



# Virginia Medicaid's Preferred Drug List (PDL)/ Common Core Formulary

7/1/21

## Virginia Medicaid's Pharmacy Benefits Management System

Phone: 800-932-6648 Fax: 800-932-6651

### General Information:

- Virginia Medicaid's Preferred Drug List (PDL)/ Common Core Formulary only includes select drug classes, other classes will pay such as but not limited to diuretics, many cardiac agents, many antibiotics etc.
- PDL preferred drugs do not require Service Authorizations (SA) unless subject to additional clinical criteria (e.g., long acting opioids, hepatitis C therapies, growth hormone)
- Non-preferred drugs require a SA
- Drugs not on the PDL are subject to Virginia's mandatory generic substitution requirements.
- SAs may be submitted by fax, phone or WebPA. For urgent requests, please call **800-932-6648**. Fax requests receive a response within 24 hours.

PDL drug coverage information can be found at <http://www.VirginiaMedicaidPharmacyServices.com>. The following "routine" PDL criteria guidelines will be applied to all non-preferred drugs.

1. Is there any reason the member cannot be changed to a preferred drug within the same class? Acceptable reasons include:
  - Allergy to preferred drug.
  - Contraindication to or drug-to-drug interaction with preferred drug.
  - History of unacceptable/toxic side effects to preferred drug.
  - Member's condition is clinically stable; changing to a preferred drug might cause deterioration of the member's condition.
2. The requested drug may be approved if both of the following are true:
  - There has been a therapeutic failure of at least **two** preferred drugs **within the same class as appropriate for diagnosis unless otherwise noted in the clinical criteria**. A therapeutic failure of only one preferred drug is required when there is only one preferred drug within a therapeutic class.
  - The requested drug's corresponding generic (if a generic is available **and** covered by the State) has been attempted and failed or is contraindicated.

All changes from last posting will be highlighted in yellow.

\*\*Members currently receiving aripiprazole oral solution, Geodon® (IM), Nuplazid or olanzapine/fluoxetine will be "grandfathered" for a period not to exceed one year. After that time, the prescriber will need to submit a service authorization request documenting the medical necessity of the non-preferred drug.

### LEGEND

AG = age edit

CE = clinical edit

ST = step edit

QL = quantity limit

cap = capsule

cr = cream

ER = extended release

inj = injection

IR = immediate release

ODT = oral disintegrating tablet

oint = ointment

soln = solution

supp = suppository

susp = suspension

tab = tablet



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Analgesics</b>		
<b>* Opioids – Long Acting (LAO)</b>		
<b>Preferred (Sch III-VI)</b>	<b>Non-Preferred</b>	*All Long Acting Opioids (preferred and non-preferred) require submission of a Clinical SA. Refer to combined short/long-acting opioid SA form <a href="#">(Short &amp; Long Acting Opioid SA Form)</a>
<b>Butrans<sup>®</sup></b> (buprenorphine) Transdermal Patch	<i>Belbuca</i> (buprenorphine buccal film) <i>buprenorphine</i> (generic <b>Butrans<sup>®</sup></b> ) <i>ConZip<sup>®</sup></i> (tramadol ER) <i>Ryzolt<sup>TM</sup></i> (tramadol ER) <i>tramadol ER</i> <i>Ultram ER<sup>®</sup></i> (tramadol ER)	
<b>Preferred (Sch II)</b>	<b>Non-Preferred</b>	<b>LENGTH OF AUTHORIZATIONS</b>
fentanyl 12, 25, 50, 75 & 100 mcg patches morphine sulfate ER tab	<i>Arymo<sup>TM</sup> ER</i> <i>Duragesic<sup>®</sup></i> <i>Embeda</i> <i>Exalgo<sup>®</sup></i> <i>fentanyl 37.5 mcg, 62.5 mcg, and 87.5 mcg patches</i> <i>hydromorphone ER</i> <i>Hysingla ER<sup>TM</sup></i> <i>Kadian<sup>®</sup> ER</i> <i>Morphabond<sup>TM</sup> ER</i> <i>morphine ER cap (generic Avinza<sup>®</sup>)</i> <i>morphine ER cap (generic Kadian<sup>®</sup>)</i> <i>MS Contin<sup>®</sup></i> <i>Nucynta<sup>®</sup> ER</i> <i>Oramorph<sup>®</sup> SR<sup>®</sup></i> <i>oxycodone-long acting</i> <i>OxyContin<sup>®</sup></i> <i>oxymorphone ER</i> <i>Xartemis<sup>TM</sup> XR</i> <i>Xtampza ER<sup>®</sup></i> <i>Zohydro ER<sup>TM</sup></i>	• Up to 3 months for (includes HIV/AIDS, Chronic back pain, Arthritis, Fibromyalgia, Diabetic neuropathy, Postherpetic Neuralgia). • Up to 6 months for chronic pain (includes Cancer pain, Sickle cell disease, Palliative care, End-of-Life Care, Hospice).  Daily dose limits have been established for all LAO. Quantity limits can be found at : <a href="#">Daily Dose Limits for Short &amp; Long Acting Opioids</a>



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<b>*Methadone Drugs</b>			
		<i>Dolophine<sup>®</sup></i> <i>Methadose<sup>®</sup> oral soln &amp; tab</i> <i>methadone oral soln &amp; tab</i>	*Methadone requires the completion of the Clinical SA form ( <a href="#">Methadone SA Form</a> ) unless prescribed for neonatal abstinence syndrome for an infant under the age of one.
<b>*Opioids – Short Acting</b>			
<b>*Transmucosal Immediate Release Fentanyl</b>			
		<i>Actiq<sup>®</sup></i> <i>Fentora<sup>®</sup></i> <i>fentanyl citrate</i> <i>Lazanda<sup>®</sup></i> <i>Subsys<sup>®</sup></i>	<b>LENGTH OF AUTHORIZATIONS:</b> <ul style="list-style-type: none"> <li>Up to 3 months for (includes HIV/AIDS, Chronic back pain, Arthritis, Fibromyalgia, Diabetic neuropathy, Postherpetic Neuralgia).</li> <li>Up to 6 months for chronic pain (includes Cancer pain, Sickle cell disease, Palliative care, End-of-Life Care, Hospice).</li> </ul>
<b>Short-Acting Opioids</b>			
codeine/APAP hydrocodone/APAP hydrocodone/ibuprofen hydromorphone morphine IR oxycodone IR oxycodone/APAP tramadol HCl 50mg tramadol HCl/APAP		<i>Abstral<sup>®</sup></i> <i>Apadaz<sup>™</sup></i> <i>codeine tab/soln</i> <i>butalbital comp with codeine</i> <i>butalbital/caffeine/APAP w/codeine</i> <i>butorphanol tartrate nasal</i> <i>dihydrocodeine/APAP/caffeine</i> <i>dihydrocodeine/ASA/caffeine</i> <i>hydromorphone liq/supp</i> <i>meperidine tab</i> <i>morphine supp</i> <i>Nucynta<sup>®</sup></i> <i>Oxaydo<sup>®</sup></i> <i>oxycodone/APAP(generic</i> <i>PrimLev<sup>™</sup>)</i> <i>oxycodone conc</i> <i>oxycodone oral syringe</i> <i>oxycodone/ASA</i> <i>oxycodone/ibuprofen</i> <i>oxymorphone HCl</i> <i>Panlor<sup>®</sup></i> <i>pentazocine/naloxone</i> <i>PrimLev<sup>™</sup></i>	*All Short-Acting Opioids ( <b>preferred and non-preferred</b> ) require the submission of a Clinical SA if prescribed for > 7 days or if more than two 7 day supply prescriptions within 60 days. Refer to combined short/long-acting opioid SA form ( <a href="#">Short &amp; Long Acting Opioid SA Form</a> )

