

General Preferred Drug List Information

- Gainwell Technologies' DUR+ process is a proprietary electronic prior authorization system used for Medicaid pharmacy claims. MSCAN plans may/may not have electronic PA functionality. However, they must adhere to Medicaid's PA criteria.
- Drug coverage subject to the rules and regulations set forth in Sec. 1927 of Social Security Act. This is not an all-inclusive list of available covered drugs and includes only managed categories. Unless otherwise stated, the listing of a particular brand or generic name includes all dosage forms of that drug. NR indicates a new drug that has not yet been reviewed by the P&T Committee.
- **PREFERRED BRANDS** will not count toward the two-brand monthly Rx Limit.
- Drugs highlighted in **yellow** denote change in PDL status.
- An * denotes existing users will be grandfathered; grandfathering is defined as approving a Non-Preferred agent for an existing user; all other changes will not qualify for grandfathering.
- A # denotes existing users will NOT be grandfathered.
- To search the PDL, **press CTRL + F**.

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|-------------------------------------|---|
| ACNE AGENTS | | |
| ANTI-INFECTIVES | | <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 21 years: all acne agents except isotretinoin products <p>Note:</p> <ul style="list-style-type: none"> • Isotretinoin products available for all ages |
| clindamycin gel (generic CLEOCIN-T) | azelaic acid | |
| clindamycin lotion, medicated swab, solution | CLEOCIN T (clindamycin) | |
| | CLINDACIN (clindamycin) | |
| | CLINDAGEL (clindamycin) | |
| | clindamycin foam | |
| | clindamycin gel (generic CLINDAGEL) | |
| | dapsone | |
| | ERY (erythromycin) | |
| | ERYGEL (erythromycin) | |
| | erythromycin | |
| | EVOCLIN (clindamycin) | |
| | KLARON (sulfacetamide) | |
| | MORGIDOX (doxycycline) | |
| | sulfacetamide sodium suspension | |
| | WINLEVI (clascoterone) cream | |
| ISOTRETINOIN PRODUCTS | | |
| AMNESTEEM (isotretinoin) | ABSORBICA (isotretinoin) | |
| CLARAVIS (isotretinoin) | isotretinoin | |
| ZENATANE (isotretinoin) | | |
| KERATOLYTICS (BENZOYL PEROXIDES) | | |
| ACNE MEDICATION (benzoyl peroxide) | BPO towelette (benzoyl peroxide) | |
| benzoyl peroxide | | |
| LINTERA (benzoyl peroxide) | | |
| RETINOIDS | | |
| adapalene gel, gel with pump | adapalene cream | |
| RETIN-A (tretinoin) | AKLIEF (trifarotene) | |
| tretinoin cream | ALTRENO (tretinoin) | |
| | ARAZLO (tazarotene) | |
| | ATRALIN (tretinoin) | |
| | DIFFERIN (adapalene) | |
| | FABIOR (tazarotene) | |
| | RETIN-A MICRO (tretinoin) | |
| | RETIN-A MICRO PUMP (tretinoin) | |
| | tretinoin gel | |
| | tretinoin microsphere | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|--|
| ACNE AGENTS (continued) | | |
| OTHERS/COMBINATION PRODUCTS | | <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 21 years: all acne agents except isotretinoin products |
| adapalene/benzoyl peroxide gel | ACANYA (benzoyl peroxide/clindamycin) gel | |
| clindamycin/benzoyl peroxide 1.2(1)%-5% gel | BENZAMYCIN (benzoyl peroxide/erythromycin) gel | |
| clindamycin/benzoyl peroxide 1%-5% gel w/pump | CABTREO (clindamycin/adapalene/benzoyl peroxide) gel | |
| sodium sulfacetamide w/sulfur 8%-4%, 9%-4.25%, 10-5% suspension | CLEANSING WASH (sulfacetamide sodium/sulfur/urea) cleanser | |
| | clindamycin phosphate/benzoyl peroxide 1.2%-2.5% gel | |
| | clindamycin phosphate/tretinoin 1.2%-0.025% gel | |
| | clindamycin/benzoyl peroxide 1%-5% gel | |
| | clindamycin/benzoyl peroxide 1.2%-3.75% gel w/pump (generic ONEXTON) | |
| | EPIDUO FORTE (adapalene/benzoyl peroxide) gel | |
| | erythromycin/benzoyl peroxide gel | |
| | NEUAC (benzoyl peroxide/clindamycin) cream, gel | |
| | ONEXTON (benzoyl peroxide/clindamycin) gel | |
| | sodium sulfacetamide w/sulfur 8%-4% cleanser | |
| | sodium sulfacetamide w/sulfur 10%-2% cream | |
| | sodium sulfacetamide w/sulfur 10%-5% cream, lotion | |
| | SSS (sodium sulfacetamide/sulfur)10-5 cream, foam | |
| | TWYNEO (benzoyl peroxide/tretinoin) cream | |
| | ZIANA (clindamycin/tretinoin) gel | |
| | ZMA CLEAR (sodium sulfacetamide/sulfur) suspension | |
| ALPHA-1 PROTEINASE INHIBITORS | | |
| ARALAST NP | | |
| GLASSIA | | |
| PROLASTIN C | | |
| ZEMAIRA | | |
| ALZHEIMER'S AGENTS ^{DUR+} | | |
| CHOLINESTERASE INHIBITORS | | <p>Preferred Criteria</p> <ul style="list-style-type: none"> • Documented approvable diagnosis <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented approvable diagnosis AND • Have tried 2 different preferred agents in the past 6 months <p>NAMZARIC</p> <ul style="list-style-type: none"> • Documented approvable diagnosis AND • 30 days of concurrent therapy with both donepezil and memantine in the past 6 months |
| donepezil 5 mg, 10 mg ODT, tablets | ADLARITY (donepezil) | |
| galantamine | ARICEPT (donepezil) | |
| galantamine ER | donepezil 23 mg tablet | |
| rivastigmine | EXELON (rivastigmine) | |
| NMDA RECEPTOR ANTAGONISTS | | |
| memantine | memantine ER | |
| | NAMENDA (memantine) | |
| | NAMENDA XR (memantine ER) | |
| COMBINATION AGENTS | | |
| | NAMZARIC (memantine/donepezil) | |
| | memantine/donepezil ER ^{NR} | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| ANALGESICS, OPIOID-SHORT ACTING DUR+ | | |
| acetaminophen/caffeine/dihydrocodeine | ACTIQ (fentanyl) | <p>MS DOM Opioid Initiative – Criteria details found here</p> <ul style="list-style-type: none"> Morphine Equivalent Daily Dose Concomitant use of Opioids and Benzodiazepines <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 18 years: codeine-containing products and tramadol-containing products <p>Quantity Limit (per 31 rolling days)</p> <ul style="list-style-type: none"> 62 tablets: butalbital/codeine combinations, codeine combinations, dihydrocodeine combinations, fentanyl, hydrocodone, hydromorphone, levorphanol, meperidine, morphine, oxycodone, oxymorphone, pentazocine, tapentadol, tramadol 186 tablets: butalbital/acetaminophen, butalbital/aspirin 5 mL: butorphanol nasal 180 mL: oxycodone liquid 280 mL: QDOL0 <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months |
| acetaminophen/codeine | aspirin/butalbital/caffeine/codeine | |
| codeine | butalbital/acetaminophen/caffeine/codeine | |
| ENDOCET (oxycodone/acetaminophen) | butorphanol | |
| hydrocodone/acetaminophen | DILAUDID (hydromorphone) | |
| hydromorphone | DSUVIA (sufentanil) | |
| morphine sulfate | fentanyl citrate | |
| oxycodone | FENTORA (fentanyl) | |
| oxycodone/acetaminophen (325 mg acetaminophen formulations) | FIORICET W/CODEINE (butalbital/acetaminophen/codeine) | |
| tramadol 50 mg tablet | hydrocodone/ibuprofen | |
| tramadol/acetaminophen | meperidine | |
| | NALOCET (oxycodone/acetaminophen) | |
| | levorphanol | |
| | oxymorphone | |
| | pentazocine/naloxone | |
| | PERCOCET (oxycodone/acetaminophen) | |
| | PROLATE (oxycodone/acetaminophen) | |
| | ROXICODONE (oxycodone) | |
| | ROXYBOND (oxycodone) | |
| | SEGLENTIS (tramadol/celecoxib) | |
| | tramadol 25 mg, 75 mg, 100 mg tablet | |
| | tramadol solution | |
| ANALGESICS, OPIOID-LONG ACTING DUR+ | | |
| BUTRANS (buprenorphine) | BELBUCA (buprenorphine) | <p>MS DOM Opioid Initiative – Criteria details found here</p> <ul style="list-style-type: none"> Morphine Equivalent Daily Dose Concomitant use of Opioids and Benzodiazepines <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 18 years: BUTRANS and tramadol-containing products <p>Quantity Limit (per 31 rolling days)</p> <ul style="list-style-type: none"> 31 tablets: AVINZA, hydromorphone ER, HYSINGLA ER, tramadol ER 62 tablets: methadone, morphine ER, OXYCONTIN, oxymorphone ER, ZOHYDRO ER 62 films: BELBUCA 10 patches: fentanyl 4 patches: BUTRANS <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months |
| fentanyl patch | buprenorphine patch | |
| morphine sulfate ER tablet | CONZIP (tramadol) | |
| | hydrocodone bitartrate ER | |
| | hydromorphone ER | |
| | HYSINGLA ER (hydrocodone) | |
| | methadone | |
| | methadone intensol | |
| | METHADOSE (methadone) | |
| | morphine sulfate ER capsule | |
| | MS CONTIN (morphine) | |
| | oxycodone ER | |
| | OXYCONTIN (oxycodone) | |
| | oxymorphone ER | |
| | tramadol ER | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| ANALGESICS/ANESTHETICS (TOPICAL) | | |
| diclofenac 1%, 3% gel | DERMACINRX LIDOCAN (lidocaine) | <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 1 bottle (112 mL): diclofenac 2% solution pump • 1 bottle (150 mL): diclofenac 1.5% solution <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 preferred agents in the past 6 months <p>Lidocaine 5% Patch</p> <ul style="list-style-type: none"> • Documented diagnosis of Herpetic Neuralgia OR • Documented diagnosis of Diabetic Neuropathy <p>ZTLIDO</p> <ul style="list-style-type: none"> • Documented diagnosis of Herpetic Neuralgia |
| lidocaine 4% cream, patch, solution | DERMACINRX LIDOGEL (lidocaine) | |
| lidocaine 5% cream, ointment, patch | DERMACINRX LIDOREX (lidocaine) | |
| lidocaine 40 mg/mL solution | diclofenac epolamine | |
| lidocaine/prilocaine cream | diclofenac sodium 2% solution pump | |
| TRIDACAINE (lidocaine) patch | DICLOGEN (diclofenac/menthol/camphor) kit | |
| TRIDACAINE XL (lidocaine) patch | DOLOGESIC PAIN RELIEF (lidocaine) | |
| ULTRA LIDO (lidocaine) cream, gel | LIDAFLEX (lidocaine) | |
| | lidocaine 3% cream | |
| | lidocaine 4% kit, liquid | |
| | lidocaine/hydrocortisone | |
| | lidocaine/prilocaine kit | |
| | LIDOCAN II, III, IV, V (lidocaine) | |
| | LIDOCORT (lidocaine/hydrocortisone) | |
| | LIDODERM (lidocaine) | |
| | LIDOTRAL (lidocaine) | |
| | LIXOFEN (diclofenac) | |
| | PENNSAID (diclofenac) | |
| | PLIAGLIS (lidocaine/tetracaine) | |
| | TRIDACAINE II, III (lidocaine) patch | |
| | ZTLIDO (lidocaine) | |
| ANDROGENIC AGENTS ^{DUR+} | | |
| testosterone | ANDROGEL (testosterone) | <p>All Agents</p> <ul style="list-style-type: none"> • Limited to male gender <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months <p>TLANDO</p> <ul style="list-style-type: none"> • Requires clinical review |
| | JATENZO (testosterone undecanoate) | |
| | NATESTO (testosterone) | |
| | TESTIM (testosterone) | |
| | TLANDO (testosterone) | |
| | VOGELXO (testosterone) | |
| ANGIOTENSIN MODULATORS ^{DUR+} | | |
| ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITORS | | See next page for PA Criteria/DUR+ Rules |
| benazepril | ACCUPRIL (quinapril) | |
| captopril | ALTACE (ramipril) | |
| enalapril | EPANED (enalapril) | |
| fosinopril | LOTENSIN (benazepril) | |
| lisinopril | moexipril | |
| quinapril | perindopril | |
| ramipril | QBRELIS (lisinopril) | |
| trandolapril | VASOTEC (enalapril) | |
| | ZESTRIL (lisinopril) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA | |
|---|--|---|--|
| ANGIOTENSIN MODULATORS ^{DUR+} (continued) | | | |
| ACE INHIBITOR (ACEI) COMBINATIONS | | | |
| benazepril/amlodipine | ACCURETIC (quinapril/hydrochlorothiazide) | <p>EPANED</p> <ul style="list-style-type: none"> Automatic approval issued for 0-6 years of age <p>ENTRESTO</p> <ul style="list-style-type: none"> Age ≥ 1 year and documented diagnosis of Heart Failure with Systemic Ventricular Systolic Dysfunction OR Age ≥ 18 years and documented diagnosis of Heart Failure <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> ACEIs: <ul style="list-style-type: none"> Have tried 2 different preferred single entity agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days ACEI/CCB Combinations: <ul style="list-style-type: none"> Have tried 2 different preferred ACEI/CCB agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days ACEI/Diuretic Combinations: <ul style="list-style-type: none"> Have tried 2 different preferred ACEI/Diuretic agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days ARBs: <ul style="list-style-type: none"> Have tried 2 different preferred single entity agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days ARB/CCB and ARB/CCB/Diuretic Combinations: <ul style="list-style-type: none"> Have tried 1 preferred ARB/CCB agent in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days ARB/Diuretic Combinations: <ul style="list-style-type: none"> Have tried 2 different preferred ARB/Diuretic agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days Direct Renin Inhibitors: <ul style="list-style-type: none"> Documented diagnosis of Hypertension AND Have tried 2 different preferred ACEI or ARB single-entity agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days Direct Renin Inhibitor Combinations: <ul style="list-style-type: none"> Documented diagnosis of Hypertension AND Have tried 2 different preferred ACEI or ARB diuretic agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days | |
| benazepril/hydrochlorothiazide | LOTENSIN HCT (benazepril/hydrochlorothiazide) | | |
| captopril/hydrochlorothiazide | LOTREL (benazepril/amlodipine) | | |
| enalapril/hydrochlorothiazide | VASERETIC (enalapril/hydrochlorothiazide) | | |
| fosinopril/hydrochlorothiazide | ZESTORETIC (lisinopril/hydrochlorothiazide) | | |
| lisinopril/hydrochlorothiazide | | | |
| quinapril/hydrochlorothiazide | | | |
| trandolapril/verapamil ER | | | |
| ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs) | | | |
| irbesartan | ATACAND (candesartan) | | |
| losartan | AVAPRO (irbesartan) | | |
| olmesartan | BENICAR (olmesartan) | | |
| telmisartan | candesartan | | |
| valsartan tablet | COZAAR (losartan) | | |
| | DIOVAN (valsartan) | | |
| | EDARBI (azilsartan) | | |
| | eprosartan | | |
| | MICARDIS (telmisartan) | | |
| | valsartan solution | | |
| ARB COMBINATIONS | | | |
| ENTRESTO (valsartan/sacubitril) tablet ^{DUR+} | ATACAND HCT (candesartan/hydrochlorothiazide) | | |
| irbesartan/hydrochlorothiazide | AVALIDE (irbesartan/hydrochlorothiazide) | | |
| losartan/hydrochlorothiazide | AZOR (olmesartan/hydrochlorothiazide) | | |
| olmesartan/amlodipine | BENICAR HCT (olmesartan/hydrochlorothiazide) | | |
| olmesartan/hydrochlorothiazide | candesartan/hydrochlorothiazide | | |
| telmisartan/hydrochlorothiazide | DIOVAN-HCT (valsartan/hydrochlorothiazide) | | |
| valsartan/amlodipine | EDARBYCLOR (azilsartan/chlorthalidone) | | |
| valsartan/amlodipine/hydrochlorothiazide | ENTRESTO (valsartan/sacubitril) sprinkle capsule | | |
| valsartan/hydrochlorothiazide | EXFORGE (valsartan/amlodipine) | | |
| | EXFORGE HCT (valsartan/amlodipine/hydrochlorothiazide) | | |
| | olmesartan/amlodipine/hydrochlorothiazide | | |
| | telmisartan/amlodipine | | |
| | TRIBENZOR (olmesartan/amlodipine/hydrochlorothiazide) | | |
| | valsartan/sacubitril | | |
| DIRECT RENIN INHIBITORS | | | |
| | aliskiren | | |
| | TEKTURNA (aliskiren) | | |
| DIRECT RENIN INHIBITOR COMBINATIONS | | | |
| | TEKTURNA HCT (aliskiren/hydrochlorothiazide) | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| ANTIBIOTICS (GI) & RELATED AGENTS | | |
| metronidazole tablet | AEMCOLO (rifamycin) | |
| neomycin | DIFICID (fidaxomicin) | |
| tinidazole | FIRVANQ (vancomycin) | |
| vancomycin oral solution | FLAGYL (metronidazole) | |
| | LIKMEZ (metronidazole) | |
| | metronidazole capsule | |
| | nitazoxanide | |
| | paromomycin | |
| | REBYOTA (fecal microbiota, live-jslm) | |
| | VANCOCIN (vancomycin) | |
| | vancomycin capsule | |
| | VOWST (fecal microbio spore, live-brpk) | |
| | XIFAXAN (rifaximin) | |
| ANTIBIOTICS (MISCELLANEOUS) | | |
| LINCOSAMIDE ANTIBIOTICS | | Quantity Limit <ul style="list-style-type: none"> • 6 tablets/month: SIVEXTRO SIVEXTRO – MANUAL PA ZYVOX – MANUAL PA |
| clindamycin | CLEOCIN (clindamycin) | |
| | CELOCIN PEDIATRIC (clindamycin) | |
| MACROLIDES | | |
| azithromycin | ERYPD (erythromycin ethylsuccinate) suspension | |
| clarithromycin | ERYTHROCIN (erythromycin stearate) | |
| clarithromycin ER | ZITHROMAX (azithromycin) | |
| E.E.S (erythromycin ethylsuccinate) suspension | | |
| ERY-TAB (erythromycin) | | |
| erythromycin | | |
| erythromycin ethylsuccinate | | |
| NITROFURANTOIN DERIVATIVES | | |
| nitrofurantoin capsule | FURADANTIN (nitrofurantoin) suspension | |
| nitrofurantoin monohydrate macrocrystals | MACROBID (nitrofurantoin monohydrate macrocrystals) | |
| | nitrofurantoin suspension | |
| OXAZOLIDINONES | | |
| | Linezolid | |
| | SIVEXTRO (tedizolid) | |
| | ZYVOX (linezolid) | |
| ANTIBIOTICS (TOPICAL) | | |
| bacitracin ^{OTC} | CENTANY (mupirocin) | |
| bacitracin/polymyxin ^{OTC} | CENTANY AT (mupirocin) | |
| gentamicin sulfate | mupirocin cream | |
| mupirocin ointment | XEPI (ozenoxacin) | |
| neomycin/bacitracin/polymyxin ^{OTC} | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| ANTIBIOTICS (VAGINAL) | | |
| CLEOCIN (clindamycin) | clindamycin phosphate | |
| NUVESSA (metronidazole) | CLINDESSE (clindamycin) | |
| | SOLOSEC (secnidazole) | |
| | XACIATO (clindamycin) | |
| ANTICOAGULANTS | | |
