

**WYOMING MEDICAID**

**Preferred Drug List - Effective 9/30/09**

Listed drugs are preferred. Drugs in the PDL classes that are not listed are non-preferred and require a PA. 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.

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THERAPEUTIC CLASS	PREFERRED AGENTS	PREFERRED AGENTS REQUIRING CLINICAL CRITERIA	CLINICAL CRITERIA
<b>ALLERGY / ASTHMA THERAPIES</b>	<b>ANTIHISTAMINES, MINIMALLY SEDATING</b>		
	cetirizine fexofenadine loratadine		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.
	<b>ANTIHISTAMINE/DECONGESTANT COMBINATIONS</b>		
	cetirizine/pseudoephedrine loratadine/pseudoephedrine		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.
	<b>ANTICHOLINERGIC BRONCHODILATORS</b>		
	ATROVENT HFA ipratropium SPIRIVA		Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months.
	<b>ANTICHOLINERGIC BETA-AGONIST COMBO'S</b>		
	albuterol/ipratropium COMBIVENT		Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months.
	<b>CORTICOSTEROID / BRONCHODILATOR COMBO'S</b>		
	ADVAIR ADVAIR HFA SYMBICORT		Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months.
	<b>NASAL STEROIDS</b>		
	fluticasone NASACORT AQ VERAMYST		Trial and failure of two (2) preferred agents greater than or equal to 30 days in the last 12 months. Rhinocort will be approved for pregnancy.
	<b>LEUKOTRIENE MODIFIERS</b>		
	SINGULAIR		Trial and failure of preferred agent greater than or equal to 30 days in the last 12 months.
	<b>LONG ACTING BRONCHODILATORS</b>		
SEREVENT		Trial and failure of preferred agent greater than or equal to 30 days in the last 12 months.	
<b>SHORT ACTING BRONCHODILATORS - INHALERS</b>			
MAXAIR PROAIR HFA VENTOLIN HFA		Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months.	
<b>SHORT ACTING BRONCHODILATORS - NEBULIZERS</b>			
albuterol neb		Trial and failure of a preferred agent greater than or equal to 30 days in the last 12 months.	
<b>INHALED CORTICOSTEROIDS</b>			
ASMANEX AZMACORT budesonide FLOVENT HFA FLOVENTDISK PULMICORT		Trial and failure of three (3) preferred agents greater than or equal to 30 days in the last 12 months. Alvesco will be approved for a history of oral thrush.	
<b>ANTIBIOTICS</b>	<b>BETA-LACTAMS / CLAVULANATE COMBO'S</b>		
	AUGMENTIN XR		
<b>ANALGESICS, NARCOTICS</b>	<b>LONG-ACTING NARCOTICS</b>		
	DURAGESIC* morphine sulfate		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.

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<b>ANGIOTENSIN MODULATORS</b>	<b>ACE INHIBITORS</b>		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.
	benazepril captopril enalapril fosinopril lisinopril moexipril quinapril ramipril trandolapril		
	<b>ACE INHIBITORS AND DIURETICS</b>		
	benazepril/HCTZ captopril/HCTZ enalapril/HCTZ fosinopril/HCTZ lisinopril/HCTZ moexipril/HCTZ quinapril/HCTZ		
	<b>ANGIOTENSIN RECEPTOR BLOCKERS (ARBs)</b>		
		AVAPRO BENICAR COZAAR DIOVAN MICARDIS	
<b>ANTIDEPRESSANTS</b>	<b>ARBs AND DIURETICS</b>		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.
		AVALIDE BENICAR-HCT DIOVAN-HCT HYZAAR MICARDIS-HCT	
	<b>ARB COMBINATIONS</b>		
		AZOR	
	<b>STEP 1</b>		Naïve patients require a trial of one step 1 drug lasting 6 weeks prior to receiving approval for step 2 drug. For depression, a trial of a step 2 drug for 6 weeks is required prior to approval of non-preferred agents. <b>Cymbalta*, Lexapro** and Pristiq are Step 3 (non-preferred agents).</b> Trazadone, buspirone, fluvoxamine, MAO inhibitors, TCA's, bupropion IR and venlafaxine IR do not require prior authorization but will not count towards meeting Step Therapy requirements.  Rapid-dissolve mirtazapine tablets are non-preferred.  *Cymbalta will be approved for a diagnosis of peripheral neuropathy.  **Lexapro will be approved for adolescents between the ages of 12 - 17.
	bupropion SR citalopram fluoxetine paroxetine IR sertraline		
	<b>STEP 2</b>		
EFFEXOR XR* mirtazapine paroxetine CR WELLBUTRIN XL *			
<b>ANTIVIRALS, ORAL</b>	<b>HERPES AGENTS</b>		
acyclovir famciclovir VALTREX			

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<b>CHOLESTEROL AGENTS</b>	<b>STATINS, LOW POTENCY</b>		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.
	LESCOL/LESCOL XL lovastatin pravastatin		
	<b>STATINS, HIGH POTENCY</b>		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.
	LIPITOR simvastatin		
	<b>STATIN COMBINATIONS</b>		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.
	ADVICOR CADUET SIMCOR		
	<b>FIBRIC ACID DERIVATIVES</b>		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.
fenofibrate gemfibrozil TRICOR			
<b>NICOTINIC ACID DERIVATIVES</b>			
NIASPAN			
<b>EAR</b>	<b>MISCELLANEOUS</b>		
	CIPRODEX		
<b>FIBROMYALGIA AGENTS</b>	<b>FIBROMYALGIA AGENTS</b>		Trial and failure of amitriptyline or cyclobenzaprine greater than or equal to 6 weeks in the last 12 months for the diagnosis of fibromyalgia.
		SAVELLA CYMBALTA LYRICA	
<b>GASTROINTESTINAL</b>	<b>PROTON PUMP INHIBITORS</b>		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. Prevacid Solutabs will be approved for children ≤ 8 years of age.
	KAPIDEX omeprazole PRILOSEC OTC * PROTONIX *		
<b>GROWTH HORMONE</b>	<b>GROWTH HORMONE</b>		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 a preferred agent 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.
		GENOTROPIN NUTROPIN	