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	Preferred Agents	Non-Preferred Agents	SA Criteria																																										
		RoxyBond™ tramadol 100 Mg Ultracet® Ultram® Zamacet® soln																																											
	<b>Opioid Dependency</b> <b>CLOSED CLASS</b>																																												
	*buprenorphine SL *Suboxone® film *Sublocade™ SQ  naloxone syringe & vial Naloxone Carpuject naltrexone tab Narcan® Nasal Spray Vivitrol®	*Bunavail™ *buprenorphine/naloxone tab SL *buprenorphine/naloxone film SL *Cassipa® *Probuphine® implant *Zubsolv™  Evzio® injection	<p><b>*All Buprenorphine Containing Drugs (non-preferred) require submission of Clinical SA. Refer to <a href="#">(Sublocade Form)</a> or <a href="#">(Oral Buprenorphine SA Form)</a></b></p> <p><b>Quantity Limits</b></p> <table border="1" data-bbox="1075 643 1698 1312"> <tbody> <tr><td>Bunavail™ 2.1 0.3mg buccal film</td><td>1/day</td></tr> <tr><td>Bunavail™ 4.2 0.7mg buccal film</td><td>2/day</td></tr> <tr><td>Bunavail™ 6.3 1mg buccal film</td><td>3/day</td></tr> <tr><td>buprenorphine SL tab 2mg</td><td>3/day</td></tr> <tr><td>buprenorphine SL tab 8mg</td><td>2/day</td></tr> <tr><td>buprenorphine/naloxone SL tab 2–0.5mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL tab 8–2mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL film 2–0.5mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL film 4–1mg</td><td>1/day</td></tr> <tr><td>buprenorphine/naloxone SL film 8–2mg</td><td>3/day</td></tr> <tr><td>Cassipa® 16mg-4mg</td><td>1/day</td></tr> <tr><td>Suboxone® SL film 2–0.5mg</td><td>3/day</td></tr> <tr><td>Suboxone® SL film 4–1mg</td><td>1/day</td></tr> <tr><td>Suboxone® SL film 8–2mg</td><td>3/day</td></tr> <tr><td>Suboxone® SL film 12–3mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 0.7–0.18 mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 1.4–0.36mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 2.9–0.71mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 5.7–1.4mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 8.6–2.1mg</td><td>2/day</td></tr> <tr><td>Zubsolv™ SL tab 11.4–2.9mg</td><td>2/day</td></tr> </tbody> </table>	Bunavail™ 2.1 0.3mg buccal film	1/day	Bunavail™ 4.2 0.7mg buccal film	2/day	Bunavail™ 6.3 1mg buccal film	3/day	buprenorphine SL tab 2mg	3/day	buprenorphine SL tab 8mg	2/day	buprenorphine/naloxone SL tab 2–0.5mg	3/day	buprenorphine/naloxone SL tab 8–2mg	3/day	buprenorphine/naloxone SL film 2–0.5mg	3/day	buprenorphine/naloxone SL film 4–1mg	1/day	buprenorphine/naloxone SL film 8–2mg	3/day	Cassipa® 16mg-4mg	1/day	Suboxone® SL film 2–0.5mg	3/day	Suboxone® SL film 4–1mg	1/day	Suboxone® SL film 8–2mg	3/day	Suboxone® SL film 12–3mg	2/day	Zubsolv™ SL tab 0.7–0.18 mg	2/day	Zubsolv™ SL tab 1.4–0.36mg	2/day	Zubsolv™ SL tab 2.9–0.71mg	2/day	Zubsolv™ SL tab 5.7–1.4mg	2/day	Zubsolv™ SL tab 8.6–2.1mg	2/day	Zubsolv™ SL tab 11.4–2.9mg	2/day
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<b>Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)</b>		
<b>Oral NSAIDs</b>		<b>LENGTH OF AUTHORIZATIONS: 1 year</b>
<p><b>Children's Motrin<sup>®</sup> susp (OTC)</b>  <b>ibuprofen cap</b>  <b>ibuprofen tab (OTC &amp; Rx)</b>  <b>Infant's ibuprofen drops</b>  <b>meloxicam tab</b>  <b>naproxen tab</b>  <b>naproxen sodium (OTC)</b>  <b>naproxen EC (Rx)</b>  <b>sulindac</b></p>	<p><i>Anaprox<sup>®</sup> IR &amp; DS<sup>®</sup></i>  <i>Advil<sup>®</sup></i>  <i>Aleve<sup>®</sup></i>  <i>Arthrotec<sup>®</sup></i>  <i>Cataflam<sup>®</sup></i>  <i>*Celebrex<sup>®</sup> &amp; *celecoxib</i>  <i>Daypro<sup>®</sup></i>  <i>diclofenac potassium</i>  <i>diclofenac sodium SR</i>  <i>diclofenac sodium/misoprostol</i>  <i>diflunisal</i>  <i>Duexis<sup>®</sup></i>  <i>etodolac IR &amp; SR</i>  <i>Feldene<sup>®</sup></i>  <i>fenoprofen</i>  <i>flurbiprofen</i>  <i>ibuprofen tab chew OTC</i>  <i>Indocin<sup>®</sup> supp</i>  <i>indomethacin IR, SR &amp; rectal</i>  <i>ketoprofen IR &amp; ER</i>  <i>ketorolac</i>  <i>ketorolac tromethamine (generic for Sprix<sup>®</sup> nasal spray)</i>  <i>meclofenamate</i>  <i>mefenamic</i>  <i>meloxicam susp</i>  <i>meloxicam (generic Vivlodex<sup>™</sup>)</i>  <i>Mobic<sup>®</sup></i>  <i>Motrin<sup>®</sup></i>  <i>nabumetone</i>  <i>Nalfon<sup>®</sup></i>  <i>Naprelan<sup>®</sup></i></p>	<p><b>Routine PDL edits plus</b></p> <p><b>*Step edit required for Celebrex<sup>®</sup> and celecoxib</b></p> <ul style="list-style-type: none"> <li>• History of a trial of a minimum of two (2) different non-COX2 NSAIDs within the past year; <b>OR</b></li> <li>• Concurrent use of anticoagulants (i.e., warfarin, heparin, etc.), methotrexate, oral corticosteroids; <b>OR</b></li> <li>• History of previous GI bleed or conditions associated with GI toxicity risk factors (i.e., PUD, GERD, etc); <b>OR</b></li> <li>• Specific indication for Celebrex<sup>®</sup> for which preferred drugs are not indicated.</li> </ul>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>Naprosyn<sup>®</sup></i> <i>naproxen CR (generic Naprelan<sup>®</sup>)</i> <i>naproxen-esomeprazole mag (generic Vimovo<sup>®</sup>)DR</i> <i>naproxen sodium (RX)</i> <i>naproxen susp</i> <i>oxaprozin</i> <i>piroxicam</i> <i>Ponstel<sup>®</sup></i> <i>Prevacid Naprapac<sup>®</sup></i> <i>Sprix<sup>®</sup> nasal spray</i> <i>Tivorbex<sup>™</sup></i> <i>tolmetin sodium</i> <i>Vimovo<sup>®</sup></i> <i>Vivlodex<sup>™</sup></i> <i>Voltaren<sup>®</sup>XR</i> <i>Zipsor<sup>®</sup></i> <i>Zorvolex<sup>™</sup></i>	
	<b>Topical NSAIDs</b>		<b><u>LENGTH OF AUTHORIZATIONS:</u> 1 year</b>
	<b>diclofenac sodium 1% gel</b> <b>diclofenac sodium 1% gel (OTC)</b>	<b>**diclofenac sodium 3 % gel</b> <i>*Flector<sup>®</sup> patch (QL)</i> <i>Licart<sup>™</sup> patch (diclofenac epolamine 0.013gm (QL)</i> <i>*Pennsaid<sup>®</sup> top soln, soln pkt &amp; pump</i> <b>**Solaraze 3% top gel</b> <i>*Vopac MDS</i> <i>*Voltaren<sup>®</sup> 1% gel</i> <b>*Voltaren<sup>®</sup> 1% gel (OTC)</b> <i>*Xrylix<sup>™</sup> Kit</i>	<b>Routine PDL edits plus</b> <b><u>Clinical Criteria for Non-Preferred Topical NSAIDs; *Flector<sup>®</sup>, Pennsaid<sup>®</sup>, Vopac MDS, &amp; Xrylix<sup>™</sup> Kit:</u></b> <ul style="list-style-type: none"> <li>• Approval is based on member failing the oral generic of the desired drug and at least one other preferred NSAID (to equal a total of at least two preferred). For example, a member who failed ibuprofen or naproxen will still need to try oral diclofenac for approval of Flector<sup>®</sup>.</li> <li>• Pennsaid<sup>®</sup>, Vopac MDS, and Xrylix<sup>™</sup> Kit can only be approved for the FDA approved indication of osteoarthritis of the knee.</li> </ul> <b><u>Quantity limit for Flector<sup>®</sup> and Licart<sup>™</sup> patch = 30 patches per RX</u></b>  <b>**Solaraze<sup>®</sup> 3% &amp; Diclofenac Sodium 3 % Clinical Criteria:</b> <ul style="list-style-type: none"> <li>• Approved only for the topical treatment of actinic keratosis</li> </ul>



