

**WYOMING MEDICAID  
Preferred Drug List (PDL) - OCTOBER 12, 2011**

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.

\*Indicates **BRAND** is Preferred. May Use DAW 5. Yellow highlighted items below indicate new changes to the PDL. Red font indicates quantity/dosage limits apply.  
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),  
Epocrates, 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 <small>GENERIC MANDATORY POLICY APPLIES THIS LIST IS NOT ALL INCLUSIVE PLEASE CONTACT GHS FOR QUESTIONS</small>	
ALLERGY / ASTHMA	<b>ANTI-HISTAMINES, MINIMALLY SEDATING</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.	ALAVERT CLARINEX levocetirizine	
	cetirizine fexofenadine loratadine				<b>ANTI-HISTAMINE/DECONGESTANT COMBINATIONS</b>
	cetirizine/pseudoephedrine fexofenadine/pseudoephedrine loratadine/pseudoephedrine		<b>ANTICHOLINERGIC BRONCHODILATORS</b>	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-  <b>Spiriva 5 day STARTER package will be allowed one (1) time per recipient.</b>	ATROVENT HFA
	ipratropium SPIRIVA		<b>CORTICOSTEROID / BRONCHODILATOR COMBO'S</b>		
	ADVAIR/HFA DULERA SYMBICORT		<b>LEUKOTRIENE MODIFIERS</b>	Trial and failure of preferred agent greater than or equal to 30 days in the last 12 months will be	SINGULAIR GRANULES (use preferred) zafirlukast ZYFLO
	SINGULAIR		<b>NASAL ANTIHISTAMINES</b>		
	azelastine		<b>NASAL STEROIDS</b>	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 flunisolide OMNARIS RHINOCORT VERAMYST
	fluticasone <b>NASACORT AQ*</b> NASONEX		<b>SHORT ACTING BRONCHODILATORS - INHALERS</b>		
	PROAIR HFA PROVENTIL HFA VENTOLIN HFA XOPENEX HFA		<b>SHORT ACTING BRONCHODILATORS - NEBULIZERS</b>	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 METAPROTERENOL PROVENTIL XOPENEX
	albuterol neb		<b>STEROID INHALANTS</b>		
	ASMANEX budesonide FLOVENT HFA/DISK QVAR		<b>ALZHEIMER AGENTS</b>	Client must have a diagnosis of dementia.	ARICEPT 23MG (use preferred) ARICEPT ODT (use preferred) donepezil (BRAND IS PREFERRED)
	ALZHEIMERS		<b>ARICEPT*</b> EXELON PATCH/SOLUTION galantamine/ER NAMENDA rivastigmine capsules		
ANALGESICS		<b>SUBOXONE/FILM</b>			

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ANALGESICS <i>Continued</i>	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-III's and C-IV's 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 requires trial of preferred and client must have diagnosis of drug/substance abuse.	AVINZA BUTRANS** EMBEDA*** KADIAN NUCYNТА ER OPANA ER oxymorphone ER (7.5mg/15mg) OXYCONTIN/CR
	fentanyl patch			
	morphine sulfate			
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 NUCYNТА oxymorphone oxycodone/IBU
codeine sulfate				
hydromorphone				
			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 RYZOLT tramadol/apap tramadol ER
TRAMADOL PRODUCTS		tramadol		
ANDROGENS	TESTOSTERONE TOPICAL GELS		Testosterone agents are only allowed for diagnosis of hypogonadism or insufficient testosterone production.	
		ANDROGEL TESTIM GEL		
ANGIOTENSIN MODULATORS	ACE 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.	
	benazepril			
	captopril			
	enalapril			
	fosinopril			
	lisinopril			
moexipril				
perindopril				
quinapril				
ramipril				
trandolapril				
ACE INHIBITORS AND DIURETICS			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/HCTZ				
captopril/HCTZ				
enalapril/HCTZ				
fosinopril/HCTZ				
lisinopril/HCTZ				
moexipril/HCTZ				
quinapril/HCTZ				
ANGIOTENSIN RECEPTOR BLOCKERS (ARBs)		AVAPRO BENICAR DIOVAN losartan MICARDIS	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.	ATACAND EDARBI TEVETEN
ARBs AND DIURETICS				
		AVALIDE		ATACAND HCT
		BENICAR HCT		TEVETEN HCT
		DIOVAN HCT		
		losartan HCT		
		MICARDIS HCT		
ARB COMBINATIONS				
		AZOR		TWYNSTA (use separate agents)
		EXFORGE/EXFORGE-HCT		
ANTIBIOTICS	QUINOLONES			AVELOX ABC PROQUIN
	AVELOX			
	ciprofloxacin/ER			
	FACTIVE			
	levofloxacin			
NOROXIN				
ofloxacin				