| LOW MOLECULAR WEIGHT HEPARIN (LMWH) | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • LMWH: <ul style="list-style-type: none"> ○ Have tried 1 preferred agent in the past 6 months OR ○ 90 days of therapy with the requested agent in the past 105 days • Oral: <ul style="list-style-type: none"> ○ Have tried 2 different preferred oral agents in the past 6 months OR ○ 90 days of therapy with the requested agent in the past 105 days |
| enoxaparin | ARIXTRA (fondaparinux) | |
| | fondaparinux | |
| | FRAGMIN (dalteparin) | |
| | LOVENOX (enoxaparin) | |
| ORAL | | |
| ELIQUIS (apixaban) | dabigatran | |
| JANTOVEN (warfarin) | PRADAXA (dabigatran) pellet pack | |
| PRADAXA (dabigatran) capsule | SAVAYSA (edoxaban) | |
| warfarin | | |
| XARELTO (rivaroxaban) | | |
| ANTICONVULSANTS ^{DUR+} | | |
| ADJUVANTS | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 6 months: DIACOMIT • 1 year: BANZEL, EPIDIOLEX • 2 years: ONFI, SYMPAZAN • 6 years: VALTOCO • 12 years: NAYZILAM <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 2 years: VIGAFYDE <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 2 twin packs: DIASTAT • 2 packages: NAYZILAM • 2 cartons: VALTOCO <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months OR • Documented diagnosis of Seizure AND • 90 days of therapy with the requested agent in the past 105 days <p style="background-color: yellow;">See next page for additional PA Criteria/DUR+ Rules</p> |
| carbamazepine | APTOM (eslicarbazepine acetate) | |
| carbamazepine ER 12-hour capsule | BANZEL (rufinamide) | |
| DEPAKOTE ER (divalproex) | BRIVIACT (brivaracetam) | |
| DEPAKOTE SPRINKLE (divalproex) | carbamazepine ER 12-hour tablet | |
| divalproex | CARBATROL (carbamazepine) | |
| divalproex ER | DEPAKOTE (divalproex) | |
| divalproex sprinkle | DIACOMIT (stiripentol) | |
| EPIDIOLEX (cannabidiol) | ELEPSIA XR (levetiracetam) | |
| lacosamide | EPRONTIA (topiramate) | |
| lamotrigine | EQUETRO (carbamazepine) | |
| lamotrigine blue, green, orange dose pack | felbamate | |
| levetiracetam | FELBATOL (felbamate) | |
| levetiracetam ER | FINTEPLA (fenfluramine) | |
| oxcarbazepine tablet | FYCOMPA (perampanel) | |
| tiagabine | KEPPRA (levetiracetam) | |
| topiramate | KEPPRA XR (levetiracetam) | |
| topiramate sprinkle | LAMICTAL (lamotrigine) | |
| TRILEPTAL (oxcarbazepine) suspension | LAMICTAL XR (lamotrigine) | |
| valproic acid | lamotrigine ER | |
| zonisamide | lamotrigine ODT | |
| | lamotrigine ODT blue, green, orange dose pack | |
| | MOTPOLY XR (lacosamide) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| ANTICONVULSANTS^{DUR+} (continued) | | |
| ADJUVANTS (continued) | | <p style="background-color: yellow;">See previous page for additional PA Criteria/DUR+ Rules</p> <p>Banzel, Onfi, and Sympazan</p> <ul style="list-style-type: none"> Documented diagnosis of Lennox-Gastaut Syndrome and have tried 1 preferred agent for Lennox-Gastaut Syndrome in the past 6 months OR Documented diagnosis of Seizure and 90 days of therapy with the requested agent in the past 105 days <p>DIACOMIT</p> <ul style="list-style-type: none"> Documented diagnosis of Dravet Syndrome AND 1 claim for clobazam in the past 30 days <p>EPIDIOLEX</p> <ul style="list-style-type: none"> Documented diagnosis of Dravet Syndrome, Lennox-Gastaut Syndrome, or Seizures associated with Tuberous Sclerosis Complex OR 1 claim for EPIDIOLEX in the past 30 days <p>FINTEPLA</p> <ul style="list-style-type: none"> Requires clinical review <p>SABRIL Powder for Oral Solution</p> <ul style="list-style-type: none"> Documented diagnosis of Infantile Spasms OR Have tried 2 different preferred agents in the past 6 months OR Documented diagnosis of Seizure AND 90 days of therapy with the requested agent in the past 105 days <p>Topiramate ER</p> <ul style="list-style-type: none"> Documented diagnosis of Seizure AND 90 days of therapy with the requested agent in the past 105 days OR 30 days of therapy with topiramate IR in the past 6 months <p>VIGAFYDE</p> <ul style="list-style-type: none"> Age ≤ 2 years AND Documented diagnosis of infantile spasms |
| | oxcarbazepine suspension | |
| | oxcarbazepine ER | |
| | OXTELLAR XR (oxcarbazepine) | |
| | QUDEXY XR (topiramate) | |
| | ROWEEPRA (levetiracetam) | |
| | rufinamide | |
| | SABRIL (vigabatrin) | |
| | SPRITAM (levetiracetam) | |
| | SUBVENITE (lamotrigine) | |
| | SUBVENITE (lamotrigine) blue, green, orange dose pack | |
| | TEGRETOL (carbamazepine) | |
| | TEGRETOL XR (carbamazepine) | |
| | TOPAMAX (topiramate) | |
| | topiramate ER | |
| | TRILEPTAL (oxcarbazepine) tablet | |
| | TROKENDI XR (topiramate) | |
| | vigabatrin | |
| | VIGADRONE (vigabatrin) | |
| | VIGAFYDE (vigabatrin) | |
| | VIGPODER (vigabatrin) | |
| | VIMPAT (lacosamide) | |
| | XCOPRI (cenobamate) | |
| | ZONISADE (zonisamide) suspension | |
| | ZTALMY (ganaxolone) | |
| HYDANTOINS | | |
| | DILANTIN (phenytoin) | |
| | DILANTIN-125 (phenytoin) | |
| | PHENYTEK (phenytoin) | |
| | phenytoin | |
| | phenytoin ER | |
| SELECTED BENZODIAZEPINES | | |
| | clobazam | |
| | DIASTAT (diazepam) rectal gel | |
| | diazepam rectal gel | |
| | LIBERVANT (diazepam) | |
| | NAYZILAM (midazolam) | |
| | ONFI (clobazam) | |
| | VALTOCO (diazepam) | |
| | SYMPAZAN (clobazam) | |
| SUCCINIMIDES | | |
| | ethosuximide | |
| | CELONTIN (methsuximide) | |
| | methsuximide | |
| | ZARONTIN (ethosuximide) | |



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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|------------------------------------|---------------------------------------|--|
| ANTIDEPRESSANTS, OTHER DUR+ | | |
| bupropion | APLENZIN (bupropion) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 18 years: all agents <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months OR • Have tried 1 preferred agent and 1 SSRI in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>AUVELITY</p> <ul style="list-style-type: none"> • Requires clinical review <p>DRIZALMA Sprinkles and duloxetine</p> <ul style="list-style-type: none"> • Automatic approval issued with a diagnosis of Generalized Anxiety Disorder for 7-11 years of age <p>ZURZUVAE – MANUAL PA</p> |
| bupropion SR | AUVELITY (bupropion/dextromethorphan) | |
| bupropion XL | desvenlafaxine ER | |
| mirtazapine | EFFEXOR XR (venlafaxine) | |
| trazodone | EMSAM (selegiline) | |
| TRINTELLIX (vortioxetine) | FETZIMA (levomilnacipran) | |
| venlafaxine | FORFIVO XL (bupropion) | |
| venlafaxine ER capsule | MARPLAN (isocarboxazid) | |
| vilazodone | NARDIL (phenelzine) | |
| | nefazodone | |
| | phenelzine | |
| | PRISTIQ (desvenlafaxine) | |
| | REMERON (mirtazapine) | |
| | tranylcypromine | |
| | venlafaxine ER tablet | |
| | VIIIBRYD (vilazodone) | |
| | WELLBUTRIN SR (bupropion) | |
| | WELLBUTRIN XL (bupropion) | |
| | ZURZUVAE (zuranolone) | |
| ANTIDEPRESSANTS, SSRIs DUR+ | | |
| citalopram solution, tablet | CELEXA (citalopram) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 6 years: ZOLOFT • 7 years: LEXAPRO, PROZAC • 8 years: LUVOX • 18 years: CELEXA, LUVOX CR, PAXIL, PEVEVA, PROZAC 90 mg <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 60 years: CELEXA <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days |
| escitalopram | citalopram capsule | |
| fluoxetine capsule | fluoxetine solution, tablet | |
| fluvoxamine | fluoxetine DR capsule | |
| paroxetine tablet | fluvoxamine ER capsule | |
| paroxetine CR | LEXAPRO (escitalopram) | |
| paroxetine ER | paroxetine suspension, capsule | |
| sertraline tablet, solution | PAXIL (paroxetine) | |
| | PAXIL CR (paroxetine) | |
| | PROZAC (fluoxetine) | |
| | sertraline capsule | |
| | ZOLOFT (sertraline) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA | |
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| ANTIEMETICS DUR+ | | | |
| 5HT3 RECEPTOR BLOCKERS | | | |
| ondansetron solution, tablet | ANZIMET (dolasetron) | <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 6 tablets: AKYNZEO • 100 mL: ZOFRAN solution <p>Non-Preferred Agents</p> <ul style="list-style-type: none"> • Have tried 1 preferred agent in the past 6 months <p>AKYNZEO – MANUAL PA</p> <p>Note: Injectables in this class are closed to point of sale. PA required if not administered in clinic/hospital.</p> | |
| ondansetron ODT 4 mg, 8 mg | granisetron | | |
| | ondansetron ODT 16 mg tablet | | |
| | SANCUSO (granisetron) | | |
| ANTIEMETIC COMBINATIONS | | | |
| DICLEGIS (doxylamine/pyridoxine) | AKYNZEO (netupitant/palonosetron) | | |
| | BONJESTA (doxylamine/pyridoxine) | | |
| | doxylamine/pyridoxine | | |
| CANNABINOIDS | | | |
| | dronabinol | | |
| | MARINOL (dronabinol) | | |
| NMDA RECEPTOR ANTAGONISTS | | | |
| aprepitant | EMEND (aprepitant) | | |
| ANTIFUNGALS (ORAL) DUR+ | | | |
| clotrimazole | ANCOBON (flucytosine) | <p>Griseofulvin suspension</p> <ul style="list-style-type: none"> • Automatic approval issued for 0-11 years of age <p>Griseofulvin tablets</p> <ul style="list-style-type: none"> • Automatic approval issued for 12-17 years of age <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 18 years: CRESEMBA <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months <p>HIV Opportunistic Infection</p> <ul style="list-style-type: none"> • Non-Preferred agent indicated for treatment (^) AND • Documented diagnosis of HIV <p>CRESEMBA – MANUAL PA</p> <p>SPORANOX</p> <ul style="list-style-type: none"> • Requires clinical review | |
| fluconazole | BREXAFEMME (ibrexafungerp) | | |
| nystatin | CRESEMBA (isavuconazonium sulfate) | | |
| terbinafine | DIFLUCAN (fluconazole) | | |
| | flucytosine | | |
| | griseofulvin | | |
| | griseofulvin ultramicrosize | | |
| | itraconazole | | |
| | ketoconazole | | |
| | NOXAFIL (posaconazole) | | |
| | ORAVIG (miconazole) | | |
| | Posaconazole | | |
| | SPORANOX (itraconazole) | | |
| | TOLSURA (itraconazole) | | |
| | VFEND (voriconazole) | | |
| | VIVJOA (oteseconazole) | | |
| | voriconazole | | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| ANTIFUNGALS (TOPICAL) DUR+ | | |
| ANTIFUNGALS | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months <p>MICOTRIN AC, MYCOZYL, and clotrimazole 30 mL solution</p> <ul style="list-style-type: none"> Require clinical review |
| ciclopirox cream, gel, solution, suspension | BENSAL HP (salicylic acid) | |
| clotrimazole cream, solution ^{Rx & OTC} | CILODAN (ciclopirox) | |
| econazole | ciclopirox shampoo | |
| ketoconazole cream, shampoo | clotrimazole solution (NDC 50228-0502-61) | |
| LUZU (luliconazole) | ERTACZO (sertaconazole) | |
| miconazole cream, powder, solution ^{OTC} | EXTINA (ketoconazole) | |
| miconazole/zinc oxide/petrolatum ointment | JUBLIA (efinaconazole) | |
| nystatin cream, ointment, powder | ketoconazole foam | |
| terbinafine ^{OTC} | KETODAN (ketoconazole) | |
| tolnaftate cream, solution ^{OTC} | LOPROX (ciclopirox) | |
| | luliconazole | |
| | MICOTRIN AC (clotrimazole) | |
| | MYCOZYL AC (clotrimazole) | |
| | MYCOZYL AP (miconazole) | |
| | naftifine | |
| | NAFTIN (naftifine) | |
| | oxiconazole | |
| | OXISTAT (oxiconazole) | |
| | tavaborole | |
| | VOTRIZA-AL (clotrimazole) | |
| | VUSION (miconazole/zinc oxide/petrolatum) | |
| ANTIFUNGAL/STEROID COMBINATIONS | | |
| clotrimazole/betamethasone cream | clotrimazole/betamethasone lotion | |
| nystatin/triamcinolone | | |
| ANTIFUNGALS (VAGINAL) | | |
| clotrimazole cream ^{OTC} | 3-DAY VAGINAL CREAM (clotrimazole) | |
| clotrimazole-3 cream | GYNAZOLE 1 (butoconazole) | |
| miconazole kit ^{OTC} | terconazole suppository | |
| terconazole cream | | |
| ANTIHISTAMINES, MINIMALLY SEDATING AND COMBINATIONS DUR+ | | |
| MINIMALLY SEDATING ANTIHISTAMINES | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of Allergy or Urticaria AND Have tried 2 different preferred agents in the past 12 months |
| cetirizine capsule, solution, tablet ^{OTC} | cetirizine chewable tablet ^{OTC} | |
| loratadine chewable tablet, ODT, solution, tablet ^{OTC} | CLARINEX (desloratadine) | |
| | desloratadine | |
| | levocetirizine | |
| MINIMALLY SEDATING ANTIHISTAMINE/DECONGESTANT COMBINATIONS | | |
| cetirizine/pseudoephedrine | CLARINEX-D 12 HOUR (desloratadine/pseudoephedrine) | |
| loratadine/pseudoephedrine | fexofenadine/pseudoephedrine | |



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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---------------------------------|---|
| ANTIMIGRAINE AGENTS, ACUTE TREATMENT | | |
| CGRP ORAL AND NASAL | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 6 years: MAXALT 12 years: AXERT, TREXIMET, ZOMIG nasal spray 18 years: AMERGE, FROVA, IMITREX, NURTEC ODT, ONZETRA XSAIL, RELPAX, REYVOW, TOSYMRA, UBRELVY, ZEMBRACE, ZOMIG tablets <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> ORAL <ul style="list-style-type: none"> 4 tablets: REYVOW 50 mg 6 tablets: AXERT, RELPAX, ZOMIG 8 tablets: NURTEC ODT, REYVOW 100 mg 9 tablets: AMERGE, FROVA, IMITREX, TREXIMET 12 tablets: MAXALT 16 tablets: UBRELVY NASAL <ul style="list-style-type: none"> 1 box: all agents <p>CUMULATIVE Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> INJECTABLES <ul style="list-style-type: none"> 4 injections: all agents <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> ORAL <ul style="list-style-type: none"> Have tried 2 preferred oral agents in the past 90 days NASAL <ul style="list-style-type: none"> Have tried 2 preferred oral agents in the past 90 days AND Have tried a preferred nasal agent in the past 90 days <p>AXERT, TREXIMET, and ZOMIG nasal</p> <ul style="list-style-type: none"> Automatic approval for 12-17 years of age <p>NURTEC ODT and UBRELVY</p> <ul style="list-style-type: none"> Documented diagnosis of Migraine AND Have tried 2 different triptans in the past 6 months AND No concurrent therapy with another CGRP agent or strong CYP3A4 inhibitor <p>REYVOW</p> <ul style="list-style-type: none"> Documented diagnosis of Migraine AND Have tried 2 different triptans in the past 90 days AND Have tried preferred NURTEC ODT in the past 90 days <p>ZAVZPRET</p> <ul style="list-style-type: none"> Documented diagnosis of Migraine AND Have tried 2 different triptans in the past 6 months AND Have tried both NURTEC ODT and UBRELVY in the past 6 months AND No concurrent therapy with another CGRP AGENT |
| | | |
| INJECTABLES | | |
| sumatriptan | IMITREX (sumatriptan) | |
| | ZEMBRACE SYMTOUCH (sumatriptan) | |
| NASAL | | |
| sumatriptan | IMITREX (sumatriptan) | |
| | TOSYMRA (sumatriptan) | |
| | zolmitriptan | |
| | ZOMIG (zolmitriptan) | |
| TRIPTANS AND RELATED AGENTS (ORAL) ^{DUR+} | | |
| naratriptan | almotriptan | |
| rizatriptan | eletriptan | |
| sumatriptan | FROVA (frovatriptan) | |
| zolmitriptan | frovatriptan | |
| zolmitriptan ODT | IMITREX (sumatriptan) | |
| | MAXALT (rizatriptan) | |
| | MAXALT MLT (rizatriptan) | |
| | RELPAX (eletriptan) | |
| | REYVOW (lasmiditan) | |
| | sumatriptan/naproxen | |
| | ZOMIG (zolmitriptan) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| ANTIMIGRAINE AGENTS, PROPHYLAXIS | | |
| INJECTABLES | | |
| AIMOVIG Autoinjector (erenumab-aooe) ^{DUR+} | EMGALITY Syringe (galcanezumab-gnlm) 300 mg/mL | <p>Preferred Injectables</p> <ul style="list-style-type: none"> History of 3 claims with the requested agent in the past 105 days OR New starts require clinical review <p>Non-preferred Injectables</p> <ul style="list-style-type: none"> Require clinical review <p>AIMOVIG, AJOVY, and EMGALITY – MANUAL PA</p> <p>VYEPTI – MANUAL PA</p> |
| AJOVY Autoinjector (fremanezumab-vfrm) ^{DUR+} | VYEPTI (eptinezumab-jjmr) | |
| AJOVY Syringe (fremanezumab-vfrm) ^{DUR+} | | |
| EMGALITY Pen (galcanezumab-gnlm) ^{DUR+} | | |
| EMGALITY Syringe (galcanezumab-gnlm) 120 mg/mL ^{DUR+} | | |
| ORAL | | |
| | QULIPTA (atogepant) | |
| * ANTINEOPLASTICS – SELECTED SYSTEMIC ENZYME INHIBITORS | | |
| BOSULIF (bosutinib) tablet | AFINITOR (everolimus) | FARYDAK – MANUAL PA |
| CAPRESLA (vandetanib) | AFINITOR DISPERZ (everolimus) | IBRANCE |
| COMETRIQ (cabozantinib) | AKEEGA (niraparib/abiraterone) | |
| COTELLIC (cobimetinib) | ALECENSA (lectinib) | <ul style="list-style-type: none"> Documented diagnosis of WD-DDLS for retroperitoneal sarcoma OR All other indications require clinical review |
| everolimus | ALUNBRIG (brigatinib) | LENVIMA |
| GILOTRIF (afatinib) | AUGTYRO (repotrectinib) | |
| ICLUSIG (ponatinib) | AYVAKIT (avapritinib) | <ul style="list-style-type: none"> Documented diagnosis of thyroid cancer, hepatocellular carcinoma, or renal cell carcinoma AND History of 1 claim for everolimus in the past 30 days AND History of 1 anti-angiogenic agent in the past 2 years OR All other indications require clinical review |
| imatinib | BALVERSA (erdafitinib) | LYNPARZA Tablets |
| IMBRUVICA (ibrutinib) | BOSULIF (bosutinib) capsule | |
| INLYTA (axitinib) | BRAFTOVI (encorafenib) | |
| IRESSA (gefitinib) | BRUKINSA (zanubrutinib) | |
| JAKAFI (ruxolitinib) | CABOMETYX (cabozantinib) | |
| MEKINIST (trametinib) | CALQUENCE (acalabrutinib) | |