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<b>Antibiotic-Anti-Infective</b>			
	<b>*Antibiotics, Inhaled</b>	<b>CLOSED CLASS</b>	
	Bethkis <sup>®</sup> (QL, AG) Kitabis <sup>™</sup> Pak (QL, AG) **Tobi Podhaler <sup>®</sup> (QL, AG, SE)	***Arikayce <sup>®</sup> (amikacin liposome) Cayston <sup>®</sup> Tobi <sup>®</sup> inhalation neb soln (QL, AG) tobramycin (generic Bethkis <sup>®</sup> ) tobramycin Pak (generic Kitabis <sup>™</sup> Pak) (QL, AG) tobramycin inhalation neb soln (generic Tobi <sup>®</sup> inhalation) (QL, AG)	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p><b>Routine PDL edits plus</b></p> <p>**Tobi Podhaler<sup>®</sup></p> <ul style="list-style-type: none"> <li>Requires a clinical reason as to why one of the preferred tobramycin inhalation nebulizer solutions cannot be used (Bethkis<sup>®</sup> or Kitabis<sup>™</sup>).</li> </ul> <p>***<b>Clinical Criteria for Arikayce<sup>®</sup></b></p> <p>Duration of Approval: 12 months</p> <p><b>Initial Approval Criteria</b></p> <ul style="list-style-type: none"> <li>Patient is ≥ 18 years of age; AND</li> <li>Diagnosis of Mycobacterium avium complex (MAC) lung disease as determined by the following:               <ul style="list-style-type: none"> <li>chest radiography or high-resolution computed tomography (HRCT) scan; AND</li> <li>at least 2 positive sputum cultures; AND</li> <li>other conditions such as tuberculosis and lung malignancy have been ruled out; AND</li> </ul> </li> <li>Patient has failed a multi-drug regimen with a macrolide (clarithromycin or azithromycin), rifampin, and ethambutol. (Failure is defined as continual positive sputum cultures for MAC while adhering to a multi-drug treatment regimen for a minimum duration of 6 months); AND</li> <li>Patient has documented failure or intolerance to aerosolized administration of amikacin solution for injection, including pretreatment with a bronchodilator; AND</li> <li>Arikayce will be prescribed in conjunction with a multi-drug antimycobacterial regimen</li> </ul> <p>*Minimum age for use is 6 years for all tobramycin inhalation nebulizer solution (Bethkis<sup>®</sup>, Kitabis<sup>™</sup> Pak, Tobi<sup>®</sup> and Tobi Podhaler<sup>®</sup>) and 7 years for Cayston<sup>®</sup>.</p>



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		<p><b>Quantity Limits:</b>            Arikayce = 590 mg/8.4 Ml (28 vials)/28 days            Each carton contains a 28-day supply of medication (28 vials)            Bethkis® = 224Ml (56 amps)/28 days            Cayston® = 84 Ml/(56 amps)/28 days            Kitabis™ Pak = 280Ml (56 amps)/28 days            Tobi Podhaler® = 224 capsule/28 day            Tobi® inhalation neb = 280Ml (56 amps)/28 days            tobramycin = 280Ml (56 amps)/28 days</p>
<b>Antifungals, Oral</b>		
<p>fluconazole tab/susp            griseofulvin susp            nystatin tab/susp            terbinafine tab</p>	<p>Ancobon®            clotrimazole (mucous mem)            Cresemba®            Diflucan® tab/susp            flucytosine            Gris-Peg®            griseofulvin tab            griseofulvin ultramicrosize            itraconazole            itraconazole solution (generic for Sporanox® soln)            ketoconazole            Lamisil® tab/granules            Noxafil®            *Onmel®            Posaconazole tab (generic for Noxafil)            *Sporanox® cap/soln            Tolsura™            Vfend® tab/susp            voriconazole tab &amp; powder for susp</p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> Duration of the prescription (up to 12 months)</p> <p><b>Routine PDL edits plus</b></p> <p>* <b>Clinical Criteria for all Non-Preferred oral Antifungals. Requires the submission of a Clinical SA. Refer to <a href="#">Antifungal Oral SA Form</a></b></p>





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<b>Cephalosporins, Oral</b>			
<b>Second Generation Cephalosporins</b>			<b>LENGTH OF AUTHORIZATIONS:</b> Date of service only; no refills.
cefaclor capsule cefprozil tab/susp cefuroxime tab	cefaclor ER cefaclor susp Ceftin® tab/susp		<b>Routine PDL edits plus</b>
<b>Third Generation Cephalosporins</b>			<b>Clinical Criteria for Non-Preferred Cephalosporins</b>
cefdinir cap/susp	Cedax® cap/susp ceftibuten cefditoren pivoxil cefixime suspension cefpodoxime proxetil cap/susp Spectracef® Suprax® chewable tab/cap/susp		<ul style="list-style-type: none"> <li>• Infection caused by an organism resistant to preferred drugs, OR</li> <li>• A therapeutic failure to no less than a three-day trial of <b>one preferred cephalosporin</b>; OR</li> <li>• The member is completing a course of therapy with a non-preferred drug initiated in the hospital.</li> </ul>
<b>Macrolides, Oral</b>			
<b>Macrolides &amp; Ketolides</b>			<b>LENGTH OF AUTHORIZATIONS:</b> Date of service only; no refills
azithromycin pack/susp/tab clarithromycin tab/susp E.E.S.® 200 susp erythromycin base cap DR erythromycin ethylsuccinate 200mg susp erythromycin stearate	Biaxin® tab clarithromycin ER Eryped® 200 susp Eryped® 400 susp Ery-tab® E.E.S.® 400 tab Erythrocin® Stearate erythromycin base tab erythromycin ethylsuccinate 400mg tab(Generic E.E.S.® 400) *Ketek® PCE® Zithromax® pac/tab/susp ZMAX® susp		<b>Routine PDL edits plus</b>  <b>Clinical Criteria for Non-Preferred Macrolides and Ketolides</b> <ul style="list-style-type: none"> <li>• Infection caused by an organism resistant to preferred drugs; OR</li> <li>• A therapeutic failure to no less than a <b>three-day trial of one preferred drug within the same class</b>; OR</li> <li>• The member is completing a course of therapy with a non-preferred drug which was initiated in the hospital.</li> </ul> <b>* Ketek® Clinical Criteria</b> <ul style="list-style-type: none"> <li>• Treatment of community-acquired pneumonia (of mild to moderate severity) AND</li> <li>• Infection is caused by one of the following microorganism: <i>Streptococcus pneumoniae</i>, <i>Haemophilus influenzae</i>, <i>Moraxella catarrhalis</i>, <i>Chlamydia pneumoniae</i>, or <i>Mycoplasma pneumoniae</i>. AND</li> <li>• A therapeutic failure to no less than a <b>three-day trial of one preferred drug within the same class</b>; OR</li> <li>• The member is completing a course of therapy with a non-preferred drug initiated in the hospital.</li> </ul>



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Preferred Agents		Non-Preferred Agents		SA Criteria
<b>Otic</b>				
Ciprodex® ofloxacin neomycin/polymyxin/hc soln/ sus	Cetraxal® Cipro HC® ciprofloxacin/dexamethasone (generic Ciprodex) ciprofloxacin and fluocinolone acetonide (generic Otovel) Otovel	<b>LENGTH OF AUTHORIZATIONS:</b> Date of service only; no refills  <b>Routine PDL edits</b>		
<b>Quinolones, Oral</b>				
<b>Second Generation Quinolones</b>		<b>LENGTH OF AUTHORIZATIONS:</b> Date of service only; no refills		
ciprofloxacin susp/tab	Cipro® IR & XR & susp ciprofloxacin ER Noroxin® ofloxacin	<b>Routine PDL edits plus:</b>  <b>Clinical Criteria for Non-Preferred Quinolones</b> <ul style="list-style-type: none"> <li>Infection caused by an organism resistant to preferred drugs; <b>OR</b></li> <li>A therapeutic failure to no less than a <b>three-day trial of one preferred quinolone</b>; <b>OR</b></li> <li>The member is completing a course of therapy with a non-preferred drug initiated in the hospital.</li> </ul>		
<b>Third Generation Quinolones</b>				
levofloxacin tab	Baxdela™ tab Levaquin® tab/susp levofloxacin susp moxifloxacin			
<b>Topical Antibiotics</b>				
mupirocin ointment	*Altabax™ (QL) Bactroban® cr/ointment Centany® Centany AT® Kit	<b>LENGTH OF AUTHORIZATIONS:</b> Date of service only; no refills  <b>Routine PDL edits</b>  *Quantity Limit = 15 grams per 34 days		
<b>Vaginal Antibiotics</b>				
Cleocin® Ovules Clindesse® cr metronidazole gel Nuversa™ Vandazole™ gel	Cleocin® cr clindamycin cr Metrogel®	<b>LENGTH OF AUTHORIZATIONS:</b> Date of Service  <b>Routine PDL edits</b>		



