

WYOMING MEDICAID Preferred Drug List (PDL) - AUGUST 1, 2012

Drug classes not included on this list are not managed through a Preferred Drug List (PDL).
HOWEVER, THIS EXCLUSION IS NOT A GUARANTEE OF PAYMENT OR COVERAGE. Dosage limits and other requirements may apply.

Drugs new to market are non-preferred until a clinical review has been completed. PA criteria will apply to both the pediatric population,
as well as the adult population for those plans where PA/PDL limits are allowed.

Unless otherwise noted on the PDL, generic substitution is mandatory.

Yellow highlighted items below indicate new changes to the PDL. Red font indicates quantity/dosage limits apply. *Indicates BRAND is Preferred. May Use DAW 5.
Contact the GHS PA Helpdesk @ 877-207-1126 for prior authorization if client has primary insurance that will not cover the brand name medication.

**Please refer to the Additional Therapeutic Criteria Chart, Dosage Limitation List (red font indicates quantity/dosage limits apply),
Eprocates, and the Wyoming Medicaid Provider Manual at <http://wyequalitycare.org> for additional criteria.**

| THERAPEUTIC CLASS | PREFERRED AGENTS | PREFERRED AGENTS REQUIRING CLINICAL CRITERIA | CLINICAL CRITERIA | NON-PREFERRED AGENTS GENERIC MANDATORY POLICY APPLIES <small>THIS LIST IS NOT ALL INCLUSIVE PLEASE CONTACT GHS FOR QUESTIONS</small> |
|--|--|--|---|---|
| ALLERGY / ASTHMA | ANTIHISTAMINES, MINIMALLY SEDATING | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | desloratadine |
| | cetirizine fexofenadine loratadine | | | CLARINEX RDT/SYRUP levocetirizine |
| | ANTIHISTAMINE/DECONGESTANT COMBINATIONS | | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. |
| | cetirizine/pseudoephedrine fexofenadine/pseudoephedrine loratadine/pseudoephedrine | | | |
| | ANTICHOLINERGIC BRONCHODILATORS | | Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for Spiriva 5 day STARTER package will be allowed one (1) time per recipient. | ATROVENT HFA |
| | ipratropium SPIRIVA | | | |
| | CORTICOSTEROID / BRONCHODILATOR COMBO'S | | Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for Advair 7 and 14-day STARTER package will be allowed one (1) time per recipient. | |
| | ADVAIR/HFA DULERA SYMBICORT | | | |
| | LEUKOTRIENE MODIFIERS | | Trial and failure of preferred agent greater than or equal to 30 days in the last 12 months will be | zafirlukast ZYFLO |
| | SINGULAIR TABS, CHEWABLES, GRANULES | | | |
| | NASAL ANTIHISTAMINES | | Trial and failure of preferred agent greater than or equal to 90 days in the last 12 months will be required before approval can be given for a non-preferred agent. | ASTEPRO 0.15% DYMISTA (use separate agents) PATANASE |
| | azelastine | | | |
| | NASAL STEROIDS | | Trial and failure of two (2) preferred agents greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Rhinocort will be approved for pregnancy. | BECONASE AQ DYMISTA (use separate agents) flunisolide OMNARIS QNASL RHINOCORT triamcinolone (BRAND IS PREFERRED) VERAMYST ZETONNA |
| | fluticasone NASACORT AQ* NASONEX | | | |
| SHORT ACTING BRONCHODILATORS - INHALERS | | Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. | ALUPENT | |
| PROAIR HFA PROVENTIL HFA VENTOLIN HFA XOPENEX HFA | | | | |
| SHORT ACTING BRONCHODILATORS - NEBULIZERS | | Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. | ACCUNEB levalbuterol (BRAND IS PREFERRED) METAPROTERENOL | |
| albuterol neb XOPENEX neb* | | | | |
| STEROID INHALANTS | | Trial and failure of three (3) preferred agents greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Alvesco will be approved for a history of oral thrush with steroid inhalants. | AEROBID/AEROBID-M ALVESCO ASMANEX STARTER PACK (use preferred) AZMACORT PULMICORT | |
| ASMANEX budesonide FLOVENT HFA/DISK QVAR | | | | |
| ALZHEIMERS | ALZHEIMER AGENTS | | Client must have a diagnosis of dementia. | ARICEPT 23MG (use preferred) donepezil ODT (use preferred) |
| | donepezil EXELON PATCH/SOLUTION galantamine/ER NAMENDA rivastigmine capsules | | | |