| NEXAVAR (sorafenib) | COPIKTRA (duvelisib) | |
| ROZLYTREK (entrectinib) | DANZITEN (nilotinib) | |
| SPRYCEL (dasatinib) | dasatinib | |
| STIVARGA (regorafenib) | DATROWAY (datopotomab deruxtecan-dlnk) ^{NR} | |
| SUTENT (sunitinib) | DAURISMO (glasdegib) | |
| TAFINLAR (dabrafenib) | ERIVEDGE (vismodegib) | |
| TARCEVA (erlotinib) | ERLEADA (apalutamide) | |
| TASIGNA (nilotinib) | erlotinib | |
| TURALIO (pexidartinib) | FOTIVDA (tivozanib) | |
| TYKERB (lapatinib) | FURZAQLA (fruquintinib) | |
| VOTRIENT (pazopanib) | GAVRETO (pralsetinib) | |
| XALKORI (crizotinib) | gefitinib | |
| XTANDI (enzalutamide) | GLEEVEC (imatinib) | |
| ZELBORAF (vemurafenib) | IBRANCE (palbociclib) | |
| ZYDELIG (idelalisib) | IDHIFA (enasidenib) | |
| ZYKADIA (ceritinib) | IMKELDI (imatinib) | |
| | INQOVI (decitabine/cedazuridine) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| *ANTINEOPLASTICS – SELECTED SYSTEMIC ENZYME INHIBITORS <i>(continued)</i> | | |
| | INREBIC (fedratinib) | See previous page for PA Criteria/DUR+ Rules |
| | ITOVEBI (inavolisib) | |
| | IWILFIN (eflornithine) | |
| | JAYPIRCA (pirtobrutinib) | |
| | KISQALI (ribociclib) | |
| | KISQALI-FEMARA CO-PACK (ribociclib/letrozole) | |
| | KOSELUGO (selumetinib/vitamin E) | |
| | KRAZATI (adagrasib) | |
| | lapatinib | |
| | LAZCLUZE (lazertinib) | |
| | LENVIMA (lenvatinib) | |
| | LOBRENA (lorlatinib) | |
| | LUMAKRAS (sotorasib) | |
| | LYNPARZA (olaparib) | |
| | LYTGOBI (futibatinib) | |
| | MEKTOVI (binimetinib) | |
| | NERLYNX (neratinib) | |
| | NUBEQA (darolutamide) | |
| | ODOMZO (sonidegib) | |
| | OGSIVEO (nirogacestat) | |
| | OJEMDA (tovorafenib) | |
| | OJJAARA (momelotinib) | |
| | ONUREG (azacitidine) | |
| | ORGOVYX (relugolix) | |
| | pazopanib | |
| | PEMAZYRE (pemigatinib) | |
| | PIQRAY (alpelisib) | |
| | QINLOCK (ripretinib) | |
| | RETEVMO (selpercatinib) | |
| | REVUFORJ (revumenib) | |
| | REZLIDHIA (olutasidenib) | |
| | RUBRACA (rucaparib) | |
| | RYDAPT (midostaurin) | |
| | SCEMBLIX (asciminib) | |
| | sorafenib | |
| | sunitinib | |
| | TABRECTA (capmatinib) | |
| | TAGRISSO (osimertinib) | |
| | TALZENNA (talazoparib) | |
| | TAZVERIK (tazemetostat) | |
| | TECENTRIZ HYBREZA (atezolizumab-hyaluronidase) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| *ANTINEOPLASTICS – SELECTED SYSTEMIC ENZYME INHIBITORS <i>(continued)</i> | | |
| | TEPMETKO (tepotinib) | See previous page for PA Criteria/DUR+ Rules |
| | TIBSOVO (ivosidenib) | |
| | TORPENZ (everolimus) | |
| | TRUQAP (capivasertib) | |
| | TUKYSA (tucatinib) | |
| | VANFLYTA (quizartinib) | |
| | VERZENIO (abemaciclib) | |
| | VITRAKVI (larotrectinib) | |
| | VIZIMPRO (dacomitinib) | |
| | VONJO (pacritinib) | |
| | VORANIGO (vorasidenib) | |
| | WELIREG (belzutifan) | |
| | XOSPATA (gilteritinib) | |
| | XPOVIO (selinexor) | |
| | ZEJULA (niraparib) | |
| ANTIOBESITY SELECT AGENTS | | |
| SAXENDA (liraglutide) | orlistat | All agents – MANUAL PA required |
| WEGOVY (semaglutide) | XENICAL (orlistat) | |
| ANTIPARASITICS (TOPICAL) DUR+ | | |
| PEDICULICIDES | | Minimum Age Limit <ul style="list-style-type: none"> • 2 months: permethrin 1% (OTC), permethrin 5% • 6 months: NATROBA, SKLICE • 2 years: piperonyl/pyrethrins (OTC) • 4 years: NATROBA • 6 years: OVIDE • 18 years: EURAX Non-Preferred Criteria <ul style="list-style-type: none"> • Pediculicides <ul style="list-style-type: none"> ○ Have tried 2 preferred topical lice agents in the past 90 days • Scabicides <ul style="list-style-type: none"> ○ Have tried permethrin 5% in the past 90 days |
| NATROBA (spinosad) | lindane | |
| permethrin 1% cream ^{OTC} | malathion | |
| VANALICE (piperonyl butoxide/pyrethrins) | OVIDE (malathion) | |
| | SKLICE (ivermectin) | |
| | spinosad | |
| SCABICIDES | | |
| ivermectin | CROTAN (crotamiton) | |
| permethrin 5% cream | ELIMITE (permethrin) | |
| | EURAX (crotamiton) | |
| | STROMECTOL (ivermectin) | |
| ANTIPARKINSON'S AGENTS (INJECTABLE) | | |
| | VYALEV (foscarnidopa/foslevodopa) ^{NR} | VYALEV <ul style="list-style-type: none"> • Requires clinical review |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| ANTIPARKINSON'S AGENTS (ORAL) | | |
| ANTICHOLINERGICS | | <p>DUR+</p> <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented diagnosis of Parkinson's disease AND • Have tried 2 different preferred agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>XADAGO</p> <ul style="list-style-type: none"> • Documented diagnosis of Parkinson's disease AND • 30 days of therapy with a carbidopa/levodopa combination agent in the past 45 days AND • 30 days of therapy with a selegiline agent in the past 45 days <p>GOCOVRI</p> <ul style="list-style-type: none"> • Documented diagnosis of Parkinson's disease AND • 30 days of therapy with amantadine IR in the past 105 days AND • 30 days of therapy with a carbidopa/levodopa combination agent in the past 45 days <p>LODOSYN and INBRIJA</p> <ul style="list-style-type: none"> • Documented diagnosis of Parkinson's disease AND • 30 days of therapy with a carbidopa/levodopa combination agent in the past 45 days <p>NOURIANZ</p> <ul style="list-style-type: none"> • Documented diagnosis of Parkinson's Disease AND • Have tried 1 preferred carbidopa/levodopa combination agent in the past 30 days AND • 30 days of therapy with a preferred adjunctive therapy in the past 45 days |
| benztropine | | |
| trihexyphenidyl | | |
| COMT INHIBITORS | | |
| entacapone | OGENTYS (opicapone) | |
| | TASMAR (tocapone) | |
| | tolcapone | |
| DOPAMINE AGONISTS | | |
| pramipexole | NEUPRO (rotigotine) | |
| ropinirole | pramipexole ER | |
| | ropinirole ER | |
| MAO-B INHIBITORS | | |
| selegiline | AZILECT (rasagiline) | |
| | rasagiline | |
| | XADAGO (safinamide) | |
| | ZELAPAR (selegiline) | |
| OTHERS | | |
| amantadine | carbidopa/levodopa ODT | |
| bromocriptine | carbidopa/levodopa/entacapone | |
| carbidopa | CREXONT (carbidopa/levodopa) | |
| carbidopa/levodopa tablet | DHIVY (carbidopa/levodopa) | |
| carbidopa/levodopa ER | DUOPA (carbidopa/levodopa) | |
| | GOCOVRI (amantadine) | |
| | INBRIJA (levodopa) | |
| | LODOSYN (carbidopa) | |
| | NOURIANZ (istradefylline) | |
| | OSMOLEX ER (amantadine) | |
| | RYTARY (carbidopa/levodopa) | |
| | SINEMET (carbidopa/levodopa) | |
| | STALEVO (carbidopa/levodopa/entacapone) | |
| ANTIPSORIATICS (TOPICAL) | | |
| calcipotriene cream | calcipotriene foam, ointment, solution | |
| ENSTILAR (calcipotriene/betamethasone) | calcipotriene/betamethasone | |
| TACLONEX (calcipotriene/betamethasone) | calcitriol ointment | |
| | DUOBRII (halobetasol/tazarotene) | |
| | SORILUX (calcipotriene) | |
| | tazarotene | |
| | VECTICAL (calcitriol) | |
| | VTAMA (tapinarof) | |
| | ZORYVE (roflumilast) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA | |
|---|--|---|--|
| ANTIPSYCHOTICS^{DUR+} | | | |
| INJECTABLE, ATYPICALS^{DUR+} | | | |
| ABILIFY ASIMTUFII (aripiprazole) | ERZOFRI (paliperidone palmitate) ^{NR} | <p>Concurrent Therapy Limit for Age < 18 years</p> <ul style="list-style-type: none"> 90 days with ≥ 2 agents in the last 120 days will require a MANUAL PA <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 3 years: HALDOL 5 years: RISPERDAL, thioridazine 6 years: ABILIFY, trifluoperazine 10 years: LATUDA, SAPHRIS, SEROQUEL, SYMBYAX 12 years: INVEGA, molindone, perphenazine, pimozide, thiothixene 13 years: REXULTI, ZYPREXA 18 years: ABILIFY MYCITE, CAPLYTA, CLOZARIL, COBENFY, FANAPT, fluphenazine, GEODON, loxapine, LYBALVI, NUPLAZID, perphenazine/amitriptyline, SECUADO, VRAYLAR, and all injectable agents <p>Quantity Limit</p> <ul style="list-style-type: none"> 3 syringes/year: ARISTADA INITIO <p>Non-Preferred Criteria – Atypical Agents</p> <ul style="list-style-type: none"> Have tried 2 preferred agents in the past 12 months OR 30 days of therapy with the requested agent in the past 180 days <p>All Long-Acting Injectable Agents</p> <ul style="list-style-type: none"> Documented diagnosis of schizophrenia or schizoaffective disorder <p>ABILIFY MAINTENA, ABILIFY ASIMTUFII, RISPERDAL CONSTA, and RYKINDO ER</p> <ul style="list-style-type: none"> Documented diagnosis of schizophrenia, schizoaffective disorder, or bipolar disorder <p>INVEGA HAFYERA</p> <ul style="list-style-type: none"> Documented diagnosis of schizophrenia or schizoaffective disorder AND 4 claims for INVEGA SUSTENNA or ERZOFRI in the past year OR 1 claim for INVEGA TRINZA in the past year OR 1 claim for INVEGA HAFYERA in the past year <p>COBENFY and OPIPZA</p> <ul style="list-style-type: none"> Require clinical review <p>NUPLAZID</p> <ul style="list-style-type: none"> Documented diagnosis of Parkinson's disease <p>VRAYLAR</p> <ul style="list-style-type: none"> Documented diagnosis of schizophrenia, schizoaffective disorder, bipolar disorder, or major depressive disorder AND 30 days of therapy with an antidepressant in the past 45 days OR 1 claim for a 90-day supply of an antidepressant in the past 105 days | |
| ABILIFY MAINTENA (aripiprazole) | GEODON (ziprasidone) | | |
| ARISTADA, ARISTADA INITIO (aripiprazole lauroxil) | olanzapine | | |
| INVEGA HAFYERA (paliperidone) | risperidone ER | | |
| INVEGA SUSTENNA (paliperidone palmitate) | RYKINDO (risperidone) | | |
| INVEGA TRINZA (paliperidone) | ziprasidone | | |
| PERSERIS (risperidone) | ZYPREXA (olanzapine) | | |
| RISPERDAL CONSTA (risperidone) | ZYPREXA RELPREVV (olanzapine) | | |
| UZEDY (risperidone) | | | |
| ORAL | | | |
| aripiprazole tablet | ABILIFY (aripiprazole) | | |
| asenapine | ABILIFY MYCITE (aripiprazole) | | |
| clozapine tablet | ADASUVE (loxapine) | | |
| fluphenazine | aripiprazole ODT, solution | | |
| haloperidol | CAPLYTA (lumateperone) | | |
| haloperidol lactate | chlorpromazine | | |
| olanzapine | clozapine ODT | | |
| perphenazine | CLOZARIL (clozapine) | | |
| perphenazine/amitriptyline | COBENFY (xanomeline/trospium) ^{NR} | | |
| quetiapine | FANAPT (iloperidone) | | |
| quetiapine ER | GEODON (ziprasidone) | | |
| risperidone | IGALMI (dexmedetomidine) | | |
| thioridazine | INVEGA (paliperidone) | | |
| trifluoperazine | LATUDA (lurasidone) | | |
| VRAYLAR (cariprazine) | lurasidone | | |
| ziprasidone | LYBALVI (olanzapine/samidorphan) | | |
| | NUPLAZID (pimavanserin) | | |
| | olanzapine/fluoxetine | | |
| | OPIPZA (aripiprazole) | | |
| | paliperidone ER | | |
| | REXULTI (brexpiprazole) | | |
| | RISPERDAL (risperidone) | | |
| | SAPHRIS (asenapine) | | |
| | SEROQUEL (quetiapine) | | |
| | SEROQUEL XR (quetiapine ER) | | |
| | SYMBYAX (olanzapine/fluoxetine) | | |
| | VERSACLOZ (clozapine) | | |
| | ZYPREXA, ZYPREXA ZYDIS (olanzapine) | | |
| TRANSDERMAL, ATYPICALS | | | |
| | SECUADO (asenapine) | | |



MISSISSIPPI DIVISION OF MEDICAID UNIVERSAL PREFERRED DRUG LIST

EFFECTIVE 01/01/2025
Version 2025_2
Updated 02/01/2025

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|--|--|
| ANTIRETROVIRALS ^{DUR+} | | |
| CAPSID INHIBITORS | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> 1 claim with the requested agent in the past 105 days <p>STRIBILD – MANUAL PA</p> <p>SUNLENCA</p> <ul style="list-style-type: none"> Requires clinical review <p>TYBOST – MANUAL PA</p> |
| | SUNLENCA (lenacapavir) | |
| CD4 DIRECTED ATTACHMENT INHIBITORS | | |
| | RUKOBIA (fostemsavir) | |
| CD4 DIRECTED HIV-1 INHIBITORS | | |
| | TROGARZO (ibalizumab-uiyk) | |
| COMBINATION PRODUCTS – NRTIs | | |
| abacavir/lamivudine | COMBIVIR (lamivudine/zidovudine) | |
| CABENUVA (cabotegravir/rilpivirine) | EPZICOM (abacavir/lamivudine) | |
| DOVATO (dolutegravir/lamivudine) | | |
| lamivudine/zidovudine | | |
| COMBINATION PRODUCTS – NUCLEOSIDE AND NUCLEOTIDE ANALOG RTIs | | |
| DESCOVY (emtricitabine/tenofovir alafenamide) | TRUVADA (emtricitabine/tenofovir) | |
| emtricitabine/tenofovir | | |
| COMBINATION PRODUCTS – NUCLEOSIDE AND NUCLEOTIDE ANALOG AND NON-NUCLEOSIDE RTIs | | |
| DELSTRIGO (doravirine/lamivudine/tenofovir) | ATRIPLA (efavirenz/emtricitabine/tenofovir) | |
| efavirenz/emtricitabine/tenofovir | CIMDUO (lamivudine/tenofovir) | |
| ODEFSEY (emtricitabine/rilpivirine/tenofovir) | COMPLERA (emtricitabine/rilpivirine/tenofovir) | |
| COMBINATION PRODUCTS – PROTEASE INHIBITORS | | |
| lopinavir/ritonavir | KALETRA (lopinavir/ritonavir) | |
| ENTRY INHIBITORS – CCR5 CO-RECEPTOR ANTAGONISTS | | |
| | maraviroc | |
| | SELZENTRY (maraviroc) | |
| ENTRY INHIBITORS – FUSION INHIBITORS | | |
| | FUZEON (enfuvirtide) | |
| INTEGRASE STRAND TRANSFER INHIBITORS | | |
| APRETUDE (cabotegravir) | cabotegravir ER | |
| ISENTRESS (raltegravir) | ISENTRESS HD (raltegravir) | |
| TIVICAY, TIVICAY PD (dolutegravir) | VOCABRIA (cabotegravir) | |
| NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS (NNRTI) | | |
| EDURANT (rilpivirine) | etravirine | |
| efavirenz | INTELENCE (etravirine) | |
| | nevirapine, nevirapine ER | |
| | PIFELTRO (doravirine) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|--|--|
| ANTIRETROVIRALS ^{DUR+} (continued) | | |
| NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS (NRTI) | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> 1 claim with the requested agent in the past 105 days <p>STRIBILD – MANUAL PA</p> <p>SUNLENCA</p> <ul style="list-style-type: none"> Requires clinical review <p>TYBOST – MANUAL PA</p> |
| abacavir | didanosine | |
| EMTRIVA (emtricitabine) | emtricitabine | |
| lamivudine | EPIVIR (lamivudine) | |
| ZIAGEN (abacavir) | RETROVIR (zidovudine) | |
| zidovudine | stavudine | |
| | VIREAD (tenofovir disoproxil fumarate) | |
| PHARMACOENHANCER – CYTOCHROME P450 INHIBITORS | | |
| | TYBOST (cobicistat) | |
| PROTEASE INHIBITORS (NON-PEPTIDIC) | | |
| PREZISTA (darunavir) | APTIVUS (tipranavir) | |
| | darunavir | |
| | PREZCOBIX (darunavir/cobicistat) | |
| PROTEASE INHIBITORS (PEPTIDIC) | | |
| atazanavir | fosamprenavir | |
| EVOTAZ (atazanavir/cobicistat) | LEXIVA (fosamprenavir) | |
| ritonavir | NORIVIR (ritonavir) | |
| | REYATAZ (atazanavir) | |
| | VIRACEPT (nelfinavir) | |
| SINGLE PRODUCT REGIMENS | | |
| BIKTARVY (bictegravir/emtricitabine/tenofovir) | efavirenz/lamivudine/tenofovir | |
| GENVOYA (elvitegravir/cobicistat/emtricitabine/ tenofovir alafenamide) | JULUCA (dolutegravir/rilpivirine) | |
| SYMFI (efavirenz/lamivudine/tenofovir) | rilpivirine ER | |
| SYMFI LO (efavirenz/lamivudine/tenofovir) | STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir disoproxil fumarate) | |
| TRIUMEQ (abacavir/dolutegravir/lamivudine) | SYMTUZA (darunavir/cobicistat/emtricitabine/tenofovir alafenamide) | |
| TRIUMEQ PD (abacavir/dolutegravir/lamivudine) | | |
| ANTIVIRALS, ORAL | | |
| ANTI-CYTOMEGALOVIRUS AGENTS | | <p>Valganciclovir solution</p> <ul style="list-style-type: none"> Automatic approval issued for 0-12 years of age <p>PREVYMIS</p> <ul style="list-style-type: none"> Requires clinical review |
| valganciclovir tablet | LIVTENCITY (maribavir) | |
| | PREVYMIS (letermovir) | |
| | VALCYTE (valganciclovir) | |
| | valganciclovir solution | |
| ANTI-HERPETIC AGENTS | | |
| acyclovir | SITAVIG (acyclovir) | |
| famciclovir | VALTrex (valacyclovir) | |
| valacyclovir | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| ANTIVIRALS, ORAL (continued) | | |
| ANTI-INFLUENZA AGENTS | | |
| oseltamivir | FLUMADINE (rimantadine) | |
| | RAPIVAB (peramivir) | |
| | RELENZA (zanamivir) | |
| | rimantadine | |
| | TAMIFLU (oseltamivir) | |
| | XOFLUZA (baloxavir) | |
| ANTIVIRALS, TOPICAL | | |
| ZOVIRAX (acyclovir) cream | acyclovir | |
| | DENAVIR (penciclovir) | |
| | penciclovir | |
| | XERESE (acyclovir/hydrocortisone) | |
| | ZOVIRAX (acyclovir) ointment | |
| AROMATASE INHIBITORS | | |
| anastrozole | ARIMIDEX (anastrozole) | |
| exemestane | AROMASIN (exemestane) | |
| letrozole | FEMARA (letrozole) | |
| ATOPIC DERMATITIS | | |
| ADBRY (tralokinumab-ldrm) | CIBINQO (abrocitinib) | Minimum Age Limit <ul style="list-style-type: none"> • 3 months: EUKRISA • 2 years: ELIDEL, PROTOPIC 0.03% • 12 years: OPZELURA • 16 years: PROTOPIC 0.1% |
| ADBRY Autoinjector (tralokinumab-ldrm) | EBGLYSS Pen (lebrikizumab-lbkz) ^{NR} | |
| DUPIXENT (dupilumab) ^{DUR+} | OPZELURA (ruxolitinib) | |
| ELIDEL (pimecrolimus) | ZORYVE (roflumilast) 0.15% cream | |
| EUKRISA (crisaborole) ^{DUR+} | | |
| pimecrolimus | | |
| tacrolimus | | |
| <p>ADBRY – MANUAL PA</p> <p>CIBINQO</p> <ul style="list-style-type: none"> • Requires clinical review <p>DUPIXENT</p> <ul style="list-style-type: none"> • 1 claim with DUPIXENT in the past 60 days OR • New starts require clinical review (see manual PA links below) <ul style="list-style-type: none"> ○ Asthma – MANUAL PA ○ Atopic Dermatitis – MANUAL PA ○ Eosinophilic Esophagitis – MANUAL PA ○ Nasal Polyposis – MANUAL PA ○ Prurigo Nodularis – MANUAL PA | | <p>EBGLYSS</p> <ul style="list-style-type: none"> • Requires clinical review <p>EUKRISA</p> <ul style="list-style-type: none"> • 30 days of therapy with a calcineurin inhibitor or topical steroid in the past 6 months <p>OPZELURA</p> <ul style="list-style-type: none"> • 30 days of therapy with ELIDEL, EUKRISA or PROTOPIC |