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<b>Antivirals</b>		
*Hepatitis C Agents		<b>CLOSED CLASS</b>
Interferon		<b>LENGTH OF AUTHORIZATIONS:</b> 8 weeks (initial approval)
Peg-Intron®	Pegasis® Proclick/syringe/kit/vial	*ALL Hepatitis C Drugs ( <b>Preferred and Non-Preferred</b> ) require the submission of a Clinical SA. Refer to <a href="#">Hepatitis C Antivirals Preferred SA Form</a> or <a href="#">Hepatitis C Antivirals Non-Preferred SA Form</a>
Protease Inhibitor		
	Olysio™ (discontinued)	
*Nucleotide Analog NS5A & NS5B Polymerase Inhibitors & Combinations		
sofosbuvir /velpatasvir (generic Epclusa®)	Epclusa® Sovaldi® Vosevi™	
*NS5A, NS3/4A Inhibitor Combinations		
Mavyret™	Technivie™ Viekira Pak™ Viekira XR™ Zepatier®	
*NS5B & Protease Inhibitor combinations		
	Harvoni® Ledipasvir/Sofosbuvir (generic Harvoni®)	
<b>Herpes Oral</b>		
acyclovir cap/tab/susp famciclovir valacyclovir	Famvir® Sitavig® buccal tab Valtrex® Zovirax® tab/susp	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  Routine PDL edits
<b>Herpes Topical</b>		
Abreva OTC®	acyclovir cr (generic Zovirax cr)	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
	<b>docosanol</b> Zovirax® cr	acyclovir oint Denavir® Xerese® cr Zovirax® oint	Routine PDL edits
<b>Influenza</b>			
	oseltamivir susp/ cap	Flumadine® tab rimantadine Relenza Disk® Tamiflu® susp/cap Xofluza™	<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only Routine PDL edits
<b>Blood Modifiers</b>			
<b>Bile Salts</b>			
	ursodiol mg cap, tab	Actigal® Chenodal® Cholbam® Ocaliva® Urso® Urso® Forte tab	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits
<b>Phosphate Binders</b>			
	calcium acetate 667mg cap sevelamer carbonate tab	Auryxia™ calcium acetate 667mg tab Eliphos Fosrenol® chewable tab lanthanum carbonate chewable tab Phoslyra® powder, tab Renagel® Renvela® sevelamer carbonate powder packet sevelamer HCL (generic Renagel) Velphoro® chewable tab	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits
<b>Bone Resorption Suppression and Related Agents</b>			
<b>Bisphosphonates</b>			
	alendronate tab	Actonel®	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year



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ibandronate		alendronate soln Atelvia DR® Boniva® Binosto™ etidronate Fosamax® tab & Fosamax® plus D risedronate DR	Routine PDL edits
<b>Calcitonins</b>			
calcitonin-salmon nasal		Miacalcin®	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year Routine PDL edits
<b>Others</b>			
raloxifene		Evista® *Forteo® teriparatide *Tymlos™	<b>LENGTH OF AUTHORIZATIONS:</b> Initial approval will be for 1 year Routine PDL edits for Evista® *Clinical SA must be completed for <a href="#">(Forteo® OR Tymlos™ SA Form)</a>
<b>Cardiac</b>			
<b>Anticoagulants</b> <b>CLOSED CLASS</b>			
Low Molecular Weight Heparin includes FactorXA Inhibitor			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
enoxaparin		Arixtra® fondaparinux Fragmin® syringe & vial Lovenox®	Routine PDL edits plus
Oral Anticoagulants			<b>Clinical Criteria for Savaysa™</b>
Eliquis™ Eliquis™ Dose Pack Jantoven Pradaxa® Xarelto® Xarelto® Starter Pack warfarin		apixaban (generic Eliquis™) <del>Coumadin®</del> *Savaysa™	<ul style="list-style-type: none"> <li>• Diagnosis of:</li> <li>• Non-valvular Atrial Fibrillation, <b>OR</b></li> <li>• deep vein thrombosis, <b>OR</b></li> <li>• pulmonary embolism ; <b>AND</b></li> <li>• Documentation that CrCl is NOT ≥ 95mL/min calculated by Cockcroft-Gault equation</li> </ul> <p><b>Bristol-Myers Squibb announced it will discontinue sales of all strengths of Coumadin due to unexpected manufacturing issues that could not be resolved expeditiously. Supply is expected to be depleted by 8/31/2020.</b></p>



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Antihypertensive Agents</b>		
<b>ACE Inhibitors</b>		<b>LENGTH OF AUTHORIZATIONS: 1 year</b>
<b>benazepril</b> <b>enalapril</b> <b>lisinopril</b> <b>ramipril</b> <b>quinapril</b>	<i>Accupril<sup>®</sup></i> <i>Altace<sup>®</sup></i> <i>captopril</i> <i>Epaned<sup>™</sup> soln</i> <i>fosinopril</i> <i>Lotensin<sup>®</sup></i> <i>Mavik<sup>®</sup></i> <i>moexipril</i> <i>Monopril<sup>®</sup></i> <i>perindopril</i> <i>Qbrelis<sup>™</sup></i> <i>ramipril</i> <i>trandolapril</i> <i>Univasc<sup>®</sup></i> <i>Vasotec<sup>®</sup></i> <i>Zestril<sup>®</sup></i>	Routine PDL edits
<b>ACE Inhibitors + Calcium Channel Blocker Combinations</b>		
<b>amlodipine/benazepril</b>	<i>Lotrel<sup>®</sup></i> <i>Tarka<sup>®</sup></i> <i>trandolapril-verapamil ER</i>	
<b>ACE Inhibitors + Diuretic Combinations</b>		
<b>benazepril/HCTZ</b> <b>lisinopril/HCTZ</b> <b>enalapril/HCTZ</b>	<i>Accuretic<sup>®</sup></i> <i>captopril/HCTZ</i> <i>fosinopril/HCTZ</i> <i>Lotensin HCT<sup>®</sup></i> <i>moexipril/HCTZ</i> <i>quinapril/HCTZ</i> <i>Vaseretic<sup>®</sup></i> <i>Zestoretic<sup>®</sup></i>	



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<i>Preferred Agents</i>		<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
<b>Angiotensin Receptor Blockers</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year <b>Routine PDL edits plus</b>  <i>Quantity Limit = 2 per day for Entresto™</i>
*Entresto™ (QL) irbesartan losartan olmesartan valsartan	Atacand® Avapro® Benicar® candesartan Cozaar® Diovan® Edarbi® eprosartan mesylate Micardis® Teveten®		
<b>Angiotensin Receptor Blockers + Calcium Channel Blocker Combinations</b>			
amlodipine/olmesartan amlodipine/valsartan	Azor® amlodipine/olmesartan/HCTZ amlodipine/valsartan/HCTZ Exforge® & Exforge® HCT Tribenzor®		
<b>Angiotensin Receptor Blockers + Diuretic Combinations</b>			
irbesartan/HCTZ losartan/HCTZ olmesartan/HCTZ valsartan/HCTZ	Atacand HCT® Avalide® Benicar HCT® candesartan/HCTZ Diovan HCT® Edarbyclor® Hyzaar® Micardis HCT® telmisartan/HCTZ Teveten HCT®		
<b>Antihypertensives, Sympatholytics</b>		<b>CLOSED CLASS</b>	
Catapres®-TTS clonidine tab	Catapres® clonidine (transdermal)		