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| ANALGESICS | BUPRENORPHINE COMBINATIONS | | Client must have a diagnosis of opioid dependence or abuse. This is not to be used to for the treatment of chronic pain. Only one (1) narcotic prescription will be allowed between fills. Subutex will be approved for clients pregnant or nursing or with a documented allergy to naloxone. Dosage limits apply (Max Dose: 24mg/day). | SUBUTEX |
| | LONG-ACTING | | Trial and failure of a preferred agent(s) greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. Fentanyl patches are limited to one patch every 72 hours. C-IIIs and C-IVs are not included and are available without prior authorization with the exception of Butrans (generic substitution is mandatory). **Butrans requires a trial of morphine sulfate or low dose trial of fentanyl patch. ***Embeda and Oxecta require a diagnosis of drug/substance abuse and a 14 day trial of a preferred agent. | AVINZA BUTRANS** EMBEDA*** KADIAN (10mg/200mg) morphines sulfate ER capsules NUCYNTA ER OPANA ER (5mg/10mg/20mg/30mg/40mg) OXECTA*** oxymorphone ER (7.5mg/15mg) OXYCONTIN/CR |
| | fentanyl patch morphine sulfate ER <u>tablets</u> | SUBOXONE/FILM | | |
| | SHORT-ACTING C-IIs | | Trial and failure of three (3) preferred agents greater than or equal to a 6 day supply in the last 90 days will be required before approval can be given for a non-preferred agent. | EXALGO levorphanol NUCYNTA oxymorphone oxycodone/IBU |
| | codeine sulfate hydromorphone morphine sulfate oxycodone oxycodone/APAP oxycodone/ASA | | | |
| | TRAMADOL PRODUCTS | | Trial and failure of a preferred agent(s) greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. Quantity and dosage limits apply(max 8 tabs/day). | CONZIP RYBIX ODT tramadol/apap tramadol ER |
| | tramadol | | | |
| ANDROGENS | TESTOSTERONE TOPICAL GELS | | Testosterone agents are only allowed for diagnosis of hypogonadism or insufficient testosterone production. Prior authorization required for non-preferred agent. | TESTIM GEL |
| | | ANDROGEL | | |
| ANTIBIOTICS | QUINOLONES | | | AVELOX FACTIVE NOROXIN PROQUIN |
| | ciprofloxacin/ER levofloxacin ofloxacin | | | |
| | DOXYCYCLINE | | | ADOXA (use preferred) DORYX (use preferred) ORACEA (use preferred) |
| | doxycycline | | | |
| | MINOCYCLINE | | | SOLODYN (use preferred) |
| | minocycline/ER | | | |
| ANTICOAGULANTS | LOW MOLECULAR WEIGHT HEPARIN (LMWH) | | | enoxaparin (BRAND IS PREFERRED) FRAGMIN (use preferred) LOVENOX 300MG/3ML (use preferred) |
| | LOVENOX* | | | |
| | DIRECT THROMBIN INHIBITOR | | Client must have diagnosis of non-valvular atrial fibrillation and relative contraindication to warfarin for approval. | |
| | | PRADAXA | | |
| | SELECTIVE FACTOR XA INHIBITOR | | Client must have diagnosis of non-valvular atrial fibrillation or prophylaxis of deep vein thrombosis which can lead to pulmonary embolism in clients undergoing hip or knee replacement. | |
| | | XARELTO | | |
| ANTICONVULSANTS | DIAZEPAM RECTAL GEL | | | diazepam gel (BRAND IS PREFERRED) |
| | DIASTAT* | | | |

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| ANTIDEPRESSANTS | ANTIDEPRESSANTS | | <p>Trial and failure of two (2) preferred agents greater than or equal to six (6) weeks will be required before approval can be given for a non-preferred agent.</p> <p>Trazodone, buspirone, fluvoxamine, MAO inhibitors, TCA's, bupropion IR and venlafaxine IR do not require prior authorization but will not count towards meeting preferred therapy requirements.</p> <p>*Cymbalta will be approved for a diagnosis of peripheral neuropathy and osteoarthritis of the knee.</p> <p>**Lexapro will be approved for adolescents between the ages of 12 - 17.</p> | <p><i>fluoxetine tablets (use preferred)</i> <i>mirtazapine 7.5mg and mirtazapine rapid-dissolve tablets (use preferred)</i> <i>venlafaxine ER tablets (use preferred)</i></p> <p>APLENZIN CYMBALTA* LEXAPRO** (use brand) PRISTIQ VIIBRYD</p> |
| ANTIHYPERTENSIVES | ACE INHIBITORS | | <p>Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> | |
| | ACE INHIBITORS AND DIURETICS | | <p>Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> | |
| | ANGIOTENSIN RECEPTOR BLOCKERS (ARBs) | | <p>Trial and failure of an ACE inhibitor greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for preferred ARB. Non-preferred ARBs and ARB/diuretic combinations also require a history of ALL preferred ARBs before approval can be given.</p> | <p>ATACAND EDARBI eprosartan 600mg irbesartan (BRAND IS PREFERRED) TEVETEN 400mg</p> |
| | ARBs AND DIURETICS | | | <p>ATACAND HCTZ EDARBYCLOR irbesartan HCTZ (BRAND IS PREFERRED) TEVETEN HCTZ</p> |
| | ARB COMBINATIONS | | | <p>AZOR TWINSTA (use separate agents) TRIBENZOR (use separate agents)</p> |
| | ALPHA-BLOCKERS | | | <p>clonidine patch (BRAND IS PREFERRED) NEXICLON XR (use preferred)</p> |
| | <p>CATAPRES PATCHES* clonidine</p> | | | |