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<b>INSOMNIA AGENTS</b>	<b>NON-BENZODIAZEPINES</b>		Non-preferred products require a 14 day history of a preferred product in the last 365 days prior to approval for a non-preferred product. Rozerem is non-preferred without a history of substance abuse.
	LUNESTA zaleplon zolpidem		
<b>MIGRAINE AGENTS</b>	<b>TRIPTANS</b>		Trial and failure of ALL preferred agents each greater than or equal to 14 days is required for PA approval of non-preferred agents in the last 12 months. Quantity limits apply.
	MAXALT/MLT RELPAX sumatriptan		
<b>NSAIDS</b>	<b>NON-SELECTIVE</b>		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 NSAID.
	diclofenac etodolac fenoprofen flurbiprofen ibuprofen indomethacin ketoprofen ketorolac meclofenamate mefenamic acid meloxicam nabumetone naproxen oxaprozin sulindac tolmetin		
	<b>COX 2 INHIBITORS</b>		
	CELEBREX	Trial and failure of two (2) preferred non-selective NSAIDs greater than or equal to a 14 days supply in the last 12 months required prior to PA approval.	
<b>OPHTHALMICS</b>	<b>OP. -ANTIBIOTICS- QUINOLONES</b>		Trial and failure of a preferred agent greater than or equal to 5 days in the last 12 months. Azasite will be approved for pregnancy.
	ciprofloxacin ofloxacin VIGAMOX ZYMAR		
	<b>OP. -ANTI-INFLAMMATORY- NSAIDS</b>		Trial of each preferred agent greater than or equal to 5 days in the last 12 months.
	ACULAR/LS/PF flurbiprofen diclofenac		
	<b>OP. -BETA-BLOCKERS</b>		Trial and failure of three (3) different preferred agents each greater than or equal to 30 days in the last 12 months. Betoptic S will be approved for those with heart and lung conditions.
	betaxolol carteolol levobunolol metipranolol timolol		
	<b>OP. -CARBONIC ANHYDRASE INHIBITOR</b>		Trial of a preferred agent greater than or equal to 30 days in the last 12 months.
	dorzolamide		
	<b>OP. -CARBONIC ANHYDRASE INHIBITOR COMBO</b>		Trial of a preferred agent greater than or equal to 30 days in the last 12 months.
	dorzolamide/timolol		
	<b>OP. -SYMPATHOMIMETICS</b>		Trial of a preferred agent greater than or equal to 30 days in the last 12 months.
	ALPHAGAN P brimonidine dipivefrin		
<b>OP. -PARASYMPATHOMIMETICS</b>		Trial of two (2) preferred agents greater than or equal to 30 days in the last 12 months.	
carbachol ISOPTO CARBACHOL phospholine iodide pilocarpine			
<b>OP. -MAST CELL STABILIZERS</b>		Trial and failure of two (2) preferred agents greater than or equal to 30 days in the last 12 months. Emadine, Alomide, and Alocril will be approved for pregnancy. Alomide will be approved for children under age 3.	
cromolyn ketotifen OPTIVAR PATADAY PATANOL			

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<b>OVERACTIVE BLADDER AGENTS</b>	<b>OVERACTIVE BLADDER AGENTS</b>		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. The patch will only be allowed if the patient has the inability to swallow.
	DETROL LA ENABLEX oxybutynin /ER SANCTURA / XR VESICARE		
<b>PROSTATE AGENTS</b>	<b>5-ALPHA-REDUCTASE INHIBITORS</b>		Trial and failure of a preferred agent greater than or equal to a 30 day supply in the last 12 months.
	AVODART finasteride		
	<b>ALPHA BLOCKERS</b>		
doxazosin terazosin			
<b>SKELETAL MUSCLE RELAXANTS</b>	<b>MUSCLE RELAXANTS</b>		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.
	baclofen cyclobenzaprine tizanidine		
<b>SMOKING CESSATION</b>	<b>NICOTINE REPLACEMENT</b>		Quantity limits apply. Generic bupropion SR needs to be an AB rated generic of Zyban.
	nicotine gum, lozenges, and patches		
	<b>OTHER</b>		
bupropion SR CHANTIX			
<b>STIMULANTS</b>	<b>LONG ACTING AMPHETAMINES</b>		Clients must have a diagnosis for ADD or ADHD. Prior Authorization will be required for clients under the age of 5. 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. Dosing limits will apply (150% of labeled max). Trial and failure of two (2) preferred agents (each from a different class) greater than or equal to a 30 day supply in the last 12 months.
	<b>ADDERALL XR*</b> VYVANSE		
	<b>SHORT ACTING AMPHETAMINES</b>		
	amphetamine salts combo dextroamphetamine		
	<b>STIMULANT LIKE</b>		
	STRATTERA		
	<b>LONG ACTING METHYLPHENIDATE</b>		
	CONCERTA FOCALIN XR methylin ER methylphenidate ER/CR/SR		
<b>METHYLPHENIDATE</b>			
dexmethylphenidate methylin (tabs) methylphenidate			
<b>TOPICAL AGENTS</b>	<b>IMMUNOMODULATORS</b>		
	ELIDEL PROTOPIC		