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ANTICOAGULANTS	LOW MOLECULAR WEIGHT HEPARIN (LMWH)			enoxaparin (BRAND IS PREFERRED) <b>LOVENOX 300MG/3ML (USE PREFERRED)</b>	
	ARIXTRA FRAGMIN LOVENOX*				
ANTICONVULSANTS	DIAZEPAM RECTAL GEL			diazepam gel (BRAND IS PREFERRED)	
	DIASTAT*				
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 <b>will not count</b> 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><b>fluoxetine 20mg tablets (use preferred)</b> <b>mirtazapine 7.5mg and mirtazapine rapid-dissolve tablets (use preferred)</b></p> <p>APLENZIN CYMBALTA* LEXAPRO** PRISTIQ VIIBRYD</p> <p><b>Effective 1/1/12, venlafaxine ER tablets will be non-preferred .</b></p>	
	bupropion ER/SR/XL citalopram fluoxetine mirtazapine 15, 30, and 45mg paroxetine IR/CR sertraline venlafaxine ER				
ANTIPSYCHOTICS	ATYPICAL ANTIPSYCHOTICS		<p>Non-preferred agents (Fanapt, Latuda, and Saphris) require a trial of ALL preferred agents at max doses.</p> <p>Dosage limits apply:                      ABILIFY &lt;13 years of age: 23mg/day                      ABILIFY ≥13 years of age: 45mg/day                      GEODON ≤ 17 years of age: 180mg/day                      GEODON &gt; 17 years of age: 300mg/day                      INVEGA all ages: 18mg/day                      Risperidone ≤ 17 years of age: 5mg/day                      Risperidone &gt; 17 years of age: 24mg/day                      SEROQUEL &lt;13 years of age: 600mg/day                      SEROQUEL 13-17 years of age: 900mg/day                      SEROQUEL &gt; 17 years of age: 1200mg/day                      ZYPREXA &lt; 13 years of age: 15mg/day                      ZYPREXA ≥ 13 years of age: 15mg/day</p>	<p>ABILIFY ODT (use preferred) FANAPT LATUDA SAPHRIS SEROQUEL XR (use preferred; CURRENT USERS WILL BE GRANDFATHERED)</p>	
		<p>ABILIFY GEODON INVEGA INVEGA SUSTENNA RISPERDAL CONSTA risperidone SEROQUEL ZYPREXA ZYPREXA RELPREVV</p>			
	SPECIAL ATYPICAL ANTIPSYCHOTICS				Dosage limits apply (Max Dose:1350mg/day).
ANTIVIRALS, ORAL	HERPES AGENTS			valacyclovir (BRAND IS PREFERRED)	
	acyclovir famciclovir VALTREX*				
CHOLESTEROL	STATINS, LOW POTENCY		<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>	ALTOPREV	
	LESCOL/XL lovastatin pravastatin				
	STATINS, HIGH POTENCY		<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>	CRESTOR LIVALO	
	LIPITOR simvastatin				
	STATIN COMBINATIONS		<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>ADVICOR (use separate agents) CHOLESTIN PRAVIGARD VYTORIN (use separate agents)</p>	
	CADUET SIMCOR				
TRIGLYCERIDE LOWERING AGENTS		<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>ANTARA fenofibric FENOGLIDE LOVAZA TRILIPIX</p>		
fenofibrate gemfibrozil TRICOR					
INTESTINAL CHOLESTEROL ABSORPTION INHIBITOR					
	ZETIA				

