL (PEDIATRIC MULTIVITAMIN NO.192) DROP QUFLORA (PEDI MULTIVIT NO.157/FLUORIDE) GUMMIES	Prior Authorization/Class Criteria Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class Drug specific criteria: DEKAS Plus: Approved for diagnosis of Cystic Fibrosis and does not require a trial of a preferred agent
DROPS TRI-VI-SOL (VIT A PALMITATE/VIT C/VIT D3) DROPS	NO.157/FLUORIDE) GUMMIES QUFLORA FE (PED MULTIVIT 142/IRON/FLUORIDE) CHEW QUFLORA FE (PED MULTIVIT 151/IRON/FLUORIDE) DROP	
	QUFLORA PED (PEDI MULTIVIT NO.63 W-FLUORIDE) CHEW QUFLORA PED (PEDI MULTIVIT 84 WITH FLUORIDE, PEDI MULTIVIT NO.83 W-FLUORIDE) DROP TRI-VI-FLOR (PED MVIT A,C,D3 NO.38/FLUORIDE) DROPS	

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PENICILLINS

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

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AL – Age Limit

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PRENATAL VITAMINS

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COMPLETENATE CHEW TABLET EXPECTA PRENATAL OTC FE C/FA FE C/VIT C/VIT B12/FA OTC MARNATAL-F O-CAL FA PNV2/IRON B-G SUC-P/FA/OMEGA-3 PNV 11-IRON FUM-FOLIC ACID-OM3 PNV NO.118/IRON FUMARATE/FA CHEW TABLET PNV NO.15/IRON FUM & PS CMP/FA PNV W-CA NO.40/IRON FUM/FA CMB NO.1 PNV WITH CA NO.68/IRON/FA NO.1/DHA PNV WITH CA,NO.72/IRON/FA PNV#16/IRON FUM & PS/FA/OM-3 PNV2/IRON B-G SUC-P/FA/OMEGA-3 PRENATAL VIT #76/IRON,CARB/FA PRENATAL VIT NO.78/IRON/FA PRENATAL VIT/FE FUMARATE/FA OTC PUREFE OB PLUS PUREFE PLUS STUART ONE OTC TRINATAL RX 1 VITAFOL TAB CHEW VITAFOL ULTRA VP-PNV-DHA	CITRANATAL B-CALM COMPLETENATE CHEW TABLET DERMACINRX PRENATRIX OTC DERMACINRX PRETRATE TABLET OTC ENBRACE HR MULTI-MAC OTC NESTABS NESTABS ABC NESTABS ONE OB COMPLETE ONE OB COMPLETE PETITE OB COMPLETE PREMIER OB COMPLETE TABLET OB COMPLETE WITH DHA PNV COMBO#47/IRON/FA #1/DHA PNV WITH CA,NO.72/IRON/FA PNV WITH CA,NO.74/IRON/FA OTC PNV119/IRON FUMARATE/FA/DSS PRENATAL H DHA OTC PRENATE AM PRENATE CHEWABLE TABLET PRENATE CHEWABLE TABLET PRENATE ELITE PRENATE ENHANCE PRENATE ESSENTIAL PRENATE FIXIE PRENATE STAR PRIMACARE SELECT-OB + DHA SELECT-OB TAB CHEW TENDERA-OB OTC TRICARE TRINATAL RX 1 TRISTART DHA VITAFOL NANO VITAFOL-OB VITAFOL-OB VITAFOL-ONE WESTGEL 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

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AL – Age Limit

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PROGESTERONE (hydroxyprogesterone caproate)

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

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
omeprazole (generic Prilosec) RX contoprazole (generic Protonix) ^{QL} PROTONIX SUSP (pantoprazole)	DEXILANT (dexlansoprazole) dexlansoprazole (generic Dexilant) esomeprazole magnesium (generic Nexium) RX ^{QL} esomeprazole magnesium (generic Nexium) OTC ^{QL} esomeprazole strontium lansoprazole (generic Prevacid) ^{QL} NEXIUM SUSPENSION (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) pantoprazole GRANULES ^{QL} rabeprazole (generic Aciphex)	 Non-preferred agents will be approved for patients who have failed an 8-week trial of both preferred omeprazole Rx AND pantoprazole OR Protonix SUSP. 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 of age- Only approve non-preferred for Gl diagnosis if:

with Prior Authorization Criteria

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SEDATIVE HYPNOTICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BENZODIA temazepam 15mg, 30mg (generic for Restoril) OTH zaleplon (generic for Sonata) zolpidem (generic for Ambien)	estazolam (generic for ProSom) flurazepam (generic for Dalmane) temazepam (generic for Restoril) 7.5mg, 22.5mg triazolam (generic for Halcion)	 Lunesta®/ Rozerem®/zolpidem ER: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used Edluar®: Requires a trial with generic zolpidem within the last 12 months AND Trial OR Clinical reason why zaleplon and preferred benzodiapine cannot be used and Requires documentation of swallowing disorder flurazepam/triazolam: Requires trial of preferred benzodiazepine Hetlioz®: Requires trial with generic zolpidem within last 12 months AND clinical reason why zaleplon AND preferred benzodiazepine cannot be used Silenor®: Must meet ONE of the following: Contraindication to preferred oral sedative hypnotics Medical necessity for doxepin dose < 10mg Age greater than 65 years old or hepatic impairment (3mg dose will be approved if this criteria is met) temazepam 7.5mg/22.5mg: Requires clinical reason why 15mg/30mg cannot be used zolpidem/zolpidem ER: Maximum daily dose for females: Zolpidem 5mg; Zolpidem ER® 6.25mg zolpidem SL: Requires clinical reason why half of zolpidem tablet cannot be used

with Prior Authorization Criteria

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SICKLE CELL ANEMIA TREATMENT^{AL}

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

SINUS NODE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORLANOR SOLN , 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

with Prior Authorization Criteria

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SKELETAL MUSCLE RELAXANTS

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

with Prior Authorization Criteria

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STEROIDS TOPICAL

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

with Prior Authorization Criteria

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STEROIDS, TOPICAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HIGH POTENCY		 High Potency Non-preferred
triamcinolone acetonide OINTMENT, CREAM triamcinolone LOTION	amcinonide CREAM, LOTION, OINTMENT betamethasone dipropionate	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class
tramonoro 2011014	betamethasone / propylene glycol betamethasone valerate	drug class
	desoximetasone diflorasone diacetate fluocinonide SOLN	
	fluocinonide CREAM, GEL, OINT fluocinonide emollient	
	halcinonide CREAM (generic for Halog)	
	HALOG (halcinonide) CREAM, OINT, SOLN	
	KENALOG AEROSOL (triamcinolone) SERNIVO (betamethasone	
	dipropionate) triamcinolone SPRAY (generic for	
	Kenalog spray) TRIANEX OINT (triamcinolone)	
	VANOS (fluocinonide)	
VERY HIGH	H POTENCY	 Very High Potency Non-preferred
clobetasol emollient (generic for Temovate-E) clobetasol propionate CREAM, GEL, OINT, SOLN	APEXICON-E (diflorasone) BRYHALI (halobetasol prop) LOTION clobetasol SHAMPOO, LOTION	agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class
halobetasol propionate (generic for Ultravate)	clobetasol propionate FOAM , SPRAY CLOBEX (clobetasol)	
c.a.a.a.c,	halobetasol propionate FOAM (generic for Lexette) AL,QL	
	IMPEKLO (clobetasol) LOTION ^{AL} LEXETTE(halobetasol propionate) AL,QL OLUX-E /OLUX/OLUX-E CP	
	(clobetasol)	

with Prior Authorization Criteria

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STIMULANTS AND RELATED AGENTS^{AL}