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| ANTIPSYCHOTICS | ATYPICAL ANTIPSYCHOTICS ABILIFY/ODT GEODON* INVEGA INVEGA SUSTENNA olanzapine quetiapine RISPERDAL CONSTA risperidone ZYPREXA RELPREVV | | <p>**Quetiapine doses less than 100mg will require prior authorization <u>without</u> a diagnosis of mood disorder or major depressive disorder. For titration doses, contact the GHS Pharmacy Help Desk for an override.</p> <p>Non-preferred agents (Fanapt, Latuda, and Saphris) require a trial of ALL preferred agents at max doses.</p> <p><i>Typical antipsychotics do <u>not</u> required prior authorization.</i></p> <p>Dosage limits apply: ABILIFY <13 years of age: 23mg/day ABILIFY ≥13 years of age: 45mg/day GEODON ≤ 17 years of age: 180mg/day GEODON > 17 years of age: 300mg/day INVEGA all ages: 18mg/day Risperidone ≤ 17 years of age: 5mg/day Risperidone > 17 years of age: 24mg/day Olanzapine < 13 years of age: 15mg/day Olanzapine > 13 years of age: 30mg/day Quetiapine <13 years of age: 600mg/day Quetiapine 13-17 years of age: 900mg/day Quetiapine > 17 years of age: 1200mg/day</p> <p>***Latuda will be approved for female clients of child-bearing age.</p> | <p>FANAPT LATUDA*** SAPHRIS SEROQUEL XR (<i>use preferred</i>) ziprasidone (<i>BRAND IS PREFERRED</i>)</p> |
| | SPECIAL ATYPICAL ANTIPSYCHOTICS | | <p>Dosage limits apply: 1350mg/day</p> | |
| ANTIVIRALS, ORAL | HERPES AGENTS acyclovir famciclovir VALTREX* | | | valacyclovir (<i>BRAND IS PREFERRED</i>) |
| CHOLESTEROL | BILE ACID SEQUESTRANT cholestyramine/light colestipol | | Trial and failure of ALL preferred agents greater than or equal to six (6) months in the last 12 months will be required before approval can be given for a non-preferred agent. | WELCHOL |
| | INTESTINAL CHOLESTEROL ABSORPTION INHIBITOR ZETIA | | | |
| | STATINS, LOW POTENCY lovastatin pravastatin | | <p>Trial and failure of a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> <p>If client's current medication therapy is contraindicated with the preferred statin(s) due to a drug-drug interaction, a non-preferred agent may be obtained with a prior authorization.</p> | <p>ALTOPREV fluvastatin/ER</p> |
| | STATINS, HIGH POTENCY atorvastatin simvastatin | | <p>Trial and failure of a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> <p>If client's current medication therapy is contraindicated with the preferred statin(s) due to a drug-drug interaction, a non-preferred agent may be obtained with a prior authorization.</p> | <p>CRESTOR LIVALO</p> |
| | STATIN COMBINATIONS CADUET* | | Trial and failure of a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | <p>ADVICOR (<i>use separate agents</i>) amlodopine/atorvastatin (<i>BRAND IS PREFERRED</i>) CHOLESTIN PRAVIGARD SIMCOR VYTORIN (<i>use separate agents</i>)</p> |
| | TRIGLYCERIDE LOWERING AGENTS fenofibrate gemfibrozil TRICOR | | Trial and failure of a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | <p>ANTARA fenofibric FENOGLIDE LOVAZA TRILIPIX</p> |

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|------------------------|---|--|---|---|
| CONTRACEPTIVES | <p align="center">ORAL CONTRACEPTIVES</p> altavera apri aviane azurette balzia BREVICON* briellyn cryselle emoquette enpresse errin ESTROSTEP FE* gildess FE jolessa jolivette junel/junel FE kariva kelnor lessina levora LOESTRIN 24 FE LOSEASONIQUE low-ogestrel lutera LYBREL microgestin mononessa NECON 10/11-28 nora-be norgestrel/ethinyl estradiol NORINYL 1/50-28 OGESTREL orsythia ORTHO TRI-CYCLEN LO* ORTHO-NOVUM 1/35-28, 7/7/7-28* OVCON 50 portia previfem reclipen seasonale SEASONIQUE* solia sprintec sronyx trinessa TRI-NORINYL* tri-previfem trivora velivet YASMIN* YAZ* zenchent ZOVIA | | | amethia (BRAND IS PREFERRED) amethyst (BRAND IS PREFERRED) aranelle (BRAND IS PREFERRED) BEYAZ (PA required) camila (use preferred) camrese (BRAND IS PREFERRED) caziant (use preferred) cesia (use preferred) cyclafem (BRAND IS PREFERRED) FEMCON FE (PA required) GENERESS FE CHW (PA required) gianvi (BRAND IS PREFERRED) heather (use preferred) introvale (use preferred) leena (BRAND IS PREFERRED) LO LOESTRIN (PA required) loryna (BRAND IS PREFERRED) NATAZIA (PA required) neon 0.5/35, 1/35, 7/7/7 (BRAND IS PREFERRED) NECON 1/50 (use preferred) norethindrone/ethinyl estradiol chew (PA required) norethindrone (use preferred) NORINYL 1/35 (use preferred) nortrel (BRAND IS PREFERRED) ocella (BRAND IS PREFERRED) ORTHO-NOVUM 1/50 (use preferred) quasense (use preferred) SAFYRAL (PA required) syeda (BRAND IS PREFERRED) tilia FE (BRAND IS PREFERRED) tri-legest FE (BRAND IS PREFERRED) tri-lo-sprintec (BRAND IS PREFERRED) zarah (BRAND IS PREFERRED) zenchent FE chewable (PA required) zeosa chewable (PA required) |
| CORTICOSTEROIDS | <p align="center">ORAL CORTICOSTEROIDS</p> budesonide cortisone acetate dexamethasone/intensol hydrocortisone methylprednisone prednisolone prednisone | | | CELESTONE (use preferred) |
| COUGH AND COLD | <p align="center">COMBINATION AGENTS</p> BROMFED DM <p align="center">DEXTROMETHORPHAN AGENTS</p> DELSYM* <p align="center">GUAIFENESIN AGENTS</p> guaifenesin MUCINEX TAB 1200MG MUCINEX TAB 600MG ER MUCINEX CGH LIQ 5-100MG MUCINEX/KIDS GRA 100MG | | Refer to the Over the Counter Drug Coverage at www.wyequalitycare.org for a list of covered products. | Dextromethorphan Polistirex (BRAND IS PREFERRED) |