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<i>Preferred Agents</i>		<i>Non-Preferred Agents</i>		<i>SA Criteria</i>
<b>guanfacine</b> <b>methyldopa</b> <b>reserpine</b>		<i>Clorpres</i> <sup>®</sup> <i>methyldopa/HCTZ</i> <i>Tenex</i> <sup>®</sup>		<b>*Clinical Criteria for Hemangeol™</b> <ul style="list-style-type: none"> <li>Diagnosis treatment of proliferating infantile hemangioma requiring systemic therapy; AND</li> </ul>
<b>Beta Blockers</b>				
<b>atenolol</b> <b>bisoprolol</b> <b>carvedilol</b> <b>labetalol</b> <b>metoprolol tartrate</b> <b>metoprolol succinate</b> <b>propranolol tab &amp; ER/soln</b> <b>Sorine</b> <sup>®</sup> <b>sotalol AF</b> <b>sotalol HCL</b>		<i>acebutaolol</i> <i>Betapace</i> <sup>®</sup> IR & AF <sup>®</sup> <i>betaxolol</i> <i>Bystolic</i> <sup>®</sup> <i>Carvedilol ER</i> <i>Coreg</i> <sup>®</sup> IR & CR <sup>®</sup> <i>Corgard</i> <sup>®</sup> <i>*Hemangeol</i> <sup>™</sup> <i>Inderal</i> <sup>®</sup> XL <i>Innopran</i> <sup>®</sup> XL <i>Kaspargo</i> <sup>™</sup> Sprinkle <i>Levatol</i> <sup>®</sup> <i>Lopressor</i> <sup>®</sup> <i>nadolol</i> <i>pindolol</i> <i>propranolol LA</i> <i>Sectral</i> <sup>®</sup> <i>Sotylize</i> <sup>™</sup> <i>Tenormin</i> <sup>®</sup> <i>timolol maleate</i> <i>Toprol XL</i> <sup>®</sup> <i>Trandate</i> <sup>®</sup> <i>Zebeta</i> <sup>®</sup>		
<b>Beta Blockers + Diuretic Combinations</b>				
<b>atenolol/chlorthalidone</b> <b>bisoprolol/HCTZ</b>		<i>Corzide</i> <sup>®</sup> <i>Dutoprol</i> <sup>®</sup> <i>Lopressor HCT</i> <sup>®</sup> <i>metoprolol/HCTZ</i> <i>nadolol/bendroflumethiazide</i> <i>propranolol/HCTZ</i> <i>Tenoretic</i> <sup>®</sup>		





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Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Ziac</i> <sup>®</sup>	
<b>Calcium Channel Blockers –Dihydropyridine</b>		
<b>Afeditab CR</b> <sup>®</sup> <b>amlodipine</b> <b>Nifedical XL</b> <sup>®</sup> <b>nifedipine</b> <b>nifedipine ER</b>	<i>Adalat CC</i> <sup>®</sup> <i>Consensi</i> <sup>®</sup> (amlodipine/celecoxib) <i>felodipine ER</i> <i>isradipine</i> <i>Katerzia</i> <sup>™</sup> oral suspension <i>nisoldipine</i> <i>nicardipine</i> <i>Norvasc</i> <sup>®</sup> <i>Procardia</i> <sup>®</sup> <i>Procardia XL</i> <sup>®</sup> <i>Sular</i> <sup>®</sup>	
<b>Calcium Channel Blockers- Non-Dihydropyridine</b>		
<b>Cartia XT</b> <sup>®</sup> <b>diltiazem IR, ER q12hr &amp; 24hr</b> <b>Taztia XT</b> <sup>®</sup> <b>verapamil tab IR &amp; ER</b>	<i>Calan</i> <sup>®</sup> IR & SR <i>Cardizem</i> <sup>®</sup> IR, CD & LA <i>Isoptin SR</i> <sup>®</sup> <i>diltiazem LA</i> <i>Matzim LA</i> <i>Tiazac</i> <sup>®</sup> <i>verapamil 360 cap</i> <i>verapamil ER cap</i> <i>Verelan</i> <sup>®</sup> & <i>Verelan PM</i> <sup>®</sup>	
<b>Direct Renin Inhibitors (includes combination)</b>		
	<i>aliskiren 150 &amp; 300mg (generic for Tekturna)</i> <i>Tekamlo</i> <sup>®</sup> <i>Tekturna</i> <sup>®</sup> <i>Tekturna HCT</i> <sup>®</sup> <i>Twynsta</i> <sup>®</sup> <i>telmisartan/amlodipine</i>	



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Lipotropics</b>		
<b>Bile Acid Sequestrants</b>		<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<b>cholestyramine powder reg &amp; light</b> <b>colestipol tab</b> <b>Prevalite®</b>	<i>Colestid® granule/packet/tab</i> <i>colesevelam tab and Pkt (generic Welchol)</i> <i>colestipol HCl granules</i> <i>Questran® powder/powder Light</i> <i>Welchol® pack, tab</i>	<b>Routine PDL edits plus</b>
<b>Cholesterol Absorption Inhibitor (CAI) and /or ACL Inhibitor (adenosine triphosphate citrate lyase)</b>		
<b>ezetimibe</b>	<i>Nexletol® (bempedoic acid)</i> <i>Nexlizet® (bempedoic acid/ezetimibe)</i> <i>Zetia®</i>	<b><u>Routine PDL and Clinical criteria Nexletol™ or Nexlizet™</u></b> <b><u>Initial Approval Criteria</u></b> <ul style="list-style-type: none"> <li>• Patient is ≥ 18 years of age; AND</li> <li>• Patient has diagnosis of heterozygous familial hypercholesterolemia (HeFH) or established atherosclerotic cardiovascular disease (ASCVD); AND</li> <li>• Patient has failed to achieve a target LDL-C despite physician attestation that the patient is adherent to maximally tolerated doses of statins prior to the lipid panel demonstrating suboptimal reduction; AND</li> <li>• Patient can be classified into ONE of the following risk factor groups:               <ul style="list-style-type: none"> <li>○ Extremely high risk ASCVD: (defined as extensive or active burden of ASCVD, or ASCVD with extremely high burden of adverse or poorly controlled risk cardio-metabolic risk factors including HeFH or severe hypercholesterolemia [SH] LDL-C &gt; 220 mg/dl) with an LDL-C ≥ 70 mg/dL; OR</li> <li>○ Very high risk ASCVD: (defined as less extensive ASCVD and poorly controlled cardiometabolic risk factors) with an LDL-C ≥ 100 mg/dL; OR</li> <li>○ High risk ASCVD: (defined as either less extensive ASCVD and well-controlled risk factors or primary prevention HeFH or SH &gt;220 mg/dl with poorly controlled risk factors) with LDL-C ≥ 130 mg/dL; AND</li> </ul> </li> </ul>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
			<ul style="list-style-type: none"> <li>• Therapy will be used in conjunction with maximally-tolerated doses of a statin; AND</li> <li>• Therapy will not be used with concurrent doses of simvastatin &gt; 20 mg or pravastatin &gt; 40mg;</li> </ul> <p><b>Renewal Criteria</b></p> <ul style="list-style-type: none"> <li>• Laboratory analyses demonstrate a reduction in LDL-C when compared to the baseline values (prior to initiating bempedoic acid or bempedoic acid/ezetimibe); AND</li> </ul> <p>Patient has shown continued adherence to maximally tolerated statin dosage</p>
	<b>Fibric Acid Derivatives</b>		
	<b>fenofibrate (generic Tricor® 48mg 145mg) gemfibrozil</b>	<i>Antara® fenofibrate (generics for Antara®, Fenoglide® &amp; Lipofen®) fenofibrate (generics for Triglide®) fenofibric acid Fenoglide® Fibricor® Lipofen® Lofibra® Lopid® Tricor® Triglide® Trilipix™</i>	
	<b>HMG CoA Reductase Inhibitors and Combo (High Potency Statins)</b>		
	<b>atorvastatin rosuvastatin simvastatin</b>	<i>amlodipine/atorvastatin Caduet® Crestor® Ezallor Sprinkle (rosuvastatin) Lipitor® Liptruzet®</i>	