| See below for additional PA Criteria/DUR+ Rules | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|--|
| BETA BLOCKERS, ANTIANGINALS & SINUS NODE AGENTS DUR+ | | |
| ANTIANGINALS | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>COREG CR</p> <ul style="list-style-type: none"> • Documented diagnosis of hypertension AND • Have tried generic carvedilol AND 1 preferred agent in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>CORLANOR – MANUAL PA</p> <p>HEMANGEOL</p> <ul style="list-style-type: none"> • Documented diagnosis of infantile hemangioma <p>RANEXA</p> <ul style="list-style-type: none"> • Documented diagnosis of angina AND • 1 claim for a calcium channel blocker, beta-blocker, nitrate, or combination agent in the past 30 days OR • 90 days of therapy with the requested agent in the past 105 days |
| | ASPRUZYO SPRINKLE (ranolazine) ranolazine ER | |
| BETA- AND ALPHA-BLOCKERS | | |
| carvedilol | carvedilol ER | |
| labetalol | COREG (carvedilol) COREG CR (carvedilol) | |
| BETA-BLOCKER/DIURETIC COMBINATIONS | | |
| atenolol/chlorthalidone | TENORETIC (atenolol/chlorthalidone) | |
| bisoprolol/hydrochlorothiazide | ZIAC (bisoprolol/hydrochlorothiazide) | |
| metoprolol/hydrochlorothiazide | | |
| propranolol/hydrochlorothiazide | | |
| BETA-BLOCKERS | | |
| acebutolol | BETAPACE (sotalol) | |
| atenolol | BETAPACE AF (sotalol) | |
| bisoprolol | betaxolol | |
| HEMANGEOL (propranolol) | BYSTOLIC (nebivolol) | |
| metoprolol succinate | INDERAL LA (propranolol) | |
| metoprolol tartrate | INDERAL XL (propranolol) | |
| nadolol | INNOPRAN XL (propranolol) | |
| nebivolol | KAPSPARGO SPRINKLE (metoprolol succinate) | |
| pindolol | LOPRESSOR (metoprolol tartrate) | |
| propranolol | SOTYLIZE (sotalol) | |
| propranolol ER | TENORMIN (atenolol) | |
| SORINE (sotalol) | TOPROL XL (metoprolol succinate) | |
| sotalol | | |
| sotalol AF | | |
| timolol | | |
| SINUS NODE AGENTS | | |
| | CORLANOR (ivabradine) ivabradine | |
| BILE SALTS | | |
| ursodiol | BYLVAY (odevixibat) CHENODAL (chenodiol) IQIRVO (elafibranor) LIVDELZI (seladelpar) LIVMARLI (maralixibat) OCALIVA (obeticholic acid) RELTONE (ursodiol) URSO FORTE (ursodiol) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|---|
| BLADDER RELAXANT PREPARATIONS DUR+ | | |
| MYRBETRIQ (mirabegron) | darifenacin ER | Non-Preferred Criteria <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months |
| oxybutynin | DETROL (tolterodine) | |
| oxybutynin ER | DETROL LA (tolterodine) | |
| solifenacin | fesoterodine | |
| | GEMTESA (vibegron) | |
| | mirabegron ER | |
| | tolterodine | |
| | tolterodine ER | |
| | TOVIAZ (fesoterodine) | |
| | trospium | |
| | trospium ER | |
| | VESICARE (solifenacin) | |
| | VESICARE LS (solifenacin) | |
| BONE RESORPTION SUPPRESSION AND RELATED AGENTS DUR+ | | |
| BISPHOSPHONATES | | Non-Preferred Criteria <ul style="list-style-type: none"> Documented diagnosis of osteoporosis or osteopenia AND Have tried 2 different preferred agents in the past 6 months |
| alendronate tablet | ACTONEL (risedronate) | |
| ibandronate tablet | alendronate solution | |
| risedronate | ATELVIA (risedronate) | |
| | BINOSTO (alendronate) | |
| | FOSAMAX (alendronate) | |
| | FOSAMAX PLUS D (alendronate/vitamin D3) | |
| | ibandronate syringe/vial | |
| | risedronate DR | |
| OTHERS | | |
| FORTEO (teriparatide) | calcitonin salmon | |
| raloxifene | EVENITY (romosozumab-aqgg) | |
| | EVISTA (raloxifene) | |
| | MIACALCIN (calcitonin salmon) | |
| | PROLIA (denosumab) | |
| | teriparatide | |
| | TYMLOS (abaloparatide) | |
| | XGEVA (denosumab) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|--|
| BPH AGENTS ^{DUR+} | | |
| 5-ALPHA-REDUCTASE INHIBITORS | | <p>CARDURA, FLOMAX, PROSCAR, terazosin, or UROXATRAL – Female</p> <ul style="list-style-type: none"> Documented State-accepted diagnosis <p>Non-Preferred Criteria – Male</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days <p>ENTADFI</p> <ul style="list-style-type: none"> Requires clinical review |
| dutasteride | AVODART (dutasteride) | |
| finasteride | ENTADFI (finasteride/tadalafil) | |
| | PROSCAR (finasteride) | |
| ALPHA BLOCKERS | | |
| alfuzosin ER | CARDURA (doxazosin) | |
| doxazosin | CARDURA XL (doxazosin) | |
| tamsulosin | dutasteride/tamsulosin | |
| terazosin | FLOMAX (tamsulosin) | |
| | RAPAFLO (silodosin) | |
| | silodosin | |
| PHOSPHODIESTERASE TYPE 5 (PDE5) INHIBITORS | | |
| | CIALIS (tadalafil) | |
| | tadalafil | |
| BRONCHODILATORS & COPD AGENTS | | |
| ANTICHOLINERGIC-BETA AGONIST COMBINATIONS | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 6 years: SPIRIVA RESPIMAT <p>SPIRIVA RESPIMAT</p> <ul style="list-style-type: none"> Automatic approval issued for diagnosis of asthma for ≥ 6 years of age <p>BREZTRI AEROSPHERE</p> <ul style="list-style-type: none"> 3 claims with BREZTRI AEROSPHERE in the past 105 days OR New starts require clinical review |
| ANORO ELLIPTA (umeclidinium/vilanterol) | BEVESPI AEROSPHERE (glycopyrrolate/formoterol) | |
| COMBIVENT RESPIMAT (ipratropium/albuterol) | DUAKLIR PRESSAIR (aclidinium/formoterol) | |
| ipratropium/albuterol | | |
| STIOLTO RESPIMAT (tiotropium/olodaterol) | | |
| ANTICHOLINERGIC-BATA AGONIST-GLUCOCORTICOID COMBINATIONS | | |
| | BREZTRI AEROSPHERE (budesonide/glycopyrrolate/formoterol) ^{DUR+} | |
| | TRELEGY ELLIPTA (fluticasone/umeclidinium/vilanterol) | |
| ANTICHOLINERGICS AND COPD AGENTS | | |
| ATROVENT HFA (ipratropium) | DALIRESP (roflumilast) | |
| INCRUSE ELLIPTA (umeclidinium) | OHTUVAYRE (ensifentrine) | |
| ipratropium | roflumilast | |
| SPIRIVA HANDHALER (tiotropium) | SPIRIVA RESPIMAT (tiotropium) ^{DUR+} | |
| | tiotropium | |
| | TUDORZA PRESSAIR (aclidinium) | |
| | YUPLERI (revedfenacin) | |
| BRONCHODILATORS, BETA AGONISTS | | |
| INHALATION SOLUTION ^{DUR+} | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> 1 claim for a preferred agent in the past 6 months OR 3 claims with the requested agent in the past 105 days <p style="background-color: yellow;">See next page for additional PA Criteria/DUR+ Rules</p> |
| albuterol | arformoterol | |
| | BROVANA (arformoterol) | |
| | formoterol, formoterol fumarate ^{NR} | |
| | levalbuterol | |
| | PERFOROMIST (formoterol) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---------------------------------|---|
| BRONCHODILATORS, BETA AGONISTS <i>(continued)</i> | | |
| INHALERS, LONG ACTING ^{DUR+} | | <p style="background-color: yellow;">See previous page for additional PA Criteria/DUR+ Rules</p> <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 4 years: SEREVENT, XOPENEX HFA • 6 years: XOPENEX Solution • 18 years: AIRSUPRA, BROVANA, PERFORMOMIST, STRIVERDI RESPIMAT <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 2 inhalers: AIRSUPRA <p>AIRSUPRA and PROAIR DIGIHALER – Require clinical review</p> <p>XOPENEX HFA and Solution</p> <ul style="list-style-type: none"> • 1 claim for a preferred albuterol (inhaler or vials) in the past 30 days |
| SEREVENT DISKUS (salmeterol) | | |
| STRIVERDI RESPIMAT (olodaterol) | | |
| INHALERS, SHORT ACTING | | |
| albuterol HFA | AIRSUPRA (albuterol/budesonide) | |
| VENTOLIN HFA (albuterol) | levalbuterol HFA | |
| | PROAIR DIGIHALER (albuterol) | |
| | XOPENEX HFA (levalbuterol) | |
| ORAL | | |
| albuterol IR | albuterol ER | |
| terbutaline | | |
| | | |
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| CALCIUM CHANNEL BLOCKERS ^{DUR+} | | |
| LONG-ACTING | | <p>Quantity Limit (per 21 days)</p> <ul style="list-style-type: none"> • 252 tablets: nimodipine • 2520 mL: nimodipine <p>Non-Preferred Criteria – Long Acting</p> <ul style="list-style-type: none"> • Have tried 2 different preferred Long Acting CCB agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>Non-Preferred Criteria – Short Acting</p> <ul style="list-style-type: none"> • Have tried 2 different preferred Short Acting CCB agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>Nimodipine</p> <ul style="list-style-type: none"> • Documented diagnosis of subarachnoid hemorrhage in the past 45 days AND • Duration of therapy limited to 21 days |
| amlodipine | CARDIZEM CD (diltiazem) | |
| CARTIA XT (diltiazem) | CARDIZEM LA (diltiazem) | |
| diltiazem ER 24 HR | diltiazem ER 12 HR | |
| diltiazem CD 24 HR | diltiazem LA 24 HR | |
| diltiazem XR 24 HR | KATERZIA (amlodipine) | |
| DILT-XR 24 HR (diltiazem) | levamlodipine | |
| felodipine | MATZIM LA (diltiazem) | |
| nifedipine ER | nisoldipine | |
| TAZTIA XT (diltiazem) | NORVASC | |
| verapamil ER | PROCARDIA XL | |
| verapamil SR | SULAR (nisoldipine) | |
| | TIADYLT ER (diltiazem) | |
| | TIAZAC (diltiazem) | |
| | verapamil PM | |
| | VERELAN PM (verapamil) | |
| SHORT-ACTING | | |
| diltiazem | CARDIZEM (diltiazem) | |
| nicardipine | isradipine | |
| nifedipine | nimodipine | |
| verapamil | NORLIQVA (amlodipine) | |
| | NYMALIZE (nimodipine) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA | |
|--|---|--|--|
| CALORIC AGENTS | | | |
| BOOST BREAKFAST ESSENTIALS BRIGHT BEGINNINGS DUOCAL ENSURE NUTREN OSMOLITE PEDIASURE PROMOD RESOURCE TWOCAL HN | All non-preferred caloric/nutritional agents (which are all other products except those specifically listed as preferred) require a manual prior authorization. | Non-Preferred Agents – MANUAL PA | |
| CEPHALOSPORINS AND RELATED ANTIBIOTICS (ORAL) | | | |
| BETA LACTAM/BETA-LACTAMASE INHIBITOR COMBINATIONS | | Non-Preferred Criteria – All Cephalosporin Generations <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months Maximum Age Limit <ul style="list-style-type: none"> 18 years: cefdinir suspension | |
| amoxicillin/clavulanate | amoxicillin/clavulanate ER AUGMENTIN (amoxicillin/clavulanate) | | |
| CEPHALOSPORINS – FIRST GENERATION | | | |
| cefadroxil | cephalexin tablet | | |
| cephalexin capsule, suspension | | | |
| CEPHALOSPORINS – SECOND GENERATION | | | |
| cefaclor capsule | cefaclor ER | | |
| cefprozil | cefaclor suspension | | |
| cefuroxime | | | |
| CEPHALOSPORINS – THIRD GENERATION | | | |
| cefdinir | cefixime suspension | | |
| cefixime capsule | SUPRAX (cefixime) | | |
| cefpodoxime | | | |
| COLONY STIMULATING FACTORS | | | |
| FULPHILA (pegfilgrastim-jmdb) | FYLNETRA (pegfilgrastim-pbbk) | | |
| NEUPOGEN (filgrastim) | GRANIX (tbo-filgrastim) | | |
| | LEUKINE (sargramostim) | | |
| | NEULASTA, NEULASTA ONPRO (pegfilgrastim) | | |
| | NIVESTYM (filgrastim-aafi) | | |
| | NYVEPRIA (pegfilgrastim-ppgf) | | |
| | RELEUKO (filgrastim-ayow) | | |
| | ROLVEDON (eflapeggrastim-xnst) | | |
| | STIMUFEND (pegfilgrastim-fpgk) | | |
| | UDENYCA, UDENYCA ONBODY (pegfilgrastim-cbqv) | | |
| | ZARXIO (filgrastim-sndz) | | |
| | ZIEXTENZO (pegfilgrastim-bmez) | | |



MISSISSIPPI DIVISION OF MEDICAID UNIVERSAL PREFERRED DRUG LIST

EFFECTIVE 01/01/2025
Version 2025_2
Updated 02/01/2025

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| CYSTIC FIBROSIS AGENTS ^{DUR+} | | |
| PULMOZYME (dornase alfa) | ALYFTREK (vanzacaftor/tezacaftor/deutivacaftor) ^{NR} | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 1 month: KALYDECO granules 3 months: PULMOZYME 1 year: ORKAMBI 2 years: COLY-MYCIN M, TRIKAFTA granules 6 years: ALYFTREK, BETHKIS, KALYDECO tablet, KITABIS, SYMDEKO, TOBI, TOBI PODHALER, TRIKAFTA tablet 7 years: CAYSTON 18 years: BRONCHITOL <p>Maximum Age Limit</p> <ul style="list-style-type: none"> 2 years: ORKAMBI 75-94 mg granules 5 years: KALYDECO, ORKAMBI 100-125 mg granules, ORKAMBI 200-125 mg granules, TRIKAFTA granules 11 years: TRIKAFTA tablets <p>Preferred Agents</p> <ul style="list-style-type: none"> Documented diagnosis of Cystic Fibrosis OR Require clinical review <p>ALYFTREK and TOBI PODHALER – Require clinical review</p> <p>KALYDECO – MANUAL PA</p> <p>ORKAMBI – MANUAL PA</p> <p>SYMDEKO – MANUAL PA</p> <p>TRIKAFTA – MANUAL PA</p> |
| tobramycin (generic TOBI) | BETHKIS (tobramycin) | |
| | BRONCHITOL (mannitol) | |
| | CAYSTON (aztreonam) | |
| | colistimethate | |
| | COLY-MYCIN M (colistin) | |
| | KALYDECO (ivacaftor) | |
| | KITABIS (tobramycin) | |
| | ORKAMBI (lumacaftor/ivacaftor) | |
| | SYMDEKO (tezacaftor/ivacaftor) | |
| | TOBI (tobramycin) | |
| | TOBI PODHALER (tobramycin) | |
| | tobramycin (generic BETHKIS & KITABIS) | |
| | TRIKAFTA (elxacaftor/tezacaftor/ivacaftor) | |
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| CYTOKINE & CAM ANTAGONISTS ^{DUR+} | | |
| ACTEMRA (tocilizumab) syringe, vial | ABRILADA (adalimumab-afzb) | <p>Preferred Agents – Criteria details found here</p> <p>Non-Preferred Agents</p> <ul style="list-style-type: none"> Require clinical review <p>IV Administered Agents</p> <ul style="list-style-type: none"> Require clinical review |
| AVSOLA (infliximab-axxq) | ACTEMRA ACTPEN (tocilizumab) | |
| ENBREL (etanercept) | adalimumab-aacf | |
| HUMIRA (adalimumab) | adalimumab-aaty | |
| KINERET (anakinra) | adalimumab-adaz | |
| methotrexate | adalimumab-adbm | |
| OLUMIANT (baricitinib) | adalimumab-fkjp | |
| OTEZLA (apremilast) | adalimumab-ryvk | |
| RINVOQ (upadacitinib) | AMJEVITA (adalimumab-atto) | |
| RINVOQ LQ (upadacitinib) | ARCALYST (riloncept) | |
| SIMPONI (golimumab) | BIMZELX (bimekizumab-bkzx) | |
| TALTZ (ixekizumab) | CIMZIA (certolizumab) | |
| TYENNE Syringe, Vial (tocilizumab-aazg) | COSENTYX (secukinumab) | |
| XELJANZ (tofacitinib) tablet | CYLTEZO (adalimumab-adbm) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|--|
| CYTOKINE & CAM ANTAGONISTS^{DUR+} <i>(continued)</i> | | |
| | ENTYVIO (vedolizumab) | <p>Preferred Agents – Criteria details found here</p> <p>Non-Preferred Agents</p> <ul style="list-style-type: none"> Require clinical review <p>IV Administered Agents</p> <ul style="list-style-type: none"> Require clinical review |
| | HADLIMA (adalimumab-bwwd) | |
| | HULIO (adalimumab-fkjp) | |
| | HYRIMOZ (adalimumab-adaz) | |
| | IDACIO (adalimumab-aacf) | |
| | ILARIS (canakinumab) | |
| | ILUMYA (tildrakizumab-asmn) | |
| | INFLECTRA (infliximab-dyyb) | |
| | infliximab | |
| | JYLAMVO (methotrexate) | |
| | KEVZARA (sarilumab) | |
| | LITFULO (ritlectinib) | |
| | NEMLUVIO (nemolizumab-ilto) ^{NR} | |
| | OMVOH (mirikizumab-mrkz) | |
| | ORENCIA (abatacept) | |
| | OTREXUP (methotrexate) | |
| | RASUVO (methotrexate) | |
| | REMICADE (infliximab) | |
| | RENFLEXIS (infliximab-abda) | |
| | SILIQ (brodalumab) | |
| | SIMLANDI (adalimumab-ryvk) | |
| | SIMPONI ARIA (golimumab) | |
| | SKYRIZI (risankizumab-rzaa) | |
| | SOTYKTU (deucravacitinib) | |
| | SPEVIGO (spesolimab-sbzo) | |
| | STELARA (ustekinumab) | |
| | TOFIDENCE (tocilizumab-bavi) | |
| | TREMFYA (guselkumab) | |
| | TREXALL (methotrexate) | |
| | TYENNE Autoinjector (tocilizumab-aazg) | |
| | ustekinumab-kfce ^{NR} | |
| | XATMEP (methotrexate) | |
| | XELJANZ (tofacitinib) solution | |
| | XELJANZ XR (tofacitinib) | |
| | YUFLYMA (adalimumab-aaty) | |
| | YUSIMRY (adalimumab-aqvh) | |
| | ZYMFENTRA (infliximab-dyyb) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| ERYTHROPOIESIS STIMULATING PROTEINS DUR+ | | |
| EPOGEN (epoetin alfa) | ARANESP (darbepoetin alfa) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented diagnosis of cancer or chronic renal failure OR • Antineoplastic therapy in the past 6 months AND • Have tried a preferred RETACRIT or EPOGEN in the past 6 months OR • 1 claim for the requested agent in the past 105 days <p>JESDUVROQ</p> <ul style="list-style-type: none"> • Requires clinical review <p>MIRCERA</p> <ul style="list-style-type: none"> • Documented diagnosis of chronic renal failure in the past 2 years |
| MIRCERA (methoxy polyethylene glycol-epoetin-beta) | JESDUVROQ (daprodustat) | |
| RETACRIT (epoetin alfa-epbx) | PROCRIT (epoetin alfa) | |
| | VAFSEO (vadadustat) | |
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| FACTOR DEFICIENCY PRODUCTS DUR+ | | |
| FACTOR VIII | | <p>HEMLIBRA</p> <ul style="list-style-type: none"> • 3 claims with HEMLIBRA in the past 105 days OR • New starts require clinical review – MANUAL PA |
| ADVATE | ADYNOVATE | |
| AFSTYLA | ELOCTATE | |
| ALPHANATE | ESPEROCT | |
| ALTUVIIIIO | JIVI | |
| FEIBA | KCENTRA | |
| HEMOPIL M | OBIZUR | |
| HUMATE-P | VONVENDI | |
| KOATE | | |
| KOGENATE FS | | |
| KOVALTRY | | |
| NOVOEIGHT | | |
| NUWIQ | | |
| RECOMBINATE | | |
| WILATE | | |
| XYNTHA, XYNTHA SOLOFUSE | | |
| FACTOR IX | | |
| ALPHANINE SD | BEQVEZ | |
| ALPROLIX | REBINYN | |
| BENEFIX | | |
| IDELVION | | |
| IXINITY | | |
| PROFILNINE | | |
| RIXUBIS | | |
| OTHER HEMOPHILIA PRODUCTS | | |
| COAGADEX (factor X) | ALHEMO (concizumab-mtci) ^{NR} | |
| FIBRYGA (fibrinogen) | CORIFACT (factor XIII) ^{NR} | |
| HEMLIBRA (emicizumab-kxwh) DUR+ | HYMPAVZI (marstacimab-hncg) ^{NR} | |
| RIASTAP (fibrinogen) | NOVOSEVEN RT (factor VII) | |
| | SEVENFACT (factor VII) | |
| | TRETTEN (factor XIII) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|-------------------------------------|--|