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CHOLESTEROL <i>Continued</i>	BILE ACID SEQUESTRANT		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	
	cholestyramine/light colestipol				
CONTRACEPTIVES	BIPHASIC ORAL CONTRACEPTIVES		Monophasic and triphasic oral contraceptives are not included and are available without prior authorization. (generic substitution is mandatory)		
	KARIVA LO-SEASONIQUE NECON 10/11 SEASONIQUE				
COUGH AND COLD	DEXTROMETHORPHAN POLISTIREX				
	DELSYM				
DIABETES	DIABETES AGENTS		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.	FORTAMET GLUMETZA RIOMET	
	BIGUANIDES				
	metformin/ER				
	α-GLUCOSIDASE INHIBITORS			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
	acarbose				
	MEGLITINIDES			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 (BRAND IS PREFERRED) PRANDIN
	STARLIX*				
	THIAZOLIDINEDIONES			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.	ACTOS 30MG, 45MG ( use ACTOS 15mg) ACTOSPLUS MET (use separate agents) AVANDIA AVANDAMET (use separate agents)
	ACTOS 15MG				
	SULFONYLUREAS			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.	
	glimepiride/ER glipizide/ER glyburide/ER				
	DIPEPTIDYL PEPTIDASE 4 (DPP-4) INHIBITORS			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.	KOMBIGLYZE (use separate agents) TRADJENTA
		JANUMET JANUVIA ONGLYZA			
	INCRETIN MIMETICS (GLP-1 RECEPTOR AGONISTS)			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.	VICTOZA
	BYETTA				
LONG-ACTING INSULIN					
LANTUS LEVEMIR					
RAPID-ACTING INSULIN					
APIDRA HUMALOG NOVALOG					
DIABETIC METERS/TEST STRIPS		Quantity limit applies (1 meter/365days).	ALL OTHER METERS AND TEST STRIPS		
	FREESTYLE LITE FREESTYLE FREEDOM LITE ONE TOUCH ULTRA ONE TOUCH ULTRA 2 ONE TOUCH ULTRA MINI ONE TOUCH ULTRASMART PRECISION XTRA				
EAR	ANTIBIOTIC/STEROID COMBINATION SUSPENSIONS				
	CETRAXAL CIPRODEX CIPRO HC COLY-MYCIN S CORTISPORIN-TC <small>Neomycin/Polymyxin B Sulfates/Hydrocortisone</small>				
	EPOEITIN				EPOGEN
	ARANESP PROCRT				

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FIBROMYALGIA	<b>STEP 1</b>			
	amitriptyline cyclobenzaprine		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.	
	<b>STEP 2</b>			
		SAVELLA	Trial and failure of a Step 1 agent and a Step 2 agent greater than or equal to six (6) weeks in the last 12 months is required for approval of a Step 3 agent.	
GASTROINTESTINAL	<b>DIGESTIVE ENZYMES</b>		Prior authorization required.	PANCREAZE TRI-PASE
	CREON 3000, 6000, 12000, 24000 UNIT ZENPEP		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.  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.  Pantoprazole will be allowed for clients on concurrent Plavix therapy.	ACIPHEX lansoprazole NEXIUM omeprazole <b>tablets</b> (use preferred) omeprazole/bicarbonate pantoprazole VIMOVO (use separate agents)
	<b>PROTON PUMP INHIBITORS</b>			
	DEXILANT/KAPIDEX omeprazole <b>capsules</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.	ASACOL HD (use preferred) CANASA LIALDA PENTASA 500MG (use Pentasa 250mg) ROWASA
	<b>MESALAMINE</b>			
GROWTH HORMONE	APRISO ASACOL PENTASA 250MG ONLY		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
	<b>GROWTH HORMONE</b>			
HEPATITIS C		GENOTROPIN NORDITROPIN NUTROPIN/AQ	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 will be approved for pediatric patients (aged 18 and under), for retreatment, and for dosage adjustments that cannot be achieved with Pegasys.	PEG-INTRON
	<b>INTERFERON</b>			
	PEGASYS			