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CNS STI	MULANTS mine type ADZENYS XR (amphetamine) amphetamine ER (generic Adzenys ER) SUSP amphetamine salt combination ER (generic for Adderall XR) amphetamine sulfate (generic for Evekeo) dextroamphetamine (generic for Dexedrine) dextroamphetamine ER (generic For Dexedrine) dextroamphetamine ER (generic for Dexedrine) DYANAVEL XR (amphetamine) EVEKEO ODT (amphetamine sulfate) MYDAYIS (amphetamine salt combo) MYDAYIS (dextroamphetamine) ZENZEDI (dextroamphetamine)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Procentra®: May be approved with documentation of swallowing disorder Zenzedi®: Requires clinical reason

with Prior Authorization Criteria

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STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Methylpho	enidate type	 Non-preferred agents will be approved for patients who have
METHYLIN SOLN (methylphenidate) methylphenidate (generic for Ritalin) methylphenidate SOLN (generic for Methylin) QUILLICHEW ER CHEWTAB (methylphenidate)	ADHANSIA XR (methylphenidate) QL APTENSIO XR (methylphenidate) AZSTARYS (serdexmethylphenidate and dexmethylphenidate) QL 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 Ritalin LA) methylphenidate 30/70 (generic for Metadate CD) methylphenidate ER 18mg, 27mg,	 failed a trial of TWO preferred agents within this drug class Maximum accumulated dose of 108mg per day for ages < 18 Maximum accumulated dose of 72mg per day for ages > 19

with Prior Authorization Criteria

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STIMULANTS AND RELATED ADHD DRUGS (Continued)AL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MISCELLANEOUS		Note: generic guanfacine IR and
guanfacine ER (generic for Intuniv)QL	clonidine ER (generic for Kapvay) ^{QL} QELBREE (viloxazine) ^{QL} STRATTERA (atomoxetine)	–clonidine IR are available without prior authorization
		Drug-specific criteria: armodafinil and Sunosi: Require
ANALEF	PTICS	trial of modafinil
	armodafinil (generic for Nuvigil) ^{CL}	armodafinil and modafinil: approved only for:
\$	modafanil (generic for Provigil) ^{CL} SUNOSI (solriamfetol) ^{CL,QL} WAKIX (pitolisant) ^{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

with Prior Authorization Criteria

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TETRACYCLINES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR (generic Vibramycin) doxycycline monohydrate 50MG, 100MG CAPS doxycycline monohydrate SUSP, TABLET (generic Vibramycin) minocycline HCI CAPS, 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 CAP (generic 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 a 3-day trial of TWO preferred agents within this drug class Drug-specific criteria: Demeclocycline: Approved for diagnosis of SIADH doxycycline suspension: May be approved with documented swallowing difficulty

THROMBOPOIESIS STIMULATING PROTEINSCL

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

with Prior Authorization Criteria

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THYROID HORMONES

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

with Prior Authorization Criteria

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ULCERATIVE COLITIS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine) LIALDA (mesalamine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) GIAZO (balsalazide) mesalamine ER (generic Apriso) mesalamine ER (generic Pentasa) ^{NR} mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Asacol HD®/Delzicol DR®/Pentasa®: Requires clinical reason why preferred mesalamine products cannot be used Giazo®: Requires clinical reason why generic balsalazide cannot be used
CANASA (mesalamine) ROWASA (mesalamine)	mesalamine ENEMA (generic Rowasa) mesalamine SUPPOSITORY (generic Canasa) UCERIS (budesonide)	NOT covered in females

UTERINE DISORDER TREATMENT

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

Nebraska Medicaid **Preferred Drug List** with Prior Authorization Criteria

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VASODILATORS, CORONARY

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate TABLET isosorbide dinitrate ER, SA TABLET (generic Dilatrate-SR/Isordil) isosorbide mono IR/SR TABLET nitroglycerin SUBLINGUAL, TRANSDERMAL nitroglycerin ER TABLET	BIDIL (isosorbide dinitrate/ hydralazine) ^{CL} GONITRO (nitroglycerin) isosorbide dinitrate TABLET (Oceanside Pharm MFR only) isosorbide dinitrate/hydralazine (Bidil) ^{CL,NR} NITRO-BID OINT (nitroglycerin) NITRO-DUR (nitroglycerin) nitroglycerin TRANSLINGUAL (generic Nitrolingual) VERQUVO (vericiguat) ^{AL,CL,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: BiDil: Approved for the treatment of heart failure as an adjunct therapy to standard therapy in self-identified black patients Verquvo: Approved for use in patients following a recent hospitalization for HF within the past 6 months OR need for outpatient IV diuretics, in adults with symptomatic chronic HF and EF less than 45%