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| DIABETES | DIABETES AGENTS | | | |
| | BIGUANIDES | | | |
| | metformin/ER | | | FORTAMET (<i>use preferred</i>) GLUMETZA (<i>use preferred</i>) RIOMET (<i>use preferred</i>) |
| | α -GLUCOSIDASE INHIBITORS | | | |
| | acarbose | | Trial and failure of metformin and a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | GLYSET |
| | MEGLITINIDES | | | |
| | STARLIX* | | Trial and failure of metformin and a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | nateglinide (<i>BRAND IS PREFERRED</i>) PRANDIN |
| | THIAZOLIDINEDIONES | | | |
| | ACTOS | | Trial and failure of metformin and a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | ACTOSPLUS MET (<i>use separate agents</i>) AVANDIA AVANDAMET (<i>use separate agents</i>) |
| | SULFONYLUREAS | | | |
| | glimepiride/ER glipizide/ER glyburide/ER | | Trial and failure of metformin and a preferred agent greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | |
| | DIPEPTIDYL PEPTIDASE 4 (DPP-4) INHIBITORS | | | |
| | | JANUVIA ONGLYZA TRADJENTA | Trial and failure of metformin greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a preferred agent. | |
| | DIPEPTIDYL PEPTIDASE 4 (DPP-4) INHIBITOR COMBO AGENTS | | | |
| | | JANUMET JUVISYNC KOMBIGLYZE | Trial and failure of metformin greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a preferred agent . A 90 day trial of failure of the preferred agent is required before approval can be give for a non-preferred agent. | JANUMET XR JENTADUETO |
| | INCRETIN MIMETICS (GLP-1 RECEPTOR AGONISTS) | | | |
| | | BYETTA | Trial and failure of metformin greater than or equal to a 90 day supply in the last 12 months will be required before approval can be given for a preferred agent . A 90 day trial of failure of the preferred agent is required before approval can be give for a non-preferred agent. | BYDUREON VICTOZA |
| | INTERMEDIATE-ACTING INSULIN | | | |
| | HUMULIN N HUMULIN 70/30 NOVOLOG N NOVOLOG 70/30 | | | |
| | LONG-ACTING INSULIN | | | |
| LANTUS <u>via</u> | | | LANTUS OPTICLIK/SOLOSTAR (<i>use preferred</i>) LEVEMIR (<i>use preferred</i>) | |
| RAPID-ACTING INSULIN | | | | |
| APIDRA HUMALOG NOVOLOG | | | | |
| DIABETIC METERS/TEST STRIPS | | | | |
| FREESTYLE INSULINX FREESTYLE LITE FREESTYLE FREEDOM LITE ONE TOUCH ULTRA ONE TOUCH ULTRA 2 ONE TOUCH ULTRA MINI ONE TOUCH ULTRASMART PRECISION XTRA | | Quantity limit applies (1 meter/365days). | ALL OTHER METERS AND TEST STRIPS | |
| EAR | ANTIBIOTIC/STEROID COMBINATION | | | |
| | CORTISPORIN SOL 1% OTIC* Neomycin/Polymyxin B Sulfates/Hydrocortisone <u>suspension</u> ofloxacin | | | CETRAXAL (<i>use preferred</i>) CIPRODEX (<i>use preferred</i>) CIPRO HC (<i>use preferred</i>) COLY-MYCIN S (<i>use preferred</i>) CORTISPORIN-TC (<i>use preferred</i>) dexamethasone sodium phosphate (<i>use preferred</i>) FLUOCINOLONE ACET OIL 0.01% (<i>use preferred</i>) Neomycin/Polymyxin B Sulfates/Hydrocortisone <u>solution</u> (<i>BRAND IS PREFERRED</i>) |