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IMMUNOMODULATORS	<b>IMMUNOMODULATORS (DIAGNOSIS REQUIRED)</b>		<p>Client must have <b>diagnosis prior to approval</b> for <b>preferred agents</b> (outlined below):</p> <p><b>Enbrel</b>: Ankylosing Spondylitis (AS), Juvenile Idiopathic Arthritis (JIA), Plaque Psoriasis (PP), Psoriatic Arthritis (PA), Rheumatoid Arthritis (RA)**</p> <p><b>Humira</b>: 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 <b>non-preferred agents</b>, 60-day trial and failure of a preferred agent is required and client must have diagnosis prior to approval (outlined below):</p> <p><b>Actemra</b>: RA (60-day trial of methotrexate is required)</p> <p><b>Amevive</b>: PP</p> <p><b>Cimzia</b>: Crohn's***, RA</p> <p><b>Kineret</b>: RA</p> <p><b>Orencia</b>: JIA, RA</p> <p><b>Remicade</b>: AS, Crohn's, PP, PA, RA, Ulcerative Colitis****</p> <p><b>Rituxan</b>: RA</p> <p><b>Simponi</b>: AS, PA, RA</p> <p><b>Stelara</b>: PP</p> <p><b>Tysabri</b>: 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	<b>NON-BENZODIAZEPINES</b>		<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>Rozereem 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) LUNESTA ROZEREM zolpidem ER ZOLPIMIST (additional criteria applies)</p>
MIGRAINE	<b>TRIPTANS</b>		<p>Trial and failure of a preferred agent will be required before approval can be given for a non-preferred agent.</p> <p>Quantity limits apply:</p> <p>MAXALT MLT 5mg: 27tabs/34days MAXALT MLT 10mg: 14tabs/34days 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>AXERT FROVA MAXALT RELPAX TREMIMET ZOMIG</p>
MULTIPLE SCLEROSIS	<b>MULTIPLE SCLEROSIS AGENTS</b>		<p>Trial and failure of one (1) interferon agent AND failure of Copaxone.</p> <p>For Tysabri, in addition to the above criteria, additional prior authorization criteria applies.</p>	<p>EXTAVIA GILENYA TYSABRI (additional criteria applies)</p>
NSAIDS	<b>NSAIDS</b>		<p>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.</p> <p>Dosing limits apply for ketorolac (limit 5days/34 days).</p>	<p>CALDOLOR CAMBIA POWDER CELEBREX FLECTOR (additional criteria applies) NAPRELAN NEOPROFEN PENNSAID (additional criteria applies) SOLARAZE (additional criteria applies) SPRIX (additional criteria applies) VOLTAREN (additional criteria applies) ZIPSOR</p>