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		<i>Livalo</i> <sup>®</sup> <i>simvastatin sol (generic for Flolipid sol)</i> <i>simvastatin/ezetimibe</i> <i>Vytorin</i> <sup>®</sup> <i>Zocor</i> <sup>®</sup> <i>Zypitamag</i> <sup>™</sup>	
<b>HMG CoA Reductase Inhibitors and Combinations (Statins)</b>			
lovastatin pravastatin		<i>Advicor</i> <sup>®</sup> <i>Altoprev</i> <sup>®</sup> <i>fluvastatin</i> <i>Lescol</i> <sup>®</sup> and <i>Lescol XL</i> <sup>®</sup> <i>Mevacor</i> <sup>®</sup> <i>Pravachol</i> <sup>®</sup>	
<b>Microsomal Triglyceride Transfer Protein Inhibitor</b>			
		<i>*Juxtapid</i> <sup>™</sup>	<b>*Clinical Criteria for Juxtapid<sup>™</sup>. Refer to <a href="#">Juxtapid<sup>™</sup> SA Fax Form</a></b>
<b>Niacin Derivatives</b>			
niacin ER		<i>Niaspan</i> <sup>®</sup> <i>Niacor</i> <sup>®</sup>	
<b>Omega 3 Fatty Acid Agent</b>			
*** <i>omega-3 acid ethyl esters (ST)</i> Omega-3 OTC		<i>icosapent ethyl (generic Vascepa)</i> <sup>®</sup> *** <i>Lovaza</i> <sup>®</sup> (ST) <i>Vascepa</i> <sup>®</sup> (ST)	*** <b>Clinical Criteria for Lovaza<sup>®</sup> and omega-3 acid ethyl esters</b> <ul style="list-style-type: none"> <li>• Step edit requires trial and failure of any other lipotropic; <b>OR</b></li> <li>• Documented high triglycerides of ≥ 500 mg/dL.</li> </ul>
<b>*Proprotein Convertase Subtilisin Kexin Type 9 (PCSK9) Inhibitors</b>			
		<i>Praluent</i> <sup>®</sup> <i>Repatha</i> <sup>®</sup>	<b>LENGTH OF AUTHORIZATIONS:</b> Three months for initial approval; six months for renewal  <b>*ALL PCSK9 Inhibitors require the submission of a Clinical SA. Refer to <a href="#">PCSK9 or M4V SA Form</a></b>
<b>Platelet Inhibitors</b>			
Brilinta <sup>®</sup> clopidogrel dipyridamole		<i>*Aggrenox</i> <sup>®</sup> <i>*ASA/dipyridamole</i> ** <i>ASA/omeprazole (generic Yosprala)</i> <sup>®</sup>	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL edits plus</b>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
	<b>prasugrel (generic Effient®)</b>	** <i>Durlaza ER™</i> <i>Effient®</i> <i>Persantine®</i> <i>Plavix®</i> ** <i>Yosprala® Tab</i> *** <i>Zontivity™</i>	<b><u>Clinical Criteria for Select <span style="color:red">Non-Preferred Platelet Inhibitors</span></u></b> <b>*<u>Aggrenox® &amp; ASA/dipyridamole</u></b> <ul style="list-style-type: none"> <li>Aspirin and dipyridamole are covered as separate drugs without SA; clinical reason as to why the individual drugs cannot be used separately.</li> </ul> <b>**<u>Durlaza ER™ &amp; *Yosprala® Tab</u></b> <ul style="list-style-type: none"> <li>Aspirin is covered without SA; clinical reason as to why aspirin cannot be used.</li> </ul> <b>***<u>Zontivity™</u></b> <ul style="list-style-type: none"> <li>Diagnosis of MI (myocardial infarction) or PAD (peripheral arterial disease); <b>AND</b></li> <li>Members must not have a history of stroke, TIA, ICH, GI bleed and peptic ulcer; <b>AND</b></li> <li>Must have concomitant therapy with clopidogrel, unless member has a contraindication to clopidogrel in which case member must have concomitant therapy with aspirin; <b>AND</b></li> <li>Member is 18 years of age or older; <b>AND</b></li> <li>Prescribed by or in consultation with a cardiologist.</li> </ul>
<b>*Pulmonary Arterial Hypertension Agents</b>			
<b>Inhaled Prostaglandin Analogues</b>		<b>LENGTH OF AUTHORIZATIONS: 1 year</b>	
<b>Ventavis®</b>	<i>Tyvaso®</i>	<b>Routine PDL edits plus</b>	
<b>Oral Endothelin Receptor Antagonist</b>			
<b>ambrisentan (generic Letairis) 5 &amp; 10mg</b> <b>Tracleer® tab</b>	<i>bosentan (generic Tracleer®)</i> <i>Letairis® 5 &amp; 10 mg</i> <i>Opsumit®</i> <i>Tracleer® susp</i>		
<b>*Phosphodiesterase 5 Inhibitors (PDE-5)</b>		<b>*<u>Clinical Criteria for all <span style="color:red">preferred and non-preferred PDE-5</span></u></b>	
<b>Alyq (tadalafil)</b> <b>sildenafil tab/susp</b> <b>tadalafil</b>	<i>Adcirca®</i> <i>Revatio® tab/inj/susp</i>	<ul style="list-style-type: none"> <li>Diagnosis of pulmonary hypertension in members &gt;18 years is required; <b>AND</b></li> <li>The prescriber must be a pulmonary specialist or cardiologist; <b>AND</b></li> <li>Must have a rationale for not taking the sildenafil tablet to receive a SA for injectable Revatio®</li> </ul>	



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<b>Prostacyclin Vasodilator and Receptor Agonist</b>				
		<i>Orenitram™</i> <i>Upravi®</i>		
<b>Soluble Guanylate Cyclase Stimulators</b>				
		<i>Adempas®</i>		
<b>Central Nervous System</b>				
<b>Alzheimer's Agents</b>				
<b>Cholinesterase Inhibitors</b>				<b>LENGTH OF AUTHORIZATIONS:</b> Length of prescription (up to 3 months)
<b>donepezil OTD &amp; tab</b> <b>Exelon® (transderm)</b>		<i>Aricept® ODT, tab</i> <i>Exelon® cap</i> <i>galantamine IR, ER tab/soln</i> <i>Memantine ER (generic Namenda XR)</i> <i>Namzaric® (donepezil/memantine)</i> <i>Razadyne® IR, ER</i> <b>rivastigmine cap &amp; patch</b>		<b>Routine PDL edits</b>
<b>NMDA Receptor Antagonist</b>				
<b>memantine tab</b>		<i>memantine Dose Pack</i> <i>memantine soln</i> <i>Namenda® Dose Pack/XR tab</i> <i>Namenda® tab</i>		
<b>Anticonvulsants</b>				
<b>Barbiturates</b>				<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<b>phenobarbital elixir/tab</b> <b>primidone</b>		<i>Mysoline®</i>		<b>Routine PDL edits plus</b>
<b>Benzodiazepines</b>				<b>*Clinical Criteria for Onfi®</b>
<b>*clobazam tab (generic</b> <b>Onfi®tab)</b> <b>clonazepam tab</b>		<i>clonazepam ODT</i> <i>Diastat® rectal</i> <i>Diastat® AcuDial™ rectal</i> <b>**Nayzilam®</b>		<ul style="list-style-type: none"> <li>• Patient is at least two years of age or older; <b>AND</b></li> <li>• Patient must have a diagnosis of seizures associated with Lennox-Gastaut syndrome (LGS) <b>AND</b></li> <li>• Using as adjunctive therapy with other anticonvulsants</li> </ul>



# Virginia Medicaid's Preferred Drug List (PDL)/ Common Core Formulary

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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
	<p><b>diazepam rectal &amp; Device rectal</b>  <b>***Valtoco® Nasal</b></p>	<p><i>*Onfi® susp/tab</i>  <i>Sympazan™ (clobazam)</i></p>	<p><b>** Clinical Criteria for Nayzilam®</b></p> <ul style="list-style-type: none"> <li>• <b>Patient is at least 12 years of age or older; AND</b></li> </ul> <p>Diagnosis of acute treatment of intermittent, stereotypic episodes of frequent seizure activity that are distinct from a patient's usual seizure pattern in patients with epilepsy</p> <p><b>***Clinical Criteria for Valtoco® Nasal (this is reviewed electronically, AutoPA)</b>            Duration of Approval: 1 year</p> <ul style="list-style-type: none"> <li>• <b>Patient must be 6 years of age or older; AND</b></li> <li>• <b>Patient has been diagnosed with</b> <ul style="list-style-type: none"> <li>○ <b>intermittent, stereotypic episodes of frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) that are distinct from a patient's usual seizure pattern</b></li> </ul> </li> </ul>
	<b>Cannabidiol</b>		
	<p><b>*Epidiolex® (cannabidiol)</b></p>		<p><b>*Clinical Criteria for Epidiolex®</b> (this is reviewed electronically, AutoPA)            Duration of Approval: 1 year</p> <ul style="list-style-type: none"> <li>• Patient must be ≥ 1 years of age; AND</li> <li>• Patient has been diagnosed with               <ul style="list-style-type: none"> <li>○ Epilepsy and recurrent seizures including Lennox-Gastaut syndrome, Dravet syndrome, or tuberous sclerosis complex</li> </ul> </li> </ul>
	<b>Carbamazepine Derivatives</b>		
	<p><b>carbamazepine chewable tab/susp/tab</b>  <b>carbamazepine ER</b>  <b>carbamazepine XR</b>  <b>oxcarbazepine susp &amp; tab</b></p>	<p><i>Aptiom®</i>  <i>Carbatrol®</i>  <i>Equetro® cap</i>  <i>Oxtellar™ XR</i>  <i>Tegreto® susp/tab</i>  <i>Tegreto® XR</i>  <i>Trileptal® susp/tab</i>  <i>vigabatrin powder pack</i></p>	
	<b>Hydantoins</b>		
	<p><b>phenytoin cap/chew tab/ susp</b>  <b>phenytoin ext cap</b></p>	<p><i>Dilantin® cap</i>  <i>Dilantin® Infatab, susp</i></p>	