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| FIBROMYALGIA | FIBROMYALGIA STEP 1 | | | |
| | amitriptyline cyclobenzaprine | | | |
| | FIBROMYALGIA STEP 2 | | Trial and failure of a Step 1 agent greater than or equal to six (6) weeks in the last 12 months is required for approval of a Step 2 agent. | |
| | | SAVELLA | | |
| GASTROINTESTINAL | DIGESTIVE ENZYMES | | Prior authorization required. | PANCREAZE pancrelipase <i>(BRAND IS PREFERRED)</i> PERTZYE TRI-PASE |
| | CREON 3000, 6000, 12000, 24000 UNIT ZENPEP* | | | |
| | PROTON PUMP INHIBITORS | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | ACIPHEX lansoprazole NEXIUM omeprazole <u>tablets</u> <i>(use preferred)</i> omeprazole bicarbonate VIMOVO <i>(use separate agents)</i> |
| | DEXILANT omeprazole <u>capsules</u> pantoprazole | | Lansoprazole solutabs will be approved for children less than or equal to 8 years of age. Lansoprazole capsules will be approved for children less than 1 year of age. | |
| GASTROINTESTINAL | MESALAMINE | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | APRISO ASACOL/HD CANASA LIALDA PENTASA 500MG <i>(use preferred)</i> ROWASA |
| | mesalamine enema PENTASA 250MG ONLY | | | |
| | GROWTH HORMONE | | PA is required for use outside of FDA-approved indications. Evaluation by an endocrinologist is preferred. Clinical evidence of improved growth will be required on a yearly basis to support ongoing utilization. Clinical evidence of need for growth hormone will be required for adult growth hormone deficiency and pediatric growth failure due to inadequate endogenous growth hormone. Trial and failure of two (2) preferred agents within the last 12 months will be required for the following indications: Pediatric: Growth failure due to inadequate endogenous growth hormone, Prader-Willi syndrome, children born small for gestation. Turner syndrome. Adult: Replacement for those with growth hormone deficiency. | HUMATROPE OMNITROPE SAIZEN SEROSTIM TEV-TROPIN ZORBTIVE |
| GROWTH HORMONE | GROWTH HORMONE | | PA is required for use outside of FDA-approved indications. Evaluation by an endocrinologist is preferred. | HUMATROPE OMNITROPE SAIZEN SEROSTIM TEV-TROPIN ZORBTIVE |
| | GENOTROPIN NORDITROPIN NUTROPIN AQ | | Clinical evidence of improved growth will be required on a yearly basis to support ongoing utilization. Clinical evidence of need for growth hormone will be required for adult growth hormone deficiency and pediatric growth failure due to inadequate endogenous growth hormone. Trial and failure of two (2) preferred agents within the last 12 months will be required for the following indications: Pediatric: Growth failure due to inadequate endogenous growth hormone, Prader-Willi syndrome, children born small for gestation. Turner syndrome. Adult: Replacement for those with growth hormone deficiency. | |
| | | | | |
| HEPATITIS C | INTERFERON | | Trial and failure of preferred agent greater than or equal to 30 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | PEG-INTRON |
| | PEGASYS | | Peg-Intron will be approved for pediatric patients (aged 18 and under), for retreatment, and for dosage adjustments that cannot be achieved with Pegasys. | |
| | PROTEASE INHIBITOR | | Prior authorization required for non-preferred agent. | INCIVEK |
| HEPATITIS C | VICTRELIS | | | |