| FIBROMYALGIA/NEUROPATHIC PAIN AGENTS | | |
| duloxetine | CYMBALTA (duloxetine) | |
| gabapentin | DIRZALMA SPRINKLE (duloxetine) | |
| pregabalin | gabapentin ER | |
| SAVELLA (milnacipran) | GABARONE (gabapentin) ^{NR} | |
| | GRALISE (gabapentin) | |
| | HORIZANT (gabapentin enacarbil) | |
| | LYRICA, LYRICA CR (pregabalin) | |
| | NEURONTIN (gabapentin) | |
| | pregabalin ER | |
| FLUOROQUINOLONES ^{DUR+} | | |
| ciprofloxacin tablet | BAXDELA (delafloxacin) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> 1 claim for a preferred agent in the past 30 days <p>CIPRO Suspension Criteria for Age < 12 Years</p> <ul style="list-style-type: none"> Anthrax infection or exposure, cystic fibrosis, pneumonic plague, or tularemia AND History of doxycycline in the past 3 months OR 7 days of therapy with a preferred agent from 2 of the classes below in the past 3 months: <ul style="list-style-type: none"> Penicillin 2nd or 3rd generation cephalosporin Macrolide <p>LEVAQUIN Solution Criteria for Age < 12 Years</p> <ul style="list-style-type: none"> Anthrax infection or exposure AND CIPRO suspension in the past 3 months OR 7 days of therapy with a preferred agent from 2 of the classes below in the past 3 months: <ul style="list-style-type: none"> Penicillin 2nd or 3rd generation cephalosporin Macrolide |
| levofloxacin tablet | CIPRO (ciprofloxacin) | |
| | ciprofloxacin suspension | |
| | levofloxacin solution | |
| | moxifloxacin | |
| | ofloxacin | |
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| GAUCHER'S DISEASE | | |
| ELELYSO (taliglucerase alfa) | CERDELGA (eliglustat) | |
| ZAVESCA (miglustat) | CEREZYME (imiglucerase) | |
| | miglustat | |
| | VPRIV (velaglucerase alfa) | |
| | YARGESA (miglustat) | |
| GENITAL WARTS & ACTINIC KERATOSIS AGENTS | | |
| CONDYLOX (podofilox) | CARAC (fluorouracil) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 12 years: ALDARA, ZYCLARA 18 years: CONDYLOX, PICATO, VEREGEN |
| fluorouracil | EFUDEX (fluorouracil) | |
| imiquimod | VEREGEN (sinecatechins) | |
| podofilox | ZYCLARA (imiquimod) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| GI ULCER THERAPIES | | |
| H2 RECEPTOR ANTAGONISTS | | <p>Prilosec suspension</p> <ul style="list-style-type: none"> Automatic approval issued for 0-2 years of age |
| famotidine | cimetidine | |
| | nizatidine | |
| | PEPCID (famotidine) | |
| OTHERS | | |
| CARAFATE (sucralfate) suspension | CARAFATE (sucralfate) tablet | |
| misoprostol | CYTOTEC (misoprostol) | |
| sucralfate | DARTISLA (glycopyrrolate) | |
| | VOQUEZNA (vonoprazan) | |
| PROTON PUMP INHIBITORS | | |
| esomeprazole capsule | DEXILANT (dexlansoprazole) | |
| NEXIUM (esomeprazole) packet | dexlansoprazole | |
| omeprazole | esomeprazole packet | |
| pantoprazole | KONVOMEF (omeprazole/sodium bicarbonate) | |
| | lansoprazole Rx | |
| | NEXIUM (esomeprazole) capsule | |
| | omeprazole/sodium bicarbonate | |
| | PREVACID (lansoprazole) | |
| | PRILOSEC (omeprazole) packet | |
| | PROTONIX (pantoprazole) | |
| | rabeprazole | |
| | ZEGERID (omeprazole/sodium bicarbonate) | |
| GLUCOCORTICOIDS (INHALED) | | |
| GLUCOCORTICOIDS | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Glucocorticoids <ul style="list-style-type: none"> 2 preferred single-entity agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days Glucocorticoid/Bronchodilator Combinations <ul style="list-style-type: none"> 2 preferred combination agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days Note: <ul style="list-style-type: none"> Institutional-sized products are non-preferred <p>AIRDUO DIGIHALER</p> <ul style="list-style-type: none"> Requires clinical review <p>ARMONAIR DIGIHALER</p> <ul style="list-style-type: none"> Require clinical review |
| ASMANEX (mometasone) | ALVESCO (ciclesonide) | |
| budesonide 0.25 mg and 0.5 mg | ARMONAIR DIGIHALER (fluticasone) | |
| FLOVENT DISKUS (fluticasone) | ARNUIITY ELLIPTA (fluticasone) | |
| PULMICORT FLEXHALER (budesonide) | ASMANEX HFA (mometasone) | |
| QVAR REDIHALER (beclomethasone) | budesonide 1 mg | |
| | FLOVENT HFA (fluticasone) | |
| | fluticasone diskus | |
| | fluticasone HFA | |
| | PULMICORT (budesonide) nebulizer solution | |
| GLUCOCORTICOID/BRONCHODILATOR COMBINATIONS | | |
| ADVAIR DISKUS (fluticasone/salmeterol) | AIRDUO DIGIHALER (fluticasone/salmeterol) | |
| ADVAIR HFA (fluticasone/salmeterol) | BREO ELLIPTA (fluticasone/vilanterol) | |
| DULERA (mometasone/formoterol) | BREYNA (budesonide/formoterol) | |
| fluticasone/salmeterol diskus | budesonide/formoterol | |
| fluticasone/salmeterol HFA | fluticasone/vilanterol | |
| SYMBICORT (budesonide/formoterol) | WIXELA INHUB (fluticasone/salmeterol) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| GROWTH HORMONES ^{DUR+} | | |
| GENOTROPIN (somatropin) | HUMATROPE (somatropin) | <p>All Agents</p> <ul style="list-style-type: none"> • Age ≥ 18 years <ul style="list-style-type: none"> ○ Documented diagnosis of craniopharyngioma, panhypopituitarism, Prader-Willi Syndrome, Turner Syndrome or an approvable adult diagnosis OR ○ Documented procedure of cranial irradiation • Age < 18 years <ul style="list-style-type: none"> ○ Documented diagnosis of idiopathic short stature AND ○ Documented approvable pediatric diagnosis OR ○ Documented approvable pediatric diagnosis <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 3 years: NGENLA • 18 years: SKYTROFA <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 18 years: NGENLA <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented approvable diagnosis for age as above AND • Have tried 1 preferred agent in the past 6 months OR • 84 days of therapy with the requested agent in the past 105 days <p>SKYTROFA</p> <ul style="list-style-type: none"> • ≥ 18 years AND • No history of diagnosis of Prader-Willi Syndrome AND • 28 days of therapy with a preferred short-acting growth hormone in the past 105 days |
| NORDITROPIN FLEXPPO (somatropin) | NGENLA (somatrogon-ghla) | |
| SKYTROFA (lonapegsomatropin-tcgd) | OMNITROPE (somatropin) | |
| | SEROSTIM (somatropin) | |
| | SOGROYA (somapacitan-beco) | |
| | VOXZOGO (vosoritide) | |
| | ZOMACTON (somatropin) | |
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| H. PYLORI COMBINATION TREATMENTS | | |
| PYLERA (bismuth subcitrate potassium/metronidazole/tetracycline) | bismuth subcitrate potassium/metronidazole/tetracycline lansoprazole/amoxicillin/clarithromycin OMECLAMOX (omeprazole/clarithromycin/amoxicillin) TALICIA (omeprazole/amoxicillin/rifabutin) VOQUEZNA DUAL PAK (vonoprazan/amoxicillin) VOQUEZNA TRIPLE PAK (vonoprazan/amoxicillin/clarithromycin) | <p>Quantity Limit</p> <ul style="list-style-type: none"> • 1 treatment course/year: all agents |
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| HEPATITIS B TREATMENTS | | |
| entecavir | adefovir dipivoxil | |
| lamivudine HBV | BARACLUDGE (entecavir) | |
| tenofovir disoproxil fumarate | VEMLIDY (tenofovir alafenamide) | |
| | VIREAD (tenofovir disoproxil fumarate) | |
| | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
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| HEPATITIS C TREATMENTS | | |
| MAVYRET (glecaprevir/pibrentasvir) [∞] | EPCLUSA (sofosbuvir/velpatasvir) [∞] | <p>∞ EPCLUSA, HARVONI, MAVYRET, SOVALDI, VOSEVI, ZEPATIER</p> <ul style="list-style-type: none"> Require MANUAL PA <p>Note:</p> <ul style="list-style-type: none"> EPCLUSA, HARVONI, MAVYRET and SOVALDI have FDA-approved pediatric indications |
| PEGASYS (peginterferon alfa-2a) | HARVONI (ledipasvir/sofosbuvir) [∞] | |
| ribavirin tablet | ledipasvir/sofosbuvir [∞] | |
| sofosbuvir/velpatasvir | ribavirin capsule | |
| | SOVALDI (sofosbuvir) [∞] | |
| | VIEKIRA PAK (ombitasvir/paritaprevir/ritonavir) | |
| | VOSEVI (sofosbuvir/velpatasvir/voxilaprevir) [∞] | |
| | ZEPATIER (elbasvir/grazoprevir) [∞] | |
| HEREDITARY ANGIOEDEMA | | |
| BERINERT (C1 esterase inhibitor) | CINRYZE (C1 esterase inhibitor) | |
| icatibant | FIRAZYR (icatibant) | |
| | KALBITOR (ecallantide) | |
| | ORLADEYO (berotralstat) | |
| | RUCONEST (C1 esterase inhibitor) | |
| | SAJAZIR (icatibant) | |
| | TAKHZYRO (lanadelumab-flyo) | |
| HYPERURICEMIA & GOUT ^{DUR+} | | |
| allopurinol | ALOPRIM (allopurinol) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months |
| colchicine tablet | colchicine capsule | |
| probenecid | COLCRYS (colchicine) | |
| probenecid/colchicine | febuxostat | |
| | GLOPERBA (colchicine) | |
| | MITIGARE (colchicine) | |
| | ULORIC (febuxostat) | |
| | ZYLOPRIM (allopurinol) | |
| HYPOGLYCEMIA TREATMENT | | |
| BAQSIMI (glucagon) | GVOKE (glucagon) ^{Step Edit} | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 2 years: GVOKE 4 years: BAQSIMI 6 years: ZEGALOGUE <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> 2 packs (or kits): BAQSIMI, glucagon, GVOKE, ZEGALOGUE <p>Non-Preferred Criteria – GVOKE</p> <ul style="list-style-type: none"> 1 claim with preferred BAQSIMI or ZEGALOGUE in the past 30 days |
| GLUCAGEN (glucagon) | | |
| glucagon emergency kit | | |
| glucagon vial | | |
| ZEGALOGUE (dasiglucagon) | | |
| | | |
| | | |
| HYPOGLYCEMICS, BIGUANIDES | | |
| metformin | GLUMETZA (metformin) | |
| metformin ER (generic GLUCOPHAGE XR) | metformin ER (generic FORTAMET) | |
| | metformin ER (generic GLUMETZA) | |
| | metformin solution | |
| | RIOMET (metformin) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|--|---|
| HYPOGLYCEMICS, DPP4s AND COMBINATIONS DUR+ | | |
| JANUMET (sitagliptin/metformin) | alogliptin | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred DPP4 agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days <p><u>Note:</u></p> <ul style="list-style-type: none"> Concomitant use of a GLP-1 agent and a DPP-4 agent requires clinical review. |
| JANUMET XR (sitagliptin/metformin) | alogliptin/metformin | |
| JANUVIA (sitagliptin) | JENTADUETO XR (linagliptin/metformin) | |
| JENTADUETO (linagliptin/metformin) | KAZANO (alogliptin/metformin) | |
| TRADJENTA (linagliptin) | KOMBIGLYZE XR (saxagliptin/metformin) | |
| | NESINA (alogliptin) | |
| | ONGLYZA (saxagliptin) | |
| | OSENI (alogliptin/pioglitazone) | |
| | saxagliptin | |
| | saxagliptin/metformin ER | |
| | sitagliptin | |
| | sitagliptin/metformin | |
| | ZITUVIMET (sitagliptin/metformin) ^{NR} | |
| | ZITUVIMET XR (sitagliptin/metformin) ^{NR} | |
| | ZITUVIO (sitagliptin) | |
| HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS DUR+ | | |
| BYETTA (exenatide) | BYDUREON (exenatide) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 10 years: BYDUREON BCISE, TRULICITY, VICTOZA 18 years: BYETTA, MOUNJARO, OZEMPIC, RYBELSUS <p>Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of Type 2 Diabetes and no history of SAXENDA or WEGOVY in the past 30 days OR No documented diagnosis for Type 2 Diabetes and 84 days of therapy with the requested agent in the past 105 days <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of Type 2 Diabetes AND No history of SAXENDA or WEGOVY in the past 30 days AND 84 days of therapy with TRULICITY in the past 6 months AND 84 days of therapy with either BYETTA or VICTOZA in the past 6 months OR 84 days of therapy with the requested agent in the past 105 days <p><u>Note:</u></p> <ul style="list-style-type: none"> Concomitant use of a GLP-1 agonist and a DPP-4 agent requires clinical review. Please see the PDL category Anti-obesity Select Agents for a list of covered agents. |
| TRULICITY (dulaglutide) | exenatide | |
| VICTOZA (liraglutide) | liraglutide | |
| | MOUNJARO (tirzepatide) | |
| | OZEMPIC (semaglutide) | |
| | RYBELSUS (semaglutide) | |
| | SOLIQUA (insulin glargine/lixisenatide) | |
| | SYMLINPEN (pramlintide) | |
| | XULTOPHY (insulin degludec/liraglutide) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|---|
| HYPOGLYCEMICS, INSULINS & RELATED AGENTS DUR+ | | |
| HUMALOG MIX 75/25 (insulin lispro/lispro protamine) | ADMELOG (insulin lispro) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented diagnosis of Diabetes Mellitus AND • Have tried 1 preferred agent in the past 6 months OR • 1 claim with the requested agent in the past 105 days <p>Quantity Limit</p> <ul style="list-style-type: none"> • Insulin quantity limits can be found here <p>Note:</p> <ul style="list-style-type: none"> • Insulin pen formulations are not covered for Long Term Care (LTC) beneficiaries. |
| HUMULIN 70/30 (insulin NPH/regular) | AFREZZA (insulin regular) | |
| HUMULIN N (insulin NPH) | APIDRA (insulin glulisine) | |
| HUMULIN R (insulin regular) | BASAGLAR (insulin glargine) | |
| HUMULIN R U-500 (insulin regular) | FIASP (insulin aspart/niacinamide) | |
| insulin aspart | HUMALOG; HUMALOG JUNIOR, KWIKPEN, TEMPO | |
| insulin aspart protamine mix 70/30 | PEN (insulin lispro) | |
| insulin lispro | HUMALOG MIX KWIKPEN 50/50, 75/25 (insulin lispro/lispro protamine) | |
| insulin lispro protamine mix 75/25 | HUMULIN 70/30 (insulin NPH/regular) | |
| LANTUS (insulin glargine) | HUMULIN N KWIKPEN (insulin NPH) | |
| TOUJEO (insulin glargine) | insulin degludec | |
| TOUJEO MAX (insulin glargine) | insulin glargine | |
| | insulin glargine-yfqn | |
| | LEVEMIR (insulin detemir) | |
| | LYUMJEV (insulin lispro-aabc) | |
| | NOVOLIN 70/30 (insulin NPH/regular) | |
| | NOVOLIN R (insulin regular) | |
| | NOVOLOG (insulin aspart) | |
| | NOVOLOG MIX 70/30 (insulin aspart/aspart protamine) | |
| | REZVOGLAR (insulin glargine-aglr) | |
| | SEMGLEE (insulin glargine-yfqn) | |
| | TRESIBA (insulin degludec) | |
| HYPOGLYCEMICS, MEGLITINIDES DUR+ | | |
| nateglinide | | |
| repaglinide | | |
| HYPOGLYCEMICS, SODIUM GLUCOSE COTRANSPORTER-2 (SGLT-2) INHIBITORS DUR+ | | |
| SGLT-2 INHIBITORS | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred SGLT-2 inhibitors in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days |
| FARXIGA (dapagliflozin) | dapagliflozin | |
| JARDIANCE (empagliflozin) | INPEFA (sotagliflozin) | |
| | INVOKANA (canagliflozin) | |
| | STEGLATRO (ertugliflozin) | |
| SGLT-2 INHIBITOR COMBINATIONS | | |
| GLYXAMBI (empagliflozin/linagliptin) | dapagliflozin/metformin ER | |
| SYNJARDY (empagliflozin/metformin) | INVOKAMET (canagliflozin/metformin) | |
| SYNJARDY XR (empagliflozin/metformin) | INVOKAMET XR (canagliflozin/metformin) | |
| TRIJARDY XR (empagliflozin/linagliptin/metformin) | QTERN (dapagliflozin/saxagliptin) | |
| | SEGLUROMET (ertugliflozin/metformin) | |
| | STEGLUJAN (ertugliflozin/sitagliptin) | |
| | XIGDUO XR (dapagliflozin/metformin) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---------------------------------------|--|
| HYPOGLYCEMICS, SULFONYLUREAS | | |
| glimepiride | | |
| glipizide | | |
| glipizide ER | | |
| glipizide XL | | |
| glyburide | | |
| glyburide micronized | | |
| HYPOGLYCEMICS, THIAZOLIDINEDIONES (TZDs) and TZD Combinations | | |
| pioglitazone | ACTOPLUS MET (pioglitazone/metformin) | |
| pioglitazone/metformin | ACTOS (pioglitazone) | |
| | DUETACT (pioglitazone/metformin) | |
| IDIOPATHIC PULMONARY FIBROSIS ^{DUR+} | | |
| OFEV (nintedanib) | ESBRIET (pirfenidone) | All Agents • Documented diagnosis of Idiopathic Pulmonary Fibrosis |
| | pirfenidone | |
| IMMUNE GLOBULINS | | |
| BIVIGAM | ALYGLO | |
| FLEBOGAMMA | ASCENIV | |
| GAMASTAN | CABLIVI | |
| GAMMAGARD | CUTAQUIG | |
| GAMMAGARD S-D | CUVITRU | |
| GAMUNEX-C | GAMMAKED | |
| HIZENTRA | GAMMAPLEX | |
| HYQVIA | OCTAGAM | |
| PANZYGA | | |
| PRIVIGEN | | |
| XEMBIFY | | |
| IMMUNOLOGIC THERAPIES FOR ASTHMA | | |
| DUPIXENT (dupilumab) ^{DUR+} | CINQAIR (reslizumab) | CINQAIR • Requires clinical review See below for additional PA Criteria/DUR+ Rules |
| FASENRA (benralizumab) | NUCALA (mepolizumab) | |
| XOLAIR (omalizumab) | TEZSPIRE (tezepelumab-ekko) | |
| DUPIXENT • 1 claim with DUPIXENT in the past 60 days OR • New starts require clinical review (see manual PA links below) <ul style="list-style-type: none"> ○ Asthma – MANUAL PA ○ Atopic Dermatitis – MANUAL PA ○ Eosinophilic Esophagitis – MANUAL PA ○ Nasal Polyposis – MANUAL PA ○ Prurigo Nodularis – MANUAL PA | | |
| FASENRA • Requires clinical review – MANUAL PA | | |
| NUCALA • Requires clinical review | | |
| TEZSPIRE • Requires clinical review | | |
| XOLAIR • 1 claim with XOLAIR in the past 45 days OR • New starts require clinical review – MANUAL PA | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---------------------------|--|
| IMMUNOSUPPRESSIVE AGENTS, ORAL | | |
| AZASAN (azathioprine) | ASTAGRAF XL (tacrolimus) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 13 years: RAPAMUNE • 18 years: ZORTRESS <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 12 years: PROGRAF Granules <p style="background-color: yellow;">See below for additional PA Criteria/DUR+ Rules</p> |
| azathioprine | ENVARUSUS XR (tacrolimus) | |
| CELLCEPT (mycophenolate) | MYFORTIC (mycophenolate) | |
| cyclosporine | PROGRAF (tacrolimus) | |
| everolimus | REZUROCK (belumosudil) | |