Preferred Agents	Non-Preferred Agents	SA Criteria
	Peganone® Phenytek®	
<b>Succinimides</b>		
ethosuximide cap/syrup	Celontin® Zarontin® cap/syrup	
<b>Valproic Acid and Derivatives</b>		
divalproex tab/sprinkle divalproex ER valproic acid cap, sol	Depakene® cap/syrup Depakote® ER & sprinkle	
<b>Other Anticonvulsants</b>		
<b>Gabitril®</b> lamotrigine tab lamotrigine chew tab lamotrigine XR levetiracetam soln/tab levetiracetam ER roweepra (generic version levetiracetam) Vimpat® soln/tab topiramate tab/sprinkle cap zonisamide cap	Banzel® susp/tab Briviact® Diacomit® felbamate susp/tab Felbatol® susp/tab * Fintepla® Fycompa® susp/tab Keppra® soln/tab Keppra® XR Lamictal® XR Lamictal® ODT/ODT dose pk Lamictal® tab/dose pk Lamictal® XR dose pk lamotrigine tab dose pk & ODT Potiga® rufinamide(generic Banzel® susp) Qudexy™ XR Sabril® powder pack/tab <b>tiagabine</b> Topamax® tab/sprinkle Trokendi™ XR vigabatrin (generic Sabril® tab) Xcopri® Zonegran®	<b>*Clinical Criteria for Fintepla®</b> <ul style="list-style-type: none"> <li>• Patient is at least two years of age or older; AND</li> <li>• Patient must have a diagnosis of Dravet syndrome</li> </ul>





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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Antidepressants</b>		
<b>Other</b>		<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
bupropion IR, SR & XL desvenlafaxine ER tab (generic for Pristiq®) mirtazapine ODT/tab trazodone tab venlafaxine IR tab & ER cap	Aplenzin® Brintellix® bupropion XL (generic Forfivo® XL) desvenlafaxine ER tablet (generic for Khedezla™) Effexor® XR Emsam® transdermal Fetzima® Forfivo® XL Khedezla™ Marplan® Nardil® nefazodone Oleptro® ER Parnate® phenelzine Pristiq® Remeron® ODT/tab tranylcypromine sulfate Trintellix venlafaxine ER tab Viibryd® tab/dose pk Wellbutrin® IR, SR & XL	Routine PDL edits
<b>SSRI</b>		
citalopram soln/tab escitalopram tab fluoxetine cap/soln fluvoxamine tab paroxetine tab sertraline conc, sol, tab	Brisdelle® Celexa® tab escitalopram soln fluoxetine DR cap/tab fluvoxamine ER Lexapro® tab Luvox® CR paroxetine CR Paxil® tab/susp & Paxil® CR	



# Virginia Medicaid's Preferred Drug List (PDL)/ Common Core Formulary

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Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Pexeva<sup>®</sup></i> <i>Prozac<sup>®</sup> cap/weekly</i> <i>Sarafem<sup>®</sup></i> <i>Zoloft<sup>®</sup> conc/tab</i>	
<b>Antimigraine Agents</b>		
sumatriptan succinate tab cartridge/vial/pen <b>Imitrex<sup>®</sup> nasal</b> rizatriptan tab/MLT	<i>almotriptan</i> <i>Alsuma<sup>®</sup></i> <i>Amerge<sup>®</sup></i> <i>Axert<sup>®</sup></i> <i>Cambia<sup>®</sup></i> <i>eletriptan (generic Relpax<sup>®</sup>)</i> <i>Frova<sup>®</sup></i> <i>frovatriptan (generic Frova<sup>®</sup>)</i> <i>Imitrex<sup>®</sup> cartridge /pen/tab/vial</i> <i>Maxalt<sup>®</sup> tab &amp; MLT</i> <i>Migranow<sup>TM</sup> Kit</i> <i>naratriptan</i> <i>Onzetra<sup>TM</sup> Xsail<sup>TM</sup></i> <i>Relpax<sup>®</sup></i> <i>sumatriptan KITS</i> <i>Sumavel<sup>®</sup> Dosepro</i> <b>sumatriptan nasal</b> <i>sumatriptan/naproxen (generic</i> <i>Treximet<sup>®</sup>)</i> <i>Tosymra</i> <i>Treximet<sup>®</sup></i> <i>Zembrace<sup>TM</sup> SymTouch<sup>TM</sup></i> <i>Zomig<sup>®</sup> tab/nasal spray/ZMT</i>	<b><u>LENGTH OF AUTHORIZATIONS:</u> 1 year</b>  Routine PDL edits
<b>Antimigraine Agents, Others</b>		
<b>Calcitonin Gene-related Peptide Antagonist (CGRP)</b>		
Emgality <sup>TM</sup> Syringe Emgality <sup>TM</sup> Pen <b>Ajovy<sup>®</sup></b> <b>Ajovy<sup>®</sup> Autoinjector</b>	<i>Aimovig<sup>TM</sup></i> <i>Emgality<sup>TM</sup> 100MG Syringe</i>	*All Nonpreferred CGRPs require the submission of a Clinical SA. Refer to fax form <a href="#">Antimigraine Agents, Others SA Form</a>



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Ajovy Autoinjector 3-Pk</b>		<b>Ajovy®, Emgality™, Ubrovelvy® have an AutoPA and require a trial of 2 generic triptans, no clinical SA</b>
<b>Acute treatment of migraine</b>		
Ubrovelvy®	Nurtec™ ODT Reyvow®	<b>Nurtec™ ODT has a quantity limit of 8 units per month</b>
<b>*Antipsychotics (AG)</b>		
<b>Atypical</b>		
aripiprazole tab clozapine tab Latuda® olanzapine ODT, tab, IM quetiapine tab risperidone ODT/soln/tab ziprasidone cap	Abilify® tab/IM inj ***Abilify Mycite®(with sensor) **aripiprazole ODT, soln <b>asenapine (generic for Saphris®)</b> Caplyta Capsule™ Clozaril® clozapine ODT Fanapt® tab & titration pk Fazaclo® **Geodon® tab, IM Invega® **Nuplazid™ tab, cap (QL)(AG) **olanzapine/fluoxetine paliperidone ER Rexulti® tab Risperdal® ODT/soln/tab Saphris® Secuado® Patch Seroquel® IR Seroquel® XR Versacloz™ Vraylar™ <b>ziprasidone IM (generic for Geodon®)</b> Zyprexa® tab/IM/Zydis	<p><b><u>LENGTH OF AUTHORIZATIONS:</u> 1 year or 6 months for members &lt; 18 yrs</b></p> <p><b>Routine PDL edits plus</b></p> <p><b>*ALL antipsychotics for children 0 to 17 years of age (preferred and non-preferred) require the submission of a Clinical SA. Refer to <a href="#">(Antipsychotics In Children Less Than 18 Years SA Form)</a></b></p> <p><b>**Clinical Criteria Nuplazid™</b></p> <ul style="list-style-type: none"> <li>Indicated for the treatment of hallucinations and delusions associated with Parkinson's disease psychosis.</li> </ul> <p><i>Quantity Limit Nuplazid™ = 2 per day</i></p> <p><b>**Clinical Criteria for Abilify Mycite®</b></p> <p><b>Initial Approval Criteria: For Three months SA</b></p> <p>Patient must:</p> <ul style="list-style-type: none"> <li>Be ≥ 18 years of age; AND</li> <li>Have tolerability to oral aripiprazole with suboptimal effects (as assessed by prescriber) that may be due to adherence problems; AND</li> <li>Have a smart phone compatible with the device; AND</li> <li>Give consent to a healthcare provider and caregiver (if applicable) to monitor the portal; AND</li> <li>There is a documented intervention by prescriber if nonadherence is detected</li> </ul> <p><b>Renewal Criteria: Every 3 Months Reevaluate</b></p> <ul style="list-style-type: none"> <li>Patient must:</li> </ul>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
			<ul style="list-style-type: none"> <li>• Continue to meet initial criteria; AND</li> <li>• Have prescriber attestation that patient benefited from therapy; AND</li> <li>• Have prescriber attestation that there is a continued need for device (e.g., continued suboptimal effects and/or compliance); AND</li> <li>• Have a healthcare provider and caregiver (if applicable) agree to continue to monitor device; AND</li> <li>• Not have worsened target symptoms; AND</li> <li>• Not have had any treatment-limited adverse effects (e.g.,</li> <li>• Not have had any treatment-limited adverse effects (e.g., hypersensitivity, suicidality, neuroleptic malignant syndrome, tardive dyskinesia, metabolic changes, pathological gambling and other compulsive behaviors, orthostatic hypotension, falls, seizures, cognitive and motor impairment, dysphagia, disruption in body temperature regulation, and leukopenia, neutropenia, and agranulocytosis); AND</li> <li>• Have a healthcare provider state reason why the patient cannot use long acting injectable atypical antipsychotic if there is continued nonadherence.</li> </ul>
	<b>Atypical, Long Acting Injectable</b>	<b>CLOSED CLASS</b>	<b>LENGTH OF AUTHORIZATIONS: 1 year</b>
	Abilify Maintena® Aristada® Aristada® Initio Risperdal Consta® Invega Sustenna® & Trinza®	Perseris™ (risperidone) Zyprexa® Relprev™	<b>Routine PDL edits</b>
	<b>Typical</b>		<b>LENGTH OF AUTHORIZATIONS: 1 year</b>
	amitriptyline/perphenazine chlorpromazine fluphenazine elixir/soln/tab fluphenazine decantate haloperidol decantate haloperidol lactate conc haloperidol tab loxapine perphenazine trifluoperazine thiothixene	Haldol decanoate (injection) pimozide Moban® molindone	<b>Routine PDL edits</b>