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|---------------------------|---|--|--|--|
| IMMUNOMODULATORS | IMMUNOMODULATORS | | <p>Client must have diagnosis prior to approval for preferred agents (outlined below):</p> <p>Enbrel: Ankylosing Spondylitis (AS), Juvenile Idiopathic Arthritis (JIA), Plaque Psoriasis (PP), Psoriatic Arthritis (PA), Rheumatoid Arthritis (RA)**</p> <p>Humira: AS, Crohn's, JIA, PP, PA, RA**</p> <p>**60-day trial and failure of methotrexate required prior to approval of Enbrel or Humira for diagnosis of Rheumatoid Arthritis (RA)</p> <p>For non-preferred agents, 60-day trial and failure of a preferred agent is required and client must have diagnosis prior to approval (outlined below):</p> <p>Actemra: RA (60-day trial of methotrexate is required)</p> <p>Amevive: PP</p> <p>Cimzia: Crohn's***, RA</p> <p>Kineret: RA</p> <p>Orencia: JIA, RA</p> <p>Remicade: AS, Crohn's, PP, PA, RA, Ulcerative Colitis****</p> <p>Rituxan: RA</p> <p>Simponi: AS, PA, RA</p> <p>Stelara: PP</p> <p>Tysabri: Crohn's (additional PA criteria applies)</p> <p>***Cimzia will be allowed without a preferred trial for diagnosis of Crohn's</p> <p>****Remicade will be allowed without a preferred trial for diagnosis of Ulcerative Colitis</p> | <p>ACTEMRA AMEVIVE CIMZIA KINERET ORENCIA RAPTIVA REMICADE RITUXAN SIMPONI STELARA TYSABRI (additional criteria applies)</p> |
| INSOMNIA | <p>zaleplon zolpidem</p> | NON-BENZODIAZEPINES | <p>Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> <p>Rozerem is non-preferred without a history of substance abuse.</p> <p>Dosage limits apply: zaleplon: 30mg/day zolpidem: 15mg/day</p> | <p>EDLUAR (additional criteria applies) INTERMEZZO (additional criteria applies) ROZEREM zolpidem ER ZOLPIMIST (additional criteria applies)</p> |
| MIGRAINE | <p>naratriptan sumatriptan</p> | <p>TRIPATAN STEP 1</p> <p>TRIPATAN STEP 2</p> <p>MAXALT MLT</p> | <p>Quantity limits apply:</p> <p>naratriptan 1mg: 25tabs/34days naratriptan 2.5mg: 10tabs/34days sumatriptan kit: 3kits/34days sumatriptan vials: 2vials/34days sumatriptan nasal: 6bottles/34days sumatriptan 25mg: 41tabs/34days sumatriptan 50mg: 20tabs/34days sumatriptan 100mg: 10tabs/34days</p> <p>Trial and failure of a Step 1 agent greater will be required for approval of a Step 2 agent. Trial and failure of a Step 1 agent and a Step 2 agent will be required for approval of a non-preferred agent.</p> <p>Quantity limits apply:</p> <p>MAXALT MLT 5mg: 27tabs/34days MAXALT MLT 10mg: 14tabs/34days</p> | <p>AXERT FROVA MAXALT RELPAK TREXIMET ZOMIG</p> |
| MULTIPLE SCLEROSIS | <p>IMMUNOMODULATOR (GLATIRAMER INJECTION)</p> <p>COPAXONE</p> <p>INTERFERON BETA-1A</p> <p>AVONEX REBIF</p> <p>INTERFERON BETA-1B</p> <p>BETASERON</p> | | <p>Trial and failure of one (1) interferon agent AND failure of Copaxone before approval can be give for a non-preferre agent.</p> <p>For Tysabri, in addition to the above criteria, additional prior authorization criteria applies.</p> | <p>EXTAVIA GILENYA TYSABRI (additional criteria applies)</p> |

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|---------------------|--|---|---|---|
| NSAIDS | <p align="center">NSAIDs</p> diclofenac etodolac fenoprofen flurbiprofen ibuprofen indomethacin ketoprofen ketorolac meclofenamate meloxicam nabumetone naproxen oxaprozin sulindac tolmetin | | Trial and failure of two (2) preferred agents each greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. Dosing limits apply for ketorolac (limit 5days/34 days). | CALDOLOR CAMBIA POWDER CELEBREX FLECTOR <i>(additional criteria applies)</i> mefenamic acid NAPRELAN NEOPROFEN PENNSAID <i>(additional criteria applies)</i> SOLARAZE <i>(additional criteria applies)</i> SPRIX <i>(additional criteria applies)</i> VOLTAREN <i>(additional criteria applies)</i> ZIPSOR |
| OPHTHALMICS | <p align="center">OP. -ANTIBIOTICS- QUINOLONES</p> ciprofloxacin ofloxacin VIGAMOX <p align="center">OP. -ANTI-INFLAMMATORY- NSAIDS</p> flurbiprofen diclofenac ketorolac <p align="center">OP. -BETA-BLOCKERS</p> betaxolol carteolol levobunolol metipranolol timolol <p align="center">OP. -CARBONIC ANHYDRASE INHIBITOR</p> dorzolamide <p align="center">OP. -CARBONIC ANHYDRASE INHIBITOR COMBO</p> dorzolamide/timolol <p align="center">OP. -ANTIHISTAMINE/MAST CELL STABILIZERS STEP 1</p> azelastine cromolyn <p align="center">OP. -ANTIHISTAMINE/MAST CELL STABILIZERS STEP 2</p> PATADAY PATANOL <p align="center">OP. -PROSTAGLANDINS</p> latanoprost TRAVATAN Z <p align="center">OP. -SYMPATHOMIMETICS</p> ALPHAGAN P brimonidine dipivefrin | <p align="center">OP. -BETA-BLOCKERS</p> betaxolol carteolol levobunolol metipranolol timolol <p align="center">OP. -ANTIHISTAMINE/MAST CELL STABILIZERS STEP 2</p> PATADAY PATANOL | Trial and failure of a preferred agent greater than or equal to 5 days in the last 12 months will be required before approval can be given for a non-preferred agent. Azasite will be approved for pregnancy. Trial and failure of ALL preferred agents each greater than or equal to 5 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. Trial and failure of three (3) preferred agents each greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Betoptic S will be approved for those with heart and lung conditions. Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Trial and failure of a Step 1 agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a Step 2 agent. Trial and failure of a Step 1 agent and a Step 2 agent greater than or equal to 30 days in the last 12 months will be required for approval of a non-preferred agent. Emadine, Alomide, and Alocriil will be approved for pregnancy. Alomide will be approved for children under the Trial and failure of ALL preferred agents each greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. Trial of a preferred agent greater than or equal to 30 days in the last 12 months will be required before approval can be given for a non-preferred agent. | AZASITE BESIVANCE IQUIX levofloxacin MOXEZA ZYMAR ZYMAXID ACULAR/LS/PF <i>(use preferred)</i> ACUVAIL BROMDAY bromfenac NEVANAC BETIMOL BETOPTIC S ISTALOL AZOPT ALAMAST ALOCRIL ALOMIDE ALREX BEPREVE ELESTAT EMADINE ketotifen LASTACAPT LUMIGAN ZIOPTAN COMBIGAN <i>(use separate agents)</i> |
| OSTEOPOROSIS | <p align="center">BISPHOSPHONATES</p> alendronate <p align="center">NASAL CALCITONIN</p> calcitonin-salmon fortical | | Trial and failure of a preferred agent greater than or equal to 12 months will be required before approval can be given for a non-preferred agent. Fosamax liquid will be approved for clients that have difficulty swallowing. | ACTONEL ATELVIA FOSAMAX-D ibandronate |