| mycophenolate | ZORTRESS (everolimus) | |
| mycophenolic acid | | |
| NEORAL (cyclosporine) | | |
| RAPAMUNE (sirolimus) | | |
| SANDIMMUNE (cyclosporine) | | |
| sirolimus | | |
| tacrolimus | | |
| <p>Preferred Criteria</p> <ul style="list-style-type: none"> • AZASAN <ul style="list-style-type: none"> ◦ Documented diagnosis of kidney transplant, RA, or a State-accepted diagnosis • CELLCEPT <ul style="list-style-type: none"> ◦ Documented diagnosis of heart, kidney, or liver transplant or a State-accepted diagnosis • GENGRAF, NEORAL, SANDIMMUNE <ul style="list-style-type: none"> ◦ Documented diagnosis of heart transplant, kidney transplant, liver transplant, psoriasis, RA, or a State-accepted diagnosis • Everolimus <ul style="list-style-type: none"> ◦ Documented diagnosis of kidney or liver transplant • RAPAMUNE <ul style="list-style-type: none"> ◦ Documented diagnosis of kidney transplant • Tacrolimus <ul style="list-style-type: none"> ◦ Documented diagnosis of heart, kidney, liver, or lung transplant or a State-accepted diagnosis <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • MYHIBBIN Suspension <ul style="list-style-type: none"> ◦ Documented diagnosis of heart, kidney, or liver transplant or a State-accepted diagnosis AND ◦ 30 days of therapy with mycophenolate suspension in the past 105 days OR ◦ 90 days of therapy with MYHIBBIN Suspension in the past 105 days • ASTAGRAF XR or ENVARUSUS XR <ul style="list-style-type: none"> ◦ Documented diagnosis of heart, kidney, liver, or lung transplant or a State-accepted diagnosis AND ◦ 30 days of therapy with tacrolimus IR in the past 105 days OR ◦ 90 days of therapy with the requested agent in the past 105 days • PROGRAF Granules <ul style="list-style-type: none"> ◦ Age ≤ 11 years AND ◦ Documented diagnosis of heart, kidney, liver, or lung transplant or a State-accepted diagnosis • MYFORTIC <ul style="list-style-type: none"> ◦ Documented diagnosis of kidney transplant or psoriasis • ZORTRESS <ul style="list-style-type: none"> ◦ Documented diagnosis of kidney or liver transplant | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|---|
| INTRANASAL RHINITIS AGENTS | | |
| ANTICHOLINERGICS | | <p>Non-Preferred Criteria – Corticosteroids</p> <ul style="list-style-type: none"> Documented diagnosis of allergic rhinitis AND Have tried 1 different preferred agent in the past 6 months |
| ipratropium | | |
| ANTI-HISTAMINE/CORTICOSTEROID COMBINATIONS | | |
| | azelastine/fluticasone | |
| | DYMISTA (azelastine/fluticasone) | |
| | RYALTRIS (olopatadine/mometasone) | |
| ANTI-HISTAMINES | | |
| azelastine | olopatadine | |
| | PATANASE (olopatadine) | |
| CORTICOSTEROIDS | | |
| fluticasone | BECONASE AQ (beclomethasone) | |
| | flunisolide | |
| | mometasone | |
| | NASONEX (mometasone) | |
| | OMNARIS (ciclesonide) | |
| | QNASL (beclomethasone) | |
| | XHANCE (fluticasone) | |
| | ZETONNA (ciclesonide) | |
| IRON CHELATING AGENTS | | |
| deferasirox (all manufacturers except those listed as non-preferred) | deferasirox (manufacturers starting with 45963, 62332) | JADENU – MANUAL PA |
| deferiprone 500 mg tablet | deferiprone 1,000 mg tablet | |
| FERRIPROX (deferiprone) | EXJADE (deferasirox) | |
| | JADENU, JADENU SPRINKLE (deferasirox) | |
| IRRITABLE BOWEL SYNDROME/SHORT BOWEL SYNDROME AGENTS/SELECTED AGENTS ^{DUR+} | | |
| IRRITABLE BOWEL SYNDROME CONSTIPATION ^{DUR+} | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 1 year: GATTEX 6 years: LINZESS 72 mcg 18 years: AMITIZA, IBSRELA, LINZESS 145 mcg & 290 mcg, MOTTEGRITY, MOVANTIK, MYTESI, RELISTOR, SYMPROIC, TRULANCE, VIBERZI <p>Gender Limit</p> <ul style="list-style-type: none"> Female – AMITIZA 8 mcg <p style="background-color: yellow;">See next page for additional PA Criteria/DUR+ Rules</p> |
| LINZESS (linaclotide) | AMITIZA (lubiprostone) | |
| lubiprostone | IBSRELA (tenapanor) | |
| TRULANCE (plecanatide) | MOTTEGRITY (prucalopride) | |
| | MOVANTIK (naloxegol) | |
| | prucalopride ^{NR} | |
| | RELISTOR (methylnaltrexone) | |
| | SYMPROIC (naldemedine) | |
| IRRITABLE BOWEL SYNDROME DIARRHEA | | |
| dicyclomine | alosetron | |
| ED-SPAZ (hyoscyamine) | LOTRONEX (alosetron) ^{DUR+} | |
| hyoscyamine, hyoscyamine ER | VIBERZI (eluxadoline) ^{DUR+} | |
| HYOSYNE (hyoscyamine) | | |
| LEVSIN, LEVSIN-SL (hyoscyamine) | | |
| NULEV (hyoscyamine) | | |
| OSCIMIN, OSCIMIN SL (hyoscyamine) | | |
| SHORT BOWEL SYNDROME AND SELECTED GI AGENTS ^{DUR+} | | |
| | GATTEX (teduglutide) | |
| | MYTESI (crofelemer) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|--|--|
| IRRITABLE BOWEL SYNDROME/SHORT BOWEL SYNDROME AGENTS/SELECTED AGENTS ^{DUR+} <i>(continued)</i> | | |
| See previous page for additional PA Criteria/DUR+ Rules | | |
| IRRITABLE BOWEL SYNDROME – CONSTIPATION ^{DUR+} | | |
| <p>Chronic Idiopathic Constipation (CIC): Amitiza 24 mcg, LINZESS 72 mcg, LINZESS 145 mcg, MOTEGRITY, TRULANCE</p> <ul style="list-style-type: none"> • Preferred CIC Agents <ul style="list-style-type: none"> ○ Documented diagnosis of CIC in the past year AND ○ No history of GI or bowel obstruction • LINZESS 72 mcg <ul style="list-style-type: none"> ○ Age 6-17 years AND ○ Documented diagnosis of CIC or pediatric functional constipation in the past year AND ○ No history of GI or bowel obstruction • Non-Preferred CIC Agents <ul style="list-style-type: none"> ○ Documented diagnosis of CIC AND ○ No history of GI or bowel obstruction AND ○ Have tried 2 preferred CIC agents in the past 6 months OR ○ 1 claim with the requested agent in the past 105 days | <p>Irritable Bowel Syndrome – Constipation Dominant (IBS-C): AMITIZA 8 mcg, IBSRELA, LINZESS 290 mcg, TRULANCE</p> <ul style="list-style-type: none"> • Preferred IBS-C Agents <ul style="list-style-type: none"> ○ Documented diagnosis of IBS-C in the past year AND ○ No history of GI or bowel obstruction • Non-Preferred IBS-C Agents <ul style="list-style-type: none"> ○ Documented diagnosis of IBS-C in the past year AND ○ No history of GI or bowel obstruction AND ○ Have tried 2 preferred IBS-C agents in the past 6 months OR ○ 1 claim with the requested agent in the past 105 days | <p>Opioid Induced Constipation (OIC): AMITIZA 24 mcg, MOVANTIK, RELISTOR, SYMPROIC</p> <ul style="list-style-type: none"> • Preferred OIC Agents <ul style="list-style-type: none"> ○ Documented diagnosis of OIC and chronic pain in the past year AND ○ No history of GI or bowel obstruction AND ○ 1 claim for an opioid in the past 30 days • Non-Preferred OIC Agents <ul style="list-style-type: none"> ○ All preferred criteria met AND ○ Have tried 1 preferred OIC agents in the past 6 months OR ○ 1 claim with the requested agent in the past 105 days • Relistor Injection <ul style="list-style-type: none"> ○ Above OIC criteria OR ○ Documented diagnosis of OIC and active cancer in the past year AND ○ No history of GI or bowel obstruction AND ○ 1 claim for an opioid in the past 30 days |
| IRRITABLE BOWEL SYNDROME – DIARRHEA | | |
| <ul style="list-style-type: none"> • VIBERZI [New starts require clinical review] <ul style="list-style-type: none"> ○ Documented diagnosis of IBS – D in the past year and 1 claim for Viberzi in the past 105 days • LOTROXEX <ul style="list-style-type: none"> ○ 1 claim for LOTROXEX in the past 105 days OR ○ New starts require clinical review – MANUAL PA • XIFAXAN – (see Antibiotics, GI) | | |
| SHORT BOWEL SYNDROME AND SELECTED GI AGENTS ^{DUR+} | | |
| <p>HIV/AIDS Non-infectious Diarrhea</p> <ul style="list-style-type: none"> • MYTESI <ul style="list-style-type: none"> ○ Documented diagnosis of HIV/AIDS and non-infectious diarrhea in the past year AND ○ 1 claim for an antiretroviral in the past 30 days | <p>Short Bowel Syndrome (SBS)</p> <ul style="list-style-type: none"> • GATTEX <ul style="list-style-type: none"> ○ 1 claim for GATTEX in the past 105 days OR ○ New starts require clinical review | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|-------------------------------------|--|
| LEUKOTRIENE MODIFIERS ^{DUR+} | | |
| montelukast | ACCOLATE (zafirlukast) | Minimum Age Limit <ul style="list-style-type: none"> 12 years: ZYFLO & ZYFLO CR Non-Preferred Criteria <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months |
| zafirlukast | SINGULAIR (montelukast) | |
| | zileuton | |
| | ZYFLO (zileuton) | |
| LIPOTROPICS, OTHER (NON-STATINS) | | |
| ACL INHIBITORS AND COMBINATIONS | | Non-Preferred Criteria – Fibric Acid Derivatives <ul style="list-style-type: none"> Have tried 2 different preferred Fibric Acid Derivative agents in the past 6 months |
| | NEXLETOL (bempedoic acid) | |
| | NEXLIZET (bempedoic acid/ezetimibe) | |
| ANGIOPHOTIN-LIKE 3 INHIBITORS | | JUXTAPID – MANUAL PA |
| | EVKEEZA (evinacumab-dgnb) | |
| BILE ACID SEQUESTRANTS | | KYNAMRO • Requires clinical review |
| cholestyramine | colesevelam | |
| cholestyramine light | COLESTID (colestipol) | LEQVIO • Requires clinical review |
| colestipol tablet | colestipol packet | |
| | PREVALITE (cholestyramine) | |
| | QUESTRAN (cholestyramine) | NEXLETOL and NEXLIZET • Require clinical review |
| | QUESTRAN LIGHT (cholestyramine) | |
| | WELCHOL (colesevelam) | PRALUENT – MANUAL PA |
| CHOLESTEROL ABSORPTION INHIBITORS | | REPATHA – MANUAL PA |
| ezetimibe | ZETIA (ezetimibe) | |
| FIBRIC ACID DERIVATIVES | | WELCHOL • Documented diagnosis of Type 2 Diabetes AND • 30 days of therapy with an antidiabetic agent in the past 6 months OR • 90 days of therapy with WELCHOL in the past 105 days |
| fenofibrate | fenofibric acid | |
| gemfibrozil | FENOGLIDE (fenofibrate) | |
| | FIBRICOR (fenofibric acid) | |
| | LIPOFEN (fenofibrate) | |
| | LOPID (gemfibrozil) | |
| | TRICOR (fenofibrate) | |
| | TRILIPIX (fenofibric acid) | |
| MTP INHIBITOR | | |
| | JUXTAPID (lomitapide) | |
| NIACIN | | |
| niacin ER | | |
| OMEGA-3 FATTY ACIDS | | |
| omega-3 acid ethyl esters | icosapent ethyl | |
| | LOVAZA (omega-3 acid ethyl esters) | |
| PCSK-9 INHIBITORS | | |
| REPATHA (evolocumab) | LEQVIO (inclisiran) | |
| | PRALUENT (alirocumab) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|--|
| LIPOTROPICS, STATINS ^{DUR+} | | |
| STATINS | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 10 years: ATORVALIQ Suspension <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Have tried 2 different preferred statin or statin combination agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>Simvastatin</p> <ul style="list-style-type: none"> • Daily doses \geq 80 mg require clinical review |
| atorvastatin | ALTOPREV (lovastatin) | |
| lovastatin | ATORVALIQ (atorvastatin) | |
| pravastatin | CRESTOR (rosuvastatin) | |
| rosuvastatin | EZALLOR SPRINKLE (rosuvastatin) | |
| simvastatin | FLOLIPID (simvastatin) | |
| | fluvastatin | |
| | fluvastatin ER | |
| | LESCOL XL (fluvastatin) | |
| | LIPITOR (atorvastatin) | |
| | LIVALO (pitavastatin) | |
| | pitavastatin | |
| | ZOCOR (simvastatin) | |
| | ZYPITAMAG (pitavastatin) | |
| STATIN COMBINATIONS | | |
| ezetimibe/simvastatin | amlodipine/atorvastatin | |
| | CADUET (amlodipine/atorvastatin) | |
| | VYTORIN (ezetimibe/simvastatin) | |
| MISCELLANEOUS BRAND/GENERIC | | |
| ALLERGEN EXTRACT IMMUNOTHERAPY | | <p>CUMULATIVE quantity limit (per 31 days)</p> <ul style="list-style-type: none"> • 31 tablets: alprazolam ER <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 2 kits: epinephrine <p>EVRYSDI – MANUAL PA</p> <p>PALFORZIA – MANUAL PA</p> |
| | GRASTEK | |
| | ORALAIR | |
| | PALFORZIA | |
| | RAGWITEK | |
| EPINEPHRINE | | |
| epinephrine (Mylan) | AUVI-Q (epinephrine) | |
| | epinephrine (all other manufacturers) | |
| | EPIPEN (epinephrine) | |
| | EPIPEN JR (epinephrine) | |
| | NEFFY (epinephrine) ^{NR} | |
| MISCELLANEOUS | | |
| alprazolam | alprazolam ER | |
| hydroxyzine HCL | CAMZYOS (mavacamten) | |
| hydroxyzine pamoate | CRENESSITY (crinecerfont) ^{NR} | |
| megestrol | EVRYSDI (risdiplam) | |
| REVLIMID (lenalidomide) | KORLYM (mifepristone) | |
| | lenalidomide | |
| | VERQUVO (vericiguat) | |
| | VISTARIL (hydroxyzine pamoate) | |
| | XANAX, XANAX XR (alprazolam) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| MISCELLANEOUS BRAND/GENERIC (continued) | | |
| SUBLINGUAL NITROGLYCERIN | | |
| nitroglycerin | | |
| NITROLINGUAL (nitroglycerin) | | |
| NITROSTAT (nitroglycerin) | | |
| MOVEMENT DISORDER AGENTS ^{DUR+} | | |
| AUSTEDO (deutetrabenazine) | INGREZZA INITIATION PACK (valbenazine) | <p>AUSTEDO and AUSTEDO XR</p> <ul style="list-style-type: none"> • Documented diagnosis of Huntington's chorea OR • Documented diagnosis of tardive dyskinesia AND • 90 days of therapy with either agent in the past 105 days OR • New starts require clinical review – MANUAL PA <p>INGREZZA</p> <ul style="list-style-type: none"> • Documented diagnosis of Huntington's chorea OR • Documented diagnosis of tardive dyskinesia AND • 90 days of therapy with this agent in the past 105 days OR • New starts require clinical review – MANUAL PA |
| AUSTEOD XR (deutetrabenazine) | XENAZINE (tetrabenazine) | |
| INGREZZA (valbenazine) | | |
| INGREZZA SPRINKLE (valbenazine) | | |
| tetrabenazine | | |
| | | |
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| | | |
| MULTIPLE SCLEROSIS AGENTS ^{DUR+} | | |
| BETASERON (interferon beta-1b) | AMPYRA (dalfampridine) | <p>Preferred Agents</p> <ul style="list-style-type: none"> • Documented diagnosis of multiple sclerosis <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented diagnosis of multiple sclerosis AND • Have tried 2 different preferred agents in the past 6 months OR • 3 claims with the requested agent in the last 105 days <p>KESIMPTA, PONVORY, TASCENSO ODT, and ZEPOSIA</p> <ul style="list-style-type: none"> • Require clinical review <p>MAVENCLAD – MANUAL PA</p> <p>MAYZENT – MANUAL PA</p> <p>OCREVUS and OCREVUS ZUNOVO – MANUAL PA</p> |
| COPAXONE (glatiramer) 20 mg | AUBAGIO (teriflunomide) | |
| dalfampridine ER | AVONEX (interferon beta-1a) | |
| dimethyl fumarate | BAFIERTAM (monomethyl fumarate) | |
| fingolimod | BRIUMVI (ublituximab-xiyy) | |
| REBIF (interferon beta-1b) | COPAXONE (glatiramer) 40 mg | |
| REBIF REBIDOSE (interferon beta-1b) | GILENYA (fingolimod) | |
| teriflunomide | glatiramer | |
| TYSABRI (natalizumab) | GLATOPA (glatiramer) | |
| | KESIMPTA PEN (ofatumumab) | |
| | MAVENCLAD (cladribine) | |
| | MAYZENT (siponimod) | |
| | OCREVUS (ocrelizumab) | |
| | OCREVUS ZUNOVO (ocrelizumab/hyaluronidase-ocsq) ^{NR} | |
| | PLEGRIDY (peginterferon beta-1a) | |
| | PONVORY (ponesimod) | |
| | TASCENSO ODT (fingolimod) | |
| | TECFIDERA (dimethyl fumarate) | |
| | VUMERITY (diroximel fumarate) | |
| | ZEPOSIA (ozanimod) | |
| | | |
| | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|--|
| MUSCULAR DYSTROPHY AGENTS | | |
| EMFLAZA (deflazacort) | AGAMREE (vamorolone) | ELEVIDYS – MANUAL PA |
| | AMONDYS-45 (casimersen) | EMFLAZA – MANUAL PA |
| | deflazacort | EXONDYS – MANUAL PA |
| | DUVYZAT (givinostat) ^{NR} | VILTEPSO – MANUAL PA |
| | ELEVIDYS (delandistrogene moxeparvovec-rokl) | VYONDYS – MANUAL PA |
| | EXONDYS-51 (eteplirsen) | |
| | VILTEPSO (viltolarsen) | |
| | VYONDYS-53 (golodirsen) | |
| NSAIDS | | |
| COX II SELECTIVE | | <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 20 tablets: ketorolac tablets <p>ELYXYB</p> <ul style="list-style-type: none"> • Requires clinical review <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Non-Selective & Combinations <ul style="list-style-type: none"> ○ Have tried 2 different preferred non-selective or NSAID/GI protectant combination agents in the past 6 months • COX II Selective <ul style="list-style-type: none"> ○ Documented diagnosis of Osteoarthritis, Rheumatoid Arthritis, Familial Adenomatous Polyposis, or Ankylosing Spondylitis AND ○ 90 days of therapy with the requested agent in the past 105 days OR ○ Have tried 1 preferred COX-II Selective Agent and 1 preferred Non-Selective Agent OR ○ Documented diagnosis of GI Bleed, GERD, PUD, GI Perforation, or Coagulation Disorder AND ○ Have tried 1 preferred COX-II Selective agent |
| meloxicam | CELEBREX (celecoxib) | |
| | celecoxib | |
| | ELYXYB (celecoxib) | |
| NON-SELECTIVE | | |
| diclofenac sodium | DAYPRO (oxaprozin) | |
| diclofenac sodium ER | diclofenac potassium | |
| EC-naproxen DR 500 mg tablet | DOLOBID (diflunisal) ^{NR} | |
| etodolac tablet | etodolac capsule, etodolac ER | |
| flurbiprofen | FELDENE (piroxicam) | |
| ibuprofen | fenoprofen | |
| indomethacin capsule | indomethacin ER, indomethacin suppository | |
| ketoprofen | ketoprofen | |
| ketorolac | kiprofen | |
| nabumetone | LOFENA (diclofenac potassium) | |
| naproxen | meclufenamate | |
| piroxicam | mefenamic acid | |
| sulindac | NALFON (fenoprofen) | |
| | NAPRELAN (naproxen) | |
| | NAPROSYN (naproxen) | |
| | naproxen, naproxen CR, naproxen ER | |
| | oxaprozin | |
| | RELAFEN DS (nabumetone) | |
| | TOLECTIN 600 (tolmetin) | |
| | tolmetin | |
| NSAID/GI PROTECTANT COMBINATIONS | | |