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OPHTHALMICS	<b>OP. -ANTIBIOTICS- QUINOLONES</b>		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.	AZASITE BESIVANCE IQUIX levofloxacin ZYMAXID
	<b>OP. -ANTI-INFLAMMATORY- NSAIDS</b>		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.	ACULAR/LS/PF (use preferred) ACUVAIL BROMDAY bromfenac NEVANAC
	<b>OP. -BETA-BLOCKERS</b>		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.	BETIMOL BETOPTIC S ISTALOL
	<b>OP. -CARBONIC ANHYDRASE INHIBITOR</b>		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.	AZOPT
	<b>OP. -CARBONIC ANHYDRASE INHIBITOR COMBO</b>		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.	
	<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 will be required before approval can be given for a non-preferred agent.  Emadine, Alomide, and Alocril will be approved for pregnancy.  Alomide will be approved for children under the age of 3.	ALAMAST alaway ALOCRIL ALOMIDE ALREX BEPREVE CLARITIN OTC ELESTAT EMADINE LASTACAF ZYRTEC ITCHY EYE
	<b>OP. -PROSTAGLANDINS</b>		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.	latanoprost
OPHTHALMICS <i>Continued</i>	<b>OP. -SYMPATHOMIMETICS</b>		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.	
	<b>OP. -SYMPATHOMIMETIC COMBO</b>		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.	
	<b>BISPHOSPHONATES</b>		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 BONIVA FOSAMAX-D
	<b>NASAL CALCITONIN</b>			
OSTEOPOROSIS	<b>OP. -SYMPATHOMIMETICS</b>		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.	
OVERACTIVE BLADDER	<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.  Oxytrol will be approved for clients that have an inability to swallow.	DETROL/LA ENABLEX GELNIQUE GEL 10% OXYTROL DIS SANCTURA XR

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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>
PRENATAL VITAMINS	<b>PRENATAL VITAMINS</b> BP MULTINATL TAB PLUS COMPLETE-RF CO-NATAL FA ELITE-OB FOLIVANE-OB INATAL ULTRA LACTOCAL-F MARNATAL-F MAXINATE NATAFORT O-CAL PRENAFIRST PRENAPLUS PRENATABS RX PRENATAL 19/CHEWABLE PRENATAL LOW IRON PRENATAL PLUS/FE SE-CARE/CHEWABLE SELECT-OB CHEWABLE SE-NATAL 19/CHEWABLE SE-NATAL 90 SE-NATAL ONE TARON-BC TARON-EC CAL TRIMESIS RX TRINATAL RX TRINATE TRI RX TRIVEEN-U VINATE II VINATE AZ VINATE C VINATE CAL VINATE IC VINATE M VINATE ONE VINATE ULTRA VITASPIRE VOL-TAB RX VOL-PLUS			ALL OTHER PRENATAL VITAMINS INCLUDING OVER-THE-COUNTER FORMULATIONS
PROSTATE	<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 will be required before approval can be given for a non-preferred agent.	JALYN (use separate agents)
	<b>ALPHA BLOCKERS</b>		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 (use separate agents)
PULMONARY ANTIHYPERTENSIVES	<b>ENDOTHELIN RECEPTOR ANTAGONISTS</b>			
	LETAIRIS TRACLEER			
SKELETAL MUSCLE RELAXANTS	<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.	carisoprodol chlorzoxazone cyclobenzaprine ER metaxalone methocarbamol orphenadrine
	baclofen cyclobenzaprine tizanidine			
SMOKING CESSATION	<b>NICOTINE REPLACEMENT</b>		Generic bupropion SR needs to be an AB rated generic of Zyban.	
		nicotine gum, lozenges, and patches		
		<b>OTHER</b>	Concomitant use of Chantix with bupropion SR or other nicotine replacement therapies will not be allowed.  Quantity limits apply: NICOTINE GUM: 735pcs &/or 84 days/365days NICOTINE LOZ: 735 pcs &/or 84 days/365days NICOTINE PATCH 5mg: 14 PATCH &/or 14 days/365days NICOTINE PATCH 7mg: 14 PATCH &/or 14 days/365days NICOTINE PATCH 10mg: 14 PATCH &/or 14 days/365days NICOTINE PATCH 14mg: 14 PATCH &/or 14 days/365days NICOTINE PATCH 15mg: 84 PATCH &/or 84 days/365days NICOTINE PATCH 21mg: 42 PATCH &/or 42 days/365days bupropion SR: 168 tabs &/or 84 days/365days CHANTIX: 168 tabs &/or 84 days/365days	
	bupropion SR CHANTIX			