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|------------------------------------|-------------------------------------|--|---|--|
| OVERACTIVE BLADDER | OVERACTIVE BLADDER AGENTS | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. Oxytrol will be approved for clients that have an inability to swallow. | DETROL LA ENABLEX GELNIQUE GEL 10% OXYTROL DIS SANCTURA XR tolterodine trospium |
| PHOSPHATE BINDERS | PHOSPHATE BINDERS | | Prior authorization required for non-preferred agents. | FOSRENOL RENVELA |
| PRENATAL VITAMINS | PRENATAL VITAMINS | | Prenatal vitamins containing Omega-3 and DHA will be approved for clients at high risk for pre-term labor. | ALL OTHER PRENATAL VITAMINS INCLUDING OVER-THE-COUNTER FORMULATIONS |
| PROSTATE | 5-ALPHA-REDUCTASE INHIBITORS | | Trial and failure of a preferred agent greater than or equal to a 30 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | AVODART JALYN (<i>use separate agents</i>) |
| | ALPHA BLOCKERS | | Trial and failure of a preferred agent greater than or equal to a 30 day supply in the last 12 months will be required before approval can be given for a non-preferred agent. | alfuzosin JALYN (<i>use separate agents</i>) RAPAFLO |
| PULMONARY ANTIHYPERTENSIVES | 5-ALPHA-REDUCTASE INHIBITORS | | Client must have a diagnosis of pulmonary hypertension. | |
| | ADCIRCA REVATIO | | Prior authorization required. Client must have a diagnosis of pulmonary hypertension with documented right-heart catheterization validating the diagnosis. | TRACLEER |
| RESTLESS LEG SYNDROME | RESTLESS LEG SYNDROME | | Client must have a diagnosis of Restless Leg Syndrome (RLS). Trial and failure of gabapentin greater than or equal to 60 days and a trial and failure of a dopamine agonist greater than or equal to 60 days in the last 12 months will be required before approval can be given for a non-preferred agent. | HORIZANT |
| | | gabapentin | | |

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|----------------------------------|---|--|---|--|
| SKELETAL MUSCLE RELAXANTS | <p align="center">MUSCLE RELAXANTS</p> baclofen cyclobenzaprine tizanidine <u>tablets</u> | | Trial and failure of a preferred agent greater than or equal to a 14 day supply in the last 12 months, along with a medical diagnosis of muscle spasticity will be required before approval can be given for a non-preferred agent. | <p>carisoprodol chlorzoxazone cyclobenzaprine ER metaxalone methocarbamol orphenadrine tizanidine <u>capsules</u> (<i>use preferred</i>)</p> <p>Carisoprodol is limited to 84 tabs/365 days.</p> |
| STIMULANTS | <p align="center">AMPHETAMINES</p> <p align="center"><small>LONG ACTING AMPHETAMINES</small></p> <p align="center">ADDERALL XR * VYVANSE dextroamphetamine CR</p> <p align="center"><small>IMMEDIATE RELEASE AMPHETAMINES</small></p> <p align="center">amphetamine salts combo dextroamphetamine</p> <p align="center">METHYLPHENIDATES</p> <p align="center"><small>LONG ACTING METHYLPHENIDATES</small></p> <p align="center">DAYTRANA FOCALIN XR methylin ER methylphenidate ER/CR/SA/SR <u>tablets</u></p> <p align="center"><small>IMMEDIATE RELEASE METHYLPHENIDATES</small></p> <p align="center">FOCALIN methylin <u>tablets</u> methylphenidate</p> <p align="center">SELECTIVE ALPHA-ADRENERGIC AGONIST</p> <p align="center"><small>GUANFACINE AGENTS</small></p> guanfacine | | <p>Clients must have a diagnosis for ADD, ADHD, narcolepsy, obstructive sleep apnea, shift work sleep disturbance, MS fatigue (see MS Fatigue criteria below), or refractory depression (see refractory depression criteria below).</p> <p>Diagnosis of MS fatigue will require a fatigue severity scale score of 5.0, a 60-day trial of amantadine <u>and</u> discontinuation of medications that may contribute to drowsiness and fatigue.</p> <p>Diagnosis of refractory depression will require a 6-week trial and failure of an antidepressant (monotherapy) and continued concomitant use of an antidepressant with the stimulant.</p> <p>Prior Authorization will be required for clients under the age of 5.</p> <p>Claims will require Prior Authorization if clients have a history of the following: glaucoma, cardiac arrhythmias, arteriosclerosis, untreated hypertension, untreated hyperthyroidism, substance abuse, or current MAO inhibitor use.</p> <p>Trial and failure of two (2) preferred agents (each from a different class: methylphenidate and amphetamine) greater than or equal to a 30 day supply in the last 12 months will be required before approval can be given for a non-preferred agent.</p> <p><u>Strattera is limited to 1 tablet/day; unless the dose is greater than 40mg/day or unable to achieve a prescribed dose with 1 tablet.</u></p> <p><small>Quantity limits apply: ADDERALL XR: 60mg/day amphetamine salts combo: 60mg/day amphetamine salts combo (narcolepsy): 90mg/day CONCERTA: 135mg/day DAYTRANA: 45mg/9 hour patch dextroamphetamine: 90mg/day dextroamphetamine CR: 90mg/day dexmethylphenidate: 30mg/day FOCALIN XR ≤ 13 years of age: 45mg/day FOCALIN XR > 13 years of age: 60mg/day methylin/methylphenidate: 135mg/day methylin/methylphenidate ER/CR/SR: 135mg/day VYVANSE: 105mg/day</small></p> | <p>AMPHETAMINES: amphetamine salts combo ER (BRAND IS PREFERRED)</p> <p>METHYLPHENIDATES: methylphenidate ER/CR/SR <u>capsules</u> (METADATE CD/RITALIN LA)</p> |
| | guanfacine | | <p>To obtain the non-preferred agent, client must meet the following criteria: Client must have a diagnosis of ADHD or ADD. Prior authorization will be required for clients under the age of 5. Client must have a trial and failure of a stimulant greater than or equal to a 14 OR a trial and failure of Strattera greater than or equal to a 30 day supply AND trial and benefit of guanfacine (Tenex) in the previous 12 months OR a contraindication to ADHD medications (including stimulant and non-stimulant) OR a TIC disorder associated with stimulants (trial of stimulant required).</p> | INTUNIV |