| | ARTHROTEC 50, 75 (diclofenac/misoprostol) | |
| | diclofenac/misoprostol | |
| | ibuprofen/famotidine | |
| | naproxen/esomeprazole | |
| | VIMOVO (naproxen/esomeprazole) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| OPHTHALMIC AGENTS | | |
| ANTIBIOTICS | | |
| bacitracin/polymyxin | AZASITE (azithromycin) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 16 years: RESTASIS • 17 years: XIIDRA • 18 years: CEQUA, MIEBO, VEVYE <p>Quantity Limit (per 31 days)</p> <ul style="list-style-type: none"> • 2 mL: VEVYE • 3 mL: MIEBO • 5.5 mL: RESTASIS Multidose • 60 units: CEQUA, RESTASIS Droperette, XIIDRA <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Anti-Inflammatory Agents <ul style="list-style-type: none"> ◦ Have tried 2 different preferred agents in the past 6 months • Dry Eye Agents / CEQUA <ul style="list-style-type: none"> ◦ 4 claims for RESTASIS Droperette and XIIDRA in the past 6 months <p>EYSUVIS</p> <ul style="list-style-type: none"> • Requires clinical review <p>MIEBO</p> <ul style="list-style-type: none"> • Requires clinical review <p>RESTASIS Multidose</p> <ul style="list-style-type: none"> • Require clinical review <p>TYRVAYA</p> <ul style="list-style-type: none"> • Requires clinical review <p>VEVYE</p> <ul style="list-style-type: none"> • Requires clinical review |
| ciprofloxacin | bacitracin | |
| erythromycin | BESIVANCE (besifloxacin) | |
| gentamicin | CILOXAN (ciprofloxacin) | |
| moxifloxacin | gatifloxacin | |
| ofloxacin | NATACYN (natamycin0 | |
| polymyxin B/trimethoprim | neomycin/bacitracin/polymyxin | |
| tobramycin | OCUFLOX (ofloxacin) | |
| | sulfacetamide | |
| | TOBREX (tobramycin) | |
| | VIGAMOX (moxifloxacin) | |
| ANTIBIOTIC-STEROID COMBINATIONS | | |
| BLEPHAMIDE S.O.P. (sulfacetamide/prednisolone) | MAXITROL (neomycin/polymyxin/dexamethasone) | |
| neomycin/bacitracin/polymyxin/hydrocortisone | neomycin/polymyxin/gramicidin | |
| neomycin/polymyxin/dexamethasone | TOBRADEX ST (tobramycin/dexamethasone) | |
| PRED-G (gentamicin/prednisolone) | | |
| sulfacetamide/prednisolone | | |
| TOBRADEX (tobramycin/dexamethasone) | | |
| tobramycin/dexamethasone | | |
| ZYLET (tobramycin/loteprednol) | | |
| ANTI-INFLAMMATORY AGENTS | | |
| dexamethasone | ACULAR, ACULAR LS (ketorolac) | |
| diclofenac sodium | ACUVAIL (ketorolac) | |
| difluprednate | bromfenac | |
| FLAREX (fluorometholone) | BROMSITE (bromfenac) | |
| fluorometholone | DUREZOL (difluprednate) | |
| flurbiprofen | FML (fluorometholone) | |
| FML FORTE (fluorometholone) | ILEVRO (nepafenac) | |
| ketorolac | INVELTYS (loteprednol) | |
| MAXIDEX (dexamethasone) | LOTEMAX, LOTEMAX SM (loteprednol) | |
| PRED MILD (prednisolone) | loteprednol | |
| prednisolone acetate | NEVANAC (nepafenac) | |
| prednisolone sodium phosphate | PRED FORTE (prednisolone) | |
| | PROLENSA (bromfenac) | |
| DRY EYE AGENTS | | |
| RESTASIS Droperette (cyclosporine) | CEQUA (cyclosporine) | |
| XIIDRA (lifitegrast) | cyclosporine | |
| | EYSUVIS (loteprednol) | |
| | MIEBO (perfluorohexyloactane) | |
| | RESTASIS Multidose (cyclosporine) | |
| | TYRVAYA (varenicline) | |
| | VEVYE (cyclosporine) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|---|--|
| OPHTHALMIC, GLAUCOMA AGENTS | | |
| BETA BLOCKERS | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 18 years: IYUZEH <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days |
| BETIMOL (timolol) | betaxolol | |
| carteolol | BETOPTIC S (betaxolol) | |
| ISTALOL (timolol) | timolol droperette, daily drop, gel | |
| levobunolol | TIMOPTIC; TIMOPTIC OCUDOSE, XE (timolol) | |
| timolol drops 0.25%, 0.5% | | |
| CARBONIC ANHYDRASE INHIBITORS | | |
| dorzolamide | AZOPT (brinzolamide) | |
| | brinzolamide | |
| COMBINATION AGENTS | | |
| COMBIGAN (brimonidine/timolol) | brimonidine/timolol | |
| dorzolamide/timolol | COSOPT (dorzolamide/timolol) | |
| SIMBRINZA (brinzolamide/brimonidine) | dorzolamide/timolol PF | |
| PARASYMPATHOMIMETICS | | |
| pilocarpine | PHOSPHOLINE IODIDE (echothiophate iodide) | |
| PROSTAGLANDIN ANALOGS | | |
| latanoprost | bimatoprost | |
| | IYUZEH (latanoprost) | |
| | LUMIGAN (bimatoprost) | |
| | tafluprost | |
| | TRAVATAN Z (travoprost) | |
| | travoprost | |
| | VYZULTA (latanoprost) | |
| | XALATAN (latanoprost) | |
| | XELPROS (latanoprost) | |
| | ZIOPTAN (tafluprost) | |
| RHO KINASE INHIBITORS/COMBINATIONS | | |
| RHOPRESSA (netarsudil) | | |
| ROCKLATAN (netarsudil/latanoprost) | | |
| SYMPATHOMIMETICS | | |
| ALPHAGAN P (brimonidine) | brimonidine 0.1%, 0.15% | |
| brimonidine 0.2% | | |
| OPHTHALMICS FOR ALLERGIC CONJUNCTIVITIS | | |
| ALREX (loteprednol) | ALOCRIAL (nedocromil) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months <p>VERKAZIA</p> <ul style="list-style-type: none"> Requires clinical review |
| azelastine | ALOMIDE (lodoxamide) | |
| cromolyn | bepotastine | |
| ketotifen ^{OTC} | BEPREVE (bepotastine) | |
| olopatadine | epinastine | |
| ZADITOR (ketotifen) | LASTACAPT (alcaftadine) | |
| | VERKAZIA (cyclosporine) | |
| | ZERVIAE (cetirizine) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| OPIATE DEPENDENCE TREATMENTS | | |
| DEPENDENCE | | Buprenorphine/naloxone provider summary found here PROBUPHINE – MANUAL PA SUBLOCADE – MANUAL PA VIVITROL – MANUAL PA |
| buprenorphine/naloxone SL tablet | BRIXADI (buprenorphine) | |
| naltrexone | buprenorphine | |
| SUBOXONE (buprenorphine/naloxone) | buprenorphine/naloxone film | |
| | lofexidine ^{NR} | |
| | LUCEMYRA (lofexidine) | |
| | SUBLOCADE (buprenorphine) | |
| | VIVITROL (naltrexone) | |
| | ZUBSOLV (buprenorphine/naloxone) | |
| TREATMENT | | |
| KLOXXADO (naloxone) | LIFEMS NALOXONE (naloxone convenience kit) | |
| naloxone | | |
| NARCAN (naloxone) | | |
| OPVEE (nalmeffene) | | |
| REXTOVY (naloxone) | | |
| ZIMHI (naloxone) | | |
| OTIC ANTIBIOTICS | | |
| CIPRO HC (ciprofloxacin/hydrocortisone) | ciprofloxacin | Maximum Age Limit <ul style="list-style-type: none"> • 9 years: CIPRO HC Ciprofloxacin/Dexamethasone Suspension Criteria <ul style="list-style-type: none"> • Age ≥ 6 months AND • Experiencing otorrhea secondary to recent, post-tympanostomy tube placement AND • Continued otorrhea after 10 days of otic treatment with ciprofloxacin ophthalmic solution and dexamethasone ophthalmic suspension |
| CORTISPORIN-TC (neomycin/colistin/hydrocortisone) | ciprofloxacin/fluocinolone | |
| fluocinolone | ciprofloxacin/dexamethasone | |
| neomycin/polymyxin/hydrocortisone | DERMOTIC (fluocinolone) | |
| | FLAC OTIC OIL (fluocinolone) | |
| | hydrocortisone/acetic acid | |
| | OTOVEL (ciprofloxacin/fluocinolone) | |
| | | |
| | | |
| | | |
| PANCREATIC ENZYMES | | |
| CREON (lipase/protease/amylase) | PERTZYE (lipase/protease/amylase) | Non-Preferred Criteria <ul style="list-style-type: none"> • Have tried 2 different preferred agents in the past 6 months |
| ZENPEP (lipase/protease/amylase) | VIOKACE (lipase/protease/amylase) | |
| PARATHYROID AGENTS | | |
| calcitriol | doxercalciferol | |
| cinacalcet | RAYALDEE (calcifediol) | |
| ergocalciferol | ROCALTROL (calcitriol) | |
| paricalcitol | SENSIPAR (cinacalcet) | |
| ZEMPLAR (paricalcitol) | YORVIPATH (palopecteriparatide) ^{NR} | |
| | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|--|
| PHOSPHATE BINDERS | | |
| calcium acetate | AURYXIA (ferric citrate) | |
| CALPHRON (calcium acetate) | FOSRENOL (lanthanum) | |
| sevelamer carbonate tablet | lanthanum | |
| | MAGNEBIND (calcium carbonate/magnesium) | |
| | REVELA (sevelamer) | |
| | sevelamer carbonate packet, sevelamer HCl | |
| | VELPHORO (sucroferric oxyhydroxide) | |
| | XPHOZAH (tenapanor) | |
| PLATELET AGGREGATION INHIBITORS | | |
| aspirin/dipyridamole | EFFIENT (prasugrel) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Documented diagnosis AND • Have tried 2 different preferred agents in the past 6 months OR • 90 days of therapy with the requested agent in the past 105 days <p>ZONTIVITY – MANUAL PA</p> |
| BRILINTA (ticagrelor) | PLAVIX (clopidogrel) | |
| cilostazol | | |
| clopidogrel | | |
| dipyridamole | | |
| pentoxifylline | | |
| prasugrel | | |
| PLATELET STIMULATING AGENTS | | |
| NPLATE (romiplostim) | ALVAIZ (eltrombopag) | |
| PROMACTA (eltrombopag) tablet | DOPTELET (avatrombopag) | |
| | MULPLETA (lusutrombopag) | |
| | PROMACTA (eltrombopag) packet | |
| | TAVALISSE (fostatinib) | |
| POTASSIUM REMOVING AGENTS | | |
| LOKELMA (sodium zirconium cyclosilicate) | KIONEX (sodium polystyrene sulfonate) | |
| SPS (sodium polystyrene sulfonate) suspension | sodium polystyrene sulfonate | |
| | SPS (sodium polystyrene sulfonate) enema | |
| | VELTASSA (patiomer calcium sorbitex) | |
| PRENATAL VITAMINS | | |
| CLASSIC PRENATAL | <p>All prenatal vitamins are non-preferred except for those specifically indicated as preferred.</p> | <p>List of Preferred NDC's for Prenatal Vitamins can be found here</p> |
| COMPLETE NATAL DHA | | |
| COMPLETENATE | | |
| M-NATAL PLUS | | |
| NIVA-PLUS | | |
| PRENATAL PLUS VITAMIN-MINERAL | | |
| PNV 72, 95, 124, and 137 / IRON / FOLIC ACID | | |
| SE-NATAL-19 | | |
| STUART ONE | | |
| THRIVITE RX | | |
| TRICARE | | |
| TRINATAL RX 1 | | |
| WESNATAL DHA COMPLETE | | |
| WESTAB PLUS | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|---|
| PSEUDOBLBAR AFFECT AGENTS | | |
| | NUEDEXTA (dextromethorphan/quinidine) | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of pseudobulbar affect disorder OR 90 days of therapy with NUEDEXTA in the past 105 days |
| PULMONARY ANTIHYPERTENSIVE AGENTS | | |
| ACTIVIN SIGNALING INHIBITORS | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 18 years: ADEMPAS, OPSYNVI, TADLIQ <p>Maximum Age Limit</p> <ul style="list-style-type: none"> 12 years: REVATIO suspension <p>Preferred Criteria</p> <ul style="list-style-type: none"> PAH Agents <ul style="list-style-type: none"> Documented diagnosis of pulmonary hypertension Sildenafil tablets <ul style="list-style-type: none"> ≤ 1 year of age and documented diagnosis of pulmonary hypertension, patent ductus arteriosus, or persistent fetal circulation OR ≥ 1 year of age and documented diagnosis of pulmonary hypertension OR 90 days of therapy with the requested agent in the past 105 days <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of pulmonary hypertension AND Have tried 1 preferred PAH agent in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days <p>LIQREV, OPSUMIT, OPSYNVI, ORENITRAM ER, TYVASO, and VENTAVIS</p> <ul style="list-style-type: none"> Require clinical review <p style="text-align: center; background-color: yellow;">See below for additional PA Criteria/DUR+ Rules</p> |
| | WINREVAIR (sotatercept-csrk) | |
| COMBINATION AGENTS | | |
| | OPSYNVI (macitentan/tadalafil) | |
| ENDOTHELIN RECEPTOR ANTAGONISTS | | |
| ambrisentan | OPSUMIT (macitentan) | |
| bosentan | TRACLEER (bosentan) | |
| LETAIRIS (ambrisentan) | TRYVIO (aprocitentan) | |
| PDE5 INHIBITORS | | |
| sildenafil (generic REVATIO) tablet | ADCIRCA (tadalafil) | |
| tadalafil | ALYQ (tadalafil) | |
| | LIQREV (sildenafil) | |
| | REVATIO (sildenafil) | |
| | sildenafil (generic REVATIO) suspension | |
| | TADLIQ (tadalafil) | |
| PROSTACYCLINS | | |
| | ORENITRAM ER (treprostinil) | |
| | ORENITRAM TITRATION PAK (treprostinil) | |
| | TYVASO (treprostinil) | |
| | VENTAVIS (iloprost) | |
| SELECTIVE PROSTACYCLINE RECEPTOR AGONISTS | | |
| | UPTRAVI (selexipag) | |
| SOLUBLE GUANYLATE CYCLASE STIMULATORS | | |
| | ADEMPAS (riociguat) | |
| <p>ADEMPAS</p> <ul style="list-style-type: none"> Documented diagnosis of persistent/recurrent chronic thromboembolic pulmonary hypertension (WHO Group 4) or pulmonary arterial hypertension (WHO Group 1) AND Have tried 1 preferred PAH agent in the past 6 months OR 90 days of therapy with ADEMPAS in the past 105 days <p>REVATIO Suspension</p> <ul style="list-style-type: none"> ≤ 12 years of age AND Documented diagnosis of pulmonary hypertension, patent ductus arteriosus, or persistent fetal circulation, or a history of a heart transplant OR 90 days stable therapy with REVATIO Suspension in the past 105 days | | <p>TADLIQ</p> <ul style="list-style-type: none"> Documented diagnosis of pulmonary hypertension AND Have tried sildenafil (generic REVATIO) suspension in the past 6 months OR 90 days of therapy with TADLIQ in the past 105 days <p>UPTRAVI</p> <ul style="list-style-type: none"> Documented diagnosis of pulmonary hypertension AND Have tried 1 preferred endothelin receptor antagonist in the past 6 months AND Have tried 1 preferred PDE5 inhibitor in the past 6 months OR 90 days of therapy with UPTRAVI in the past 105 days |

MISSISSIPPI DIVISION OF MEDICAID UNIVERSAL PREFERRED DRUG LIST

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---------------------------|--|--|
| ROSACEA TREATMENTS | | |
| metronidazole | AVAR (sulfacetamide sodium/sulfur) | <p><u>Note:</u></p> <ul style="list-style-type: none"> Topical Sulfonamides used for Rosacea will require a manual PA for age > 21 years. Other labeled indications are limited to < 21 years. |
| | AVAR LS (sulfacetamide sodium/sulfur) | |
| | AVAR-E (sulfacetamide sodium/sulfur) | |
| | BP 10-1 (sulfacetamide sodium/sulfur) | |
| | brimonidine | |
| | EPSOLAY (benzoyl peroxide) | |
| | FINACEA (azelaic acid) | |
| | METROCREAM (metronidazole) | |
| | METROGEL (metronidazole) | |
| | MIRVASO (brimonidine) | |
| | NORITATE (metronidazole) | |
| | OVACE (sulfacetamide sodium) | |
| | OVACE PLUS (sulfacetamide sodium) | |
| | RHOFADE (oxymetazoline) | |
| | ROSADAN (metronidazole) | |
| | ROSULA (sulfacetamide sodium/sulfur) | |
| | sodium sulfacetamide | |
| | sodium sulfacetamide/sulfur | |
| | SOOLANTRA (ivermectin) | |
| | SUMADAN (sulfacetamide sodium/sulfur) | |
| | SUMADAN XLT (sulfacetamide sodium/sulfur/avob) | |
| | SUMAXIN (sulfacetamide sodium/sulfur) | |
| | SUMAXIN CP (sulfacetamide sodium/sulfur) | |
| | SUMAXIN TS (sulfacetamide sodium/sulfur) | |
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| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|-----------------------------------|--|
| SEDATIVE HYPNOTIC AGENTS | | |
| BENZODIAZEPINES DUR+ | | <p>MS DOM Opioid Initiative – Criteria details found here</p> <ul style="list-style-type: none"> Concomitant use of Opioids and Benzodiazepines <p>Maximum Age Limit</p> <ul style="list-style-type: none"> 64 years: zolpidem 7.5 mg, 10 mg, and 12.5 mg <p>Gender and Dose Limit</p> <ul style="list-style-type: none"> Female: AMBIEN 5 mg, AMBIEN CR 6.25 mg, INTERMEZZO 1.75 mg Male: all strengths of zolpidem <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months <p>HETLIOZ capsules</p> <ul style="list-style-type: none"> Documented diagnosis of circadian rhythm sleep disorder AND Documented diagnosis indicating total blindness OR Documented diagnosis of Smith-Magenis syndrome <p>HETLIOZ liquid</p> <ul style="list-style-type: none"> Age 3-15 years AND Documented diagnosis of Smith-Magenis syndrome <p>Note:</p> <ul style="list-style-type: none"> Single-source benzodiazepines and barbiturates are NOT covered. <ul style="list-style-type: none"> PA's will NOT be issued for these drugs. <p style="background-color: yellow; text-align: center;">See below for additional PA Criteria/DUR+ Rules</p> |
| estazolam | flurazepam | |
| temazepam 15 mg, 30 mg capsule | HALCION (triazolam) | |
| | quazepam | |
| | RESTORIL (temazepam) | |
| | temazepam 7.5 mg, 22.5 mg capsule | |
| | triazolam | |
| OTHERS DUR+ | | |
| eszopiclone | AMBIEN (zolpidem) | |
| ramelteon | AMBIEN CR (zolpidem) | |
| zaleplon | BELSOMRA (suvorexant) | |
| zolpidem tablet | DAYVIGO (lemborexant) | |
| | doxepin | |
| | EDULAR (zolpidem) | |
| | HETLIOZ LQ (tasimelteon) | |
| | LUNESTA (eszopiclone) | |
| | QUVIVIQ (daridorexant) | |
| | ROZEREM (ramelteon) | |
| | tasimelteon | |
| | zolpidem capsule | |
| | zolpidem sublingual tablet | |
| | zolpidem ER | |
| | | |
| | | |
| <p>CUMULATIVE Quantity Limit – Benzodiazepines</p> <ul style="list-style-type: none"> 31 units/31 days: Quantity limit per rolling days for all strengths. DUR+ will allow an early refill override for one dose or therapy change per year. <p>CUMULATIVE Quantity Limit – Triazolam</p> <ul style="list-style-type: none"> 10 units/31 days: Quantity limit per rolling days for all strengths. 60 units/365 days: Quantity limit per rolling days for all strengths. <p>CUMULATIVE Quantity Limit – Non-Benzodiazepines</p> <ul style="list-style-type: none"> 31 units/31 days: Quantity limit per rolling days for all strengths. DUR+ will allow an early refill override for one dose or therapy change per year. <p>CUMULATIVE Quantity Limit – HETLIOZ LQ</p> <ul style="list-style-type: none"> 1 bottle (48 mL or 158 mL): Quantity limit per rolling days for all strengths. DUR+ will allow an early refill override for one dose or therapy change per year. <p>CUMULATIVE Quantity Limit – ZOLPIMIST</p> <ul style="list-style-type: none"> 1 canister/31 days: male; Quantity limit per rolling days for all strengths. DUR+ will allow an early refill override for one dose or therapy change per year. 1 canister/62 days: female; Quantity limit per rolling days for all strengths. DUR+ will allow an early refill override for one dose or therapy change per year. | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|--|--|