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STIMULANTS	<b>AMPHETAMINES</b>		<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            FOCALIN: 30mg/day            FOCALIN XR ≤ 13 years of age: 45mg/day            FOCALIN XR &gt; 13 years of age: 60mg/day            methylin/methylphenidate: 135mg/day            methylin/methylphenidate ER/CR/SR: 135mg/day            VYVANSE: 105mg/day</small></p>	<p><b>AMPHETAMINES:</b> amphetamine salts combo ER (BRAND IS PREFERRED) <u>ADDERALL XR WILL ONLY BE PREFERRED FOR THOSE CLIENTS CURRENTLY ON THE MEDICATION.</u></p> <p><b>METHYLPHENIDATES:</b> dexmethylphenidate/ER (BRAND IS PREFERRED) METADATE CD methylphenidate ER 18,27,36mg (BRAND IS PREFERRED) RITALIN LA</p>
	<small>LONG ACTING AMPHETAMINES</small>			
		<p><b>ADDERALL XR*</b> VYVANSE dextroamphetamine CR</p>		
	<small>IMMEDIATE RELEASE AMPHETAMINES</small>			
		<p>amphetamine salts combo dextroamphetamine</p>		
	<b>METHYLPHENIDATES</b>			
	<small>LONG ACTING METHYLPHENIDATES</small>			
		<p><b>CONCERTA*</b> DAYTRANA FOCALIN XR methylin ER methylphenidate ER/CR/SR</p>		
	<small>IMMEDIATE RELEASE METHYLPHENIDATES</small>			
		<p><b>FOCALIN*</b> methylin (tabs) methylphenidate</p>		
<b>SELECTIVE ALPHA-ADRENERGIC AGONIST</b>		<p>To obtain the <b>non-preferred agent</b>, 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 <b>AND</b> 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	
<small>GUANFACINE AGENTS</small>				
guanfacine				
<small>CLONIDINE AGENTS</small>		<p>Trial and benefit of clonidine IR will be required prior to approval of the non-preferred agent.</p>	KAPVAY	
clonidine				

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TOPICAL AGENTS	<b>IMPETIGO ANTIBIOTICS</b>		Trial and failure of ALL preferred agents greater than or equal to 7 days in the past 90 days.  Use smallest size appropriate for 7 day trial.	ALTABAX
	gentamicin mupirocin			
	<b>BENZOYL PEROXIDE/CLINDAMYCIN COMBOS</b>		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.	ACANYA
		benzoyl peroxide/clindamycin		
	<b>CORTICOSTEROIS</b> <small>C=CREAM; G=GEL; L=LOTION; O=OINTMENT</small> <b>LOW POTENCY</b>		Trial and failure of ALL preferred agents greater than or equal to 14 days in the last 90 days.	PANDEL
	alclometasone desonide fluocinolone 0.01% hydrocortisone butyrate 0.1% (C) hydrocortisone 1%, 2.5% (C,L,O) prednicarbate			
	<b>MEDIUM POTENCY</b>		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
	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%			
	<b>HIGH POTENCY</b>		Trial and failure of ALL preferred agents greater than or equal to 14 days in the last 90 days.	HALOG
	amcinonide betamethasone dipropionate clobetasol desoximetasone 0.25%, 0.05% (G) diflorasone fluocinonide flurandrenolide fluticasone 0.005% (O) halobetasol triamcinolone 0.5%			
	<b>IMIQUIMODS</b>		Trial and failure of a preferred agent greater than or equal to 28 days in the last 12 months will be required before approval can be given for a non-preferred agent.	imiquimod (BRAND IS PREFERRED) ZYCLARA
	ALDARA*			
	<b>IMMUNOMODULATORS</b>			
	ELIDEL PROTOPIC			
	<b>MISC TOPICAL</b>		Tazorac is allowed for clients with the diagnosis of psoriasis for all ages.  For the treatment of acne vulgaris, acne combinations are limited to those clients under the age of 21.	
		TAZORAC		
	<b>SALICYLIC ACID</b>			All other topical salicylic acid formulations.
	aliclén shampoo 6% salacyn cream/lotion 6% Salicylic Acid Shampoo 6%			
	<b>UREA</b>			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%			