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| STIMULANTS <i>Continued</i> | CLONIDINE AGENTS | | Trial and benefit of clonidine IR will be required prior to approval of the non-preferred agent. | KAPVAY |
| | clonidine | | | |
| TOPICAL AGENTS | IMPETIGO ANTIBIOTICS | | Trial and failure of ALL preferred agents greater than or equal to 7 days in the past 90 days. | ALTABAX |
| | gentamicin mupirocin | | Use smallest size appropriate for 7 day trial. | |
| | BENZOYL PEROXIDE/CLINDAMYCIN COMBOS | | | ACANYA |
| | | benzoyl peroxide/clindamycin | Clients must be 12 to 20 years of age and have a diagnosis of acne vulgaris. Requires prior authorization for clients less than 12 years of age. Acne combinations are limited to clients under the age of 21. | |
| | CORTICOSTEROIS | | | PANDEL |
| | <small>C=CREAM; G=GEL; L=LOTION; O=OINTMENT</small> | | | |
| | <small>LOW POTENCY</small> | | | |
| | alclometasone desonide fluocinolone 0.01% hydrocortisone butyrate 0.1% (C) hydrocortisone 1%, 2.5% (C,L,O) prednicarbate | | Trial and failure of ALL preferred agents greater than or equal to 14 days in the last 90 days. | |
| | <small>MEDIUM POTENCY</small> | | | |
| | betamethasone valerate desoximetasone 0.05% (C) fluocinolone 0.025% fluticasone 0.05% (C) hydrocortisone butyrate 0.1% (O) hydrocortisone probutate 0.1% (C) mometasone triamcinolone 0.025%, 0.1% | | Trial and failure of ALL preferred agents greater than or equal to 14 days in the last 90 days. | CLODERM CORDRAN/SP TOPICORT LP TRIANEX |
| | <small>HIGH POTENCY</small> | | | |
| | amcinonide betamethasone dipropionate clobetasol desoximetasone 0.25%, 0.05% (G) diflorasone fluocinonide flurandrenolide fluticasone 0.005% (O) halobetasol triamcinolone 0.5% | | Trial and failure of ALL preferred agents greater than or equal to 14 days in the last 90 days. | HALOG |
| | IMMUNOMODULATORS | | | |
| | | ELIDEL PROTOPIC | Trial and failure of a preferred medium potency topical corticosteroid greater than or equal to a 21 day trial <u>and</u> a trial and failure of a preferred high potency topical corticosteroid greater than or equal to a 21 day trial in the last 90 days. | |
| | SALICYLIC ACID | | | All other topical salicylic acid formulations. |
| | aliclen shampoo 6% salacyn cream/lotion 6% Salicylic Acid Shampoo 6% | | | |
| | SCABICIDES/PEDICULICIDES | | | |
| | permethrin LINDANE | | Trial and failure of a preferred agent in the last 12 months. | NATROBA OVIDE ULESFIA |
| | UREA | | | All other topical urea formulations. |
| | Kerafoam Aerosol 30% Remeven Cream 50% urea hydration aerosol 35% urea emulsion 50% urea nail suspension 40% urea suspension 50% X-Viate Cream 40% | | | |