| SELECT CONTRACEPTIVE PRODUCTS | | |
| INJECTABLE CONTRACEPTIVES | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> 1 claim with the requested agent in the past 105 days |
| medroxyprogesterone | DEPO-PROVERA (medroxyprogesterone) | |
| INTRAVAGINAL CONTRACEPTIVES | | |
| ENILLORING (etonogestrel/ethinyl estradiol) | PHEXXI (lactic acid/citric acid/potassium bitartrate) | |
| ORAL CONTRACEPTIVES ^{DUR+} | | |
| All contraceptives are preferred except for those specifically indicated as non-preferred. | AMETHIA (levonorgestrel/ethinyl estradiol) | |
| | AMETHYST (levonorgestrel/ethinyl estradiol) | |
| | BALCOLTRA (levonorgestrel/ethinyl estradiol) | |
| | BEYAZ (drospirenone/ethinyl estradiol/levomefolate) | |
| | CAMRESE (levonorgestrel/ethinyl estradiol) | |
| | CAMRESE LO (levonorgestrel/ethinyl estradiol) | |
| | FEMLYV (norethindrone acetate/ethinyl estradiol) ^{NR} | |
| | JOLESSA (levonorgestrel/ethinyl estradiol) | |
| | LO LOESTRIN FE (norethindrone/ethinyl estradiol/iron) | |
| | LOESTRIN (norethindrone/ethinyl estradiol) | |
| | LOESTRIN FE (norethindrone/ethinyl estradiol/iron) | |
| | MINZOYA (levonorgestrel/ethinyl estradiol/iron) | |
| | NATAZIA (estradiol valerate/dienogest) | |
| | NEXTSTELLIS (drospirenone/estetrol) | |
| | OCELLA (ethinyl estradiol/drospirenone) | |
| | SAFYRAL (drospirenone/ethinyl estradiol/levomefolate) | |
| SIMPESSE (levonorgestrel/ethinyl estradiol) | | |
| TAYTULLA (norethindrone/ethinyl estradiol/iron) | | |
| TYDEMY (drospirenone/ethinyl estradiol/levomefolate) | | |
| YASMIN (ethinyl estradiol/drospirenone) | | |
| YAZ (ethinyl estradiol/drospirenone) | | |
| TRANSDERMAL CONTRACEPTIVES | | |
| XULANE (norelgestromin/ethinyl estradiol) | norelgestromin/ethinyl estradiol | |
| | TWIRLA (levonorgestrel/ethinyl estradiol) | |
| | ZAFEMY (norelgestromin/ethinyl estradiol) | |
| SICKLE CELL AGENTS | | |
| DROXIA (hydroxyurea) | ADAKVEO (crizanlizumab-tmca) | <p>ENDARI – <u>MANUAL PA</u></p> |
| hydroxyurea | CASGEVY (exagamglogene autotemcel) | |
| | ENDARI (glutamine) | |
| | HYDREA (hydroxyurea) | |
| | l-glutamine | |
| | LYFGENIA (lovetibeglogene autotemcel) | |
| | SIKLOS (hydroxyurea) | |
| | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|------------------------------------|---|--|
| SKELETAL MUSCLE RELAXANTS | | |
| DUR+ | | |
| baclofen 5 mg, 10 mg, 20 mg tablet | AMRIX (cyclobenzaprine) | <p>Quantity Limit</p> <ul style="list-style-type: none"> 84 tablets/180 days: carisoprodol <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of an approvable indication AND Have tried 2 different preferred agents in the past 6 months <p>Baclofen granules, solution, and suspension</p> <ul style="list-style-type: none"> Require clinical review <p>Carisoprodol</p> <ul style="list-style-type: none"> Documented diagnosis of acute musculoskeletal condition AND No history with meprobamate in the past 90 days AND 1 claim for cyclobenzaprine in the past 21 <p>Carisoprodol with codeine</p> <ul style="list-style-type: none"> Requires clinical review <p>TANLOR</p> <ul style="list-style-type: none"> Requires clinical review |
| chlorzoxazone | baclofen 15 mg tablet | |
| cyclobenzaprine 5 mg, 10 mg tablet | baclofen suspension | |
| methocarbamol | carisoprodol | |
| tizanidine tablet | carisoprodol/aspirin | |
| | cyclobenzaprine 7.5 mg tablet | |
| | cyclobenzaprine ER | |
| | DANTRIUM (dantrolene) | |
| | dantrolene | |
| | FEXMID (cyclobenzaprine) | |
| | FLEQSUVY (baclofen) | |
| | LORZONE (chlorzoxazone) | |
| | LYVISPAH (baclofen) | |
| | metaxalone | |
| | NORGESIC (orphenadrine/aspirin/caffeine) | |
| | NORGESIC FORTE (orphenadrine/aspirin/caffeine) | |
| | orphenadrine | |
| | orphenadrine/aspirin/caffeine | |
| | ORPHENGESIC FORTE (orphenadrine/aspirin/caffeine) | |
| | SOMA (carisoprodol) | |
| | TANLOR (methocarbamol) | |
| | tizanidine capsule | |
| | ZANAFLEX (tizanidine) | |
| SMOKING DETERRENTS | | |
| NICOTINE TYPE | | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> 18 years: CHANTIX <p>Quantity Limit</p> <ul style="list-style-type: none"> 336 tablets/year: CHANTIX 0.5 mg tabs, 1 mg tabs, and continuing pack 2 treatment courses/year: CHANTIX Starter Pack |
| nicotine gum ^{OTC} | NICOTROL INHALER CARTRIDGE | |
| nicotine lozenge ^{OTC} | NICOTROL NASAL SPRAY | |
| nicotine patch ^{OTC} | | |
| NON-NICOTINE TYPE | | |
| bupropion SR | | |
| CHANTIX (varenicline) | | |
| varenicline | | |
| | | |
| | | |
| | | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|--|-------------------------------------|---|
| STERIODS (TOPICAL) | | |
| LOW POTENCY | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> • Low Potency <ul style="list-style-type: none"> ○ Have tried 2 different preferred low potency agents in the past 6 months • Medium Potency <ul style="list-style-type: none"> ○ Have tried 2 different preferred medium potency agents in the past 6 months • High Potency <ul style="list-style-type: none"> ○ Have tried 2 different preferred high potency agents in the past 6 months • Very High Potency <ul style="list-style-type: none"> ○ Have tried 2 different preferred very high potency agents in the past 6 months |
| alclometasone | fluocinolone | |
| DERMA-SMOOTH-FS (fluocinolone) | hydrocortisone lotion | |
| desonide | HYDROXYM (hydrocortisone) | |
| hydrocortisone cream, ointment, solution | PROCTOCORT (hydrocortisone) | |
| MEDIUM POTENCY | | |
| fluticasone | BESER (fluticasone) | |
| mometasone | CAPEX (fluocinolone) | |
| PANDEL (hydrocortisone probutate) | clocortolone | |
| prednicarbate cream | CLODERM (clocortolone) | |
| | flurandrenolide | |
| | fluticasone lotion | |
| | LOCOID (hydrocortisone butyrate) | |
| | prednicarbate ointment | |
| | SYNALAR (fluocinolone) | |
| HIGH POTENCY | | |
| betamethasone dipropionate cream, lotion | amcinonide | |
| betamethasone dipropionate augmented | betamethasone dipropionate ointment | |
| betamethasone valerate | desoximetasone | |
| fluocinolone | diflorasone | |
| fluocinonide | halcinonide | |
| fluocinonide-E | HALOG (halcinonide) | |
| triamcinolone cream, ointment, lotion | KENALOG (triamcinolone) | |
| | TOPICORT (desoximetasone) | |
| | triamcinolone spray | |
| | VANOS (fluocinonide) | |
| VERY HIGH POTENCY | | |
| clobetasol cream, foam, gel, ointment, shampoo, solution | APEXICON E (diflorasone) | |
| clobetasol-E | BRYHALI (halobetasol) | |
| halobetasol | clobetasol emulsion | |
| | CLOBEX (clobetasol) | |
| | CLODAN (clobetasol) | |
| | DIPROLENE (betamethasone) | |
| | halobetasol | |
| | IMPEKLO (clobetasol) | |
| | LEXETTE (halobetasol) | |
| | OLUX (clobetasol) | |
| | TEMOVATE (clobetasol) | |
| | TOVET (clobetasol) | |
| | ULTRAVATE (halobetasol) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|---|--|
| STIMULANTS AND RELATED AGENTS | | |
| SHORT-ACTING | | DUR+ |
| dexmethylphenidate | ADDERALL (dextroamphetamine/amphetamine) | <p>Minimum Age Limit</p> <ul style="list-style-type: none"> • 3 years: ADDERALL, EVEKEO, PROCENTRA, ZENZEDI • 6 years: ADDERALL XR, ADHANSIA XR, ADZENYS ER SUSPENSION, ADZENYS XR ODT, APTENSIO XR, atomoxetine, AZSTARYS, clonidine ER, CONCERTA ER, COTEMPLA XR ODT, DAYTRANA, DESOXYN, DEXEDRINE, DYANAVEL XR, EVEKEO ODT, FOCALIN, FOCALIN XR, JORNAY PM, METADATE CD, METHYLIN, ONYDA XR, QELBREE, QUILLICHEW, QUILLIVANT XR, RELEXII ER, RITALIN LA, VYVANSE, XELSTRYM • 7 years: XYREM • 13 years: MYDAYIS • 16 years: modafinil • 18 years: armodafinil, SUNOSI, WAKIX <p>Maximum Age Limit</p> <ul style="list-style-type: none"> • 18 years: clonidine ER, COTEMPLA XR ODT, DAYTRANA, EVEKEO ODT, guanfacine ER <p>Quantity Limit – Stimulants (per 31 days)</p> <ul style="list-style-type: none"> • 31 tablets: ADDERALL XR, ADHANSIA XR, ADZENYS XR ODT, APTENSIO XR, AZSTARYS, CONCERTA ER 18, 27, & 54 mg, COTEMPLA XR-ODT 8.6 mg, DAYTRANA, DEXEDRINE Spansule, DYANAVEL XR Tablet, FOCALIN XR, JORNAY PM, METADATE CD, METHYLIN ER, MYDAYIS 37.5 mg & 50 mg, QUILLICHEW, RELEXII ER, RITALIN LA & SR, VYVANSE, XELSTRYM • 62 tablets: ADDERALL, CONCERTA ER 36 mg, COTEMPLA XR-ODT 17.3 & 25.9 mg, DESOXYN, EVEKEO, FOCALIN, METHYLIN, ZENZEDI • 248 mL: DYANAVEL XR Suspension • 310 mL: METHYLIN, PROCENTRA • 372 mL: QUILLIVANT XR <p>Quantity Limit – Narcolepsy (per 31 days)</p> <ul style="list-style-type: none"> • 31 tablets: armodafinil 150, 200 & 250 mg, modafinil 200 mg, SUNOSI • 46.5 tablets: modafinil 100 mg • 62 tablets: armodafinil 50 mg, WAKIX <p>Quantity Limit – Non-Stimulants (per 31 days)</p> <ul style="list-style-type: none"> • 31 tablets: atomoxetine, guanfacine ER, QELBREE 100 mg • 62 tablets: QELBREE 150 mg and 200 mg • 124 tablets: clonidine ER • 1 bottle (30 mL or 60 mL): ONYDA XR Suspension <p style="background-color: yellow;">See next page for additional PA Criteria/DUR+ Rules</p> |
| dextroamphetamine | amphetamine | |
| dextroamphetamine/amphetamine | EVEKEO (amphetamine) | |
| methylphenidate | EVEKEO ODT (amphetamine) | |
| PROCENTRA (dextroamphetamine) | FOCALIN (dexmethylphenidate) | |
| | methamphetamine | |
| | METHYLN (methylphenidate) | |
| | methylphenidate | |
| | RITALIN (methylphenidate) | |
| | ZENZEDI (dextroamphetamine) | |
| LONG-ACTING | | |
| ADDERALL XR (dextroamphetamine/amphetamine) | ADZENYS XR ODT (amphetamine) | |
| CONCERTA (methylphenidate) | APTENSIO XR (methylphenidate) | |
| dexmethylphenidate ER | AZSTARYS (serdexmethylphenidate/dexmethylphenidate) | |
| dextroamphetamine ER | COTEMPLA XR ODT (methylphenidate) | |
| dextroamphetamine/amphetamine ER | DAYTRANA (methylphenidate) | |
| DYANAVEL XR (amphetamine) suspension | DEXEDRINE (dextroamphetamine) | |
| lisdexamfetamine | dextroamphetamine/amphetamine ER | |
| methylphenidate CD | DYANAVEL XR (amphetamine) tablets | |
| methylphenidate ER tablet | FOCALIN XR (dexmethylphenidate) | |
| methylphenidate LA | JORNAY PM (methylphenidate) | |
| QUILLICHEW ER (methylphenidate) | methylphenidate patch | |
| QUILLIVANT XR (methylphenidate) | methylphenidate ER capsule | |
| VYVANSE (lisdexamfetamine) capsules | MYDAYIS (dextroamphetamine/amphetamine) | |
| | RELEXII (methylphenidate) | |
| | RITALIN LA (methylphenidate) | |
| | VYVANSE (lisdexamfetamine) chewable tablets | |
| | XELSTRYM (dextroamphetamine) | |
| NARCOLEPSY | | |
| armodafinil | NUVIGIL (armodafinil) | |
| modafinil | PROVIGIL (modafinil) | |
| SUNOSI (solriamfetol) | sodium oxybate | |
| XYREM (sodium oxybate) | WAKIX (pitolisant) | |
| | XYWAV (calcium/magnesium/potassium/sodium oxybate) | |
| NON-STIMULANTS | | |
| atomoxetine | INTUNIV (guanfacine) | |
| clonidine ER | NEXICLON XR (clonidine) | |
| guanfacine ER | ONYDA XR (clonidine) ^{NR} | |
| QELBREE (viloxazine) | STRATTERA (atomoxetine) | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|-------------|
| STIMULANTS AND RELATED AGENTS ^{DUR+} <i>(continued)</i> | | |
| See previous page for additional PA Criteria/DUR+ Rules | | |
| <p>Non-Preferred Short Acting Criteria</p> <p>ADD/ADHD</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD AND • Have tried 2 different preferred Short Acting agents in the past 6 months OR • 1 claim for a 30-day supply with the requested agent in the past 105 days <p>Narcolepsy: ADDERALL, EVEKEO, METHYLIN, PROCENTRA, RITALIN, ZENZEDI</p> <ul style="list-style-type: none"> • Documented diagnosis of narcolepsy AND • 30 days of therapy with preferred modafinil or armodafinil in the past 6 months AND • 1 preferred agent indicated for narcolepsy in the past 6 months OR • Have tried 1 claim for a 30-day supply with the requested agent in the past 105 days | <p>Non-Preferred Long Acting Criteria</p> <p>ADD/ADHD</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD AND • Have tried 2 different preferred Long-Acting agents in the past 6 months OR • 1 claim for a 30-day supply with the requested agent in the past 105 days <p>Narcolepsy: ADDERALL XR, APTENSIO XR, CONCERTA ER, DEXEDRINE, METADATE CD, METHYLIN ER, MYDAYIS, NUVIGIL, PROVIGIL, QUILLICHEW, QUILLIVANT XR, RITALIN LA</p> <ul style="list-style-type: none"> • Documented diagnosis of narcolepsy AND • 30 days of therapy with preferred modafinil or armodafinil in the past 6 months AND • 1 different preferred agent indicated for narcolepsy in the past 6 months OR • 1 claim for a 30-day supply with the requested agent in the past 105 days | |
| <p>Armodafinil</p> <ul style="list-style-type: none"> • Documented diagnosis of narcolepsy, obstructive sleep apnea, shift work sleep disorder, or bipolar depression <p>Atomoxetine</p> <ul style="list-style-type: none"> • Age ≥ 21 years AND • Documented diagnosis of ADD/ADHD <p>Clonidine ER</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD <p>Guanfacine ER</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD <p>JORNAY PM</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD AND • 84 days of therapy with 2 different preferred LA methylphenidate agents in the past 12 months AND • 84 days of therapy with 1 preferred non-methylphenidate LA stimulant agent in the past 12 months OR • Documented diagnosis of ADD/ADHD AND • 84 days of therapy with JORNAY PM in the past 105 days <p>Modafinil</p> <ul style="list-style-type: none"> • Documented diagnosis of narcolepsy, obstructive sleep apnea, shift work sleep disorder, depression, sleep deprivation or Steinert Myotonic Dystrophy Syndrome | <p>ONYDA XR</p> <ul style="list-style-type: none"> • Requires clinical review <p>QELBREE</p> <ul style="list-style-type: none"> • Documented diagnosis of ADD/ADHD AND • 30 days of therapy with a preferred ADHD agent in the past 105 days OR • 30 days of therapy with QELBREE in the past 105 days <p>SUNOSI</p> <ul style="list-style-type: none"> • Documented diagnosis of narcolepsy or obstructive sleep apnea AND • 30 days of therapy with preferred modafinil or armodafinil in the past 6 months <p>VYVANSE</p> <ul style="list-style-type: none"> • Documented diagnosis of binge eating disorder or ADD/ADHD <p>WAKIX</p> <ul style="list-style-type: none"> • Requires clinical review <p>XYREM</p> <p>Documented diagnosis of narcolepsy or excessive daytime sleepiness OR 30 days of therapy with this agent in the past 105 days</p> <p>XYWAV</p> <ul style="list-style-type: none"> • Requires clinical review | |

| PREFERRED AGENTS | NON-PREFERRED AGENTS | PA CRITERIA |
|---|--|---|
| TETRACYCLINES ^{DUR+} | | |
| doxycycline hyclate | demeclocycline | <p>Non-Preferred Agents</p> <ul style="list-style-type: none"> Have tried 2 different preferred agents in the past 6 months <p>Demeclocycline</p> <ul style="list-style-type: none"> Documented diagnosis of Syndrome of Inappropriate Antidiuretic Hormone Secretion (SIADH) will allow for automatic approval <p>ORACEA</p> <ul style="list-style-type: none"> Requires clinical review |
| doxycycline monohydrate capsule | DORYX (doxycycline hyclate) | |
| minocycline capsule | DORYX MPC (doxycycline hyclate) | |
| tetracycline capsule | doxycycline hyclate DR | |
| | doxycycline IR/DR | |
| | doxycycline monohydrate suspension, tablet | |
| | LYMEPAK (doxycycline hyclate) | |
| | MINOCIN (minocycline) | |
| | minocycline tablet | |
| | minocycline ER | |
| | MINOLIRA ER (minocycline) | |
| | MORGIDOX (doxycycline hyclate) | |
| | NUZYRA (omadacycline) | |
| | ORACEA (doxycycline monohydrate) | |
| | SOLODYN (minocycline) | |
| | tetracycline tablet | |
| ULCERATIVE COLITIS & CROHN'S AGENTS ^{DUR+} *See Cytokine & CAM Antagonists Class for Additional Agents* | | |
| ORAL | | <p>Non-Preferred Criteria</p> <ul style="list-style-type: none"> Documented diagnosis of Ulcerative Colitis AND Have tried 2 different preferred agents in the past 6 months OR 90 days of therapy with the requested agent in the past 105 days <p>VELSIPITY</p> <ul style="list-style-type: none"> Requires clinical review |
| APRISO (mesalamine) | AZULFIDINE (sulfasalazine) | |
| balsalazide | COLAZAL (balsalazide) | |
| budesonide | DELZICOL (mesalamine) | |
| PENTASA (mesalamine) | DIPENTUM (olsalazine) | |
| sulfasalazine | LIALDA (mesalamine) | |
| sulfasalazine DR | mesalamine | |
| UCERIS (budesonide) | mesalamine DR | |
| | mesalamine ER | |
| | VELSIPITY (etrasimod) | |
| RECTAL | | |
| mesalamine suppository | budesonide | |
| | CANASA (mesalamine) | |
| | mesalamine enema | |
| | ROWASA (mesalamine) | |
| | SFROWASA (mesalamine) | |
| | UCERIS (budesonide) | |
| UREA CYCLE DISORDER AGENTS | | |
| CARBAGLU (carglumic acid) | BUPHENYL (sodium phenylbutyrate) | |
| carglumic acid | OLPRUVA (sodium phenylbutyrate) | |
| | PHEBURANE (sodium phenylbutyrate) | |
| | RAVICTI (glycerol phenylbutyrate) | |