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>thioridazine</b>		
<b>Neuropathic Pain</b>		
capsaicin OTC topical duloxetine 20, 30 & 60 mg (generic for Cymbalta®) gabapentin cap/tab/soln lidocaine 5% patch pregabalin cap	Cymbalta® Dermacinrx® PHN Pak™ Kit Drizalma™ Sprinkle duloxetine 40 mg (generic for Irenka™) Gralise™ (gabapentin, extend release) Horizant™ (gabapentin enacarbil, extended release) Irenka™ Lidoderm® patch LidoPure Patch Lyrica CR Lyrica® soln Lyrica® Neurontin® cap/tab/soln pregabalin sol Qutenza Kit® (Topical) (capsaicin) Savella™ & Savella™ Dose Pak Zilacaine Patch Ztlido™ 1.8% (lidocaine topical system)	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL</b>  Plus <b>Criteria for Lyrica solution:</b> – If diagnosis is epilepsy/seizures, patient must have a problem swallowing tablets/capsules AND clinical reason why at least TWO preferred seizure medications cannot be used. – For other diagnoses, patient must have a problem swallowing tablets/capsules AND routine PDL • If Lyrica CR is requested, routine PDL applies. Note – This is NOT indicated for epilepsy/seizures.
<b>Non-Ergot Dopamine Receptor Agonist</b>		
pramipexole ropinirole HCl	Mirapex® IR & ER Neupro® pramipexole ER Requip® XR ropinirole HCl ER	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL edits</b>
<b>Sedatives / Hypnotics</b>		
temazepam 15 & 30 mg	estazolam flurazepam Halcion® Restoril®	<b>LENGTH OF AUTHORIZATIONS:</b> Length of the prescription (up to 3 months)  <b>Routine PDL edits</b>



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Preferred Agents	Non-Preferred Agents	SA Criteria
	temazepam 7.5 mg & 22.5 mg triazolam	
<b>Sedatives / Hypnotics (Non-Benzodiazepine)</b>		
eszopiclone zaleplon zolpidem	Ambien® IR & CR Belsomra® Dayvigo™ doxepin (generic for Silenor®) Edluar™ *Hetlioz™ Intermezzo® Lunesta® Rozerem® Silenor® Sonata® zolpidem CR Zolpimist™ spray zolpidem (generic Intermezzo®)	<p><b>LENGTH OF AUTHORIZATIONS:</b> 6 months. <b>For Renewal</b> - must document therapeutic benefit and confirm compliance</p> <p>Routine PDL edits plus</p> <p><b>*Clinical Criteria for Hetlioz™</b></p> <ul style="list-style-type: none"> <li>• For the treatment of Non-24-Hour Sleep-Wake Disorder (Non-24), <b>AND</b></li> <li>• Member must be age 18 years of age or older.</li> <li>• Quantity limit = 1 tablet per day.</li> </ul>
<b>Skeletal Muscle Relaxants</b>		
baclofen chlorzoxazone cyclobenzaprine HCL dantrolene sodium methocarbamol tizanidine tab	Amrix® *carisoprodol *carisoprodol/ASA *carisoprodol/ASA/codeine cyclobenzaprine ER Dantrium® Fexmid® Lorzone® metaxalone orphenadrine citrate orphenadrine/ASA/caffeine Parafon Forte® DSC Robaxin® Skelaxin® *Soma® tizanidine cap Zanaflex®	<p><b>LENGTH OF AUTHORIZATIONS:</b></p> <ul style="list-style-type: none"> <li>• 1 year for chronic conditions</li> <li>• Duration of prescription (up to 3 months) for acute conditions</li> <li>• One month per every 6 months for carisoprodol drugs</li> </ul> <p>Routine PDL edits plus</p> <p><b>*Clinical Criteria for Carisoprodol Drugs. Refer to <a href="#">Soma/carisoprodol SA Fax Form</a></b></p>



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>Smoking Cessation</b>		
bupropion SR Chantix® Chantix® DS PK nicotine gum/lozenge/patch	Nicoderm CQ® Patch Nicorette® Gum/Lozenges Nicotrol® Inhaler & NS Zyban®	<b>LENGTH OF AUTHORIZATIONS:</b> 6 months  Routine PDL edits
<b>*Stimulants/ADHD Medications (AG) <span style="color: red;">CLOSED CLASS</span></b>		
<b>Amphetamine Drugs</b>		
<b>Adderall®XR</b> amphetamine salts combo <i>(generic for Adderall IR)</i> dextroamphetamine <i>(generic for Dexedrine)</i> Vyvanse® cap/chewable tab <i>(lisdexamfetamine)</i>	Adderall® IR <i>(amphetamine salts combo)</i> Adzenys XR ODT™ Adzenys ER™ susp Adzenys® ER <b>amphetamine salts combo XR</b> amphetamine sulfate <i>(generic Evekeo®)</i> amphetamine susp <i>(generic Adzenys ER™ susp)</i> Desoxyn® Dexedrine® dextroamphetamine SR & soln Dyanavel® XR susp Evekeo® Evekeo™ ODT methamphetamine Mydayis ER™ Procentra® soln Zenedi™	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  Routine PDL edits  *All stimulants ( <span style="color: red;">preferred and non-preferred</span> ) require the submission of Clinical SA if prescribed for a child less than four or an adult eighteen years and older. Refer to Stimulant SA form ( <a href="#">Stimulant/ADHD Medications SA Form</a> )  <b>This does not include nonstimulant agents such as atomoxetine (generic for Strattera®), clonidine ER, guanfacine ER or others</b>
<b>Methylphenidate Drugs</b>		
All methylphenidate IR generic <b>Concerta®</b> Daytrana® Transdermal Focalin® IR & XR	Adhansia™XR Aptensio™ XR Cotempla XR-ODT™ dexmethylphenidate IR & XR Jornay PM Metadate CD® Metadate ER®	



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Preferred Agents		Non-Preferred Agents	SA Criteria
		<p>Methylin ER<sup>®</sup>, soln IR  methylphenidate chew &amp; soln  methylphenidate ER, LA, SR  methylphenidate ER(generic  Relexxii<sup>®</sup>)  <b>methylphenidate ER(generic  Aptensio<sup>™</sup> XR)</b>  Ritalin<sup>®</sup> IR, LA<sup>®</sup> &amp; SR<sup>®</sup>  Relexxii<sup>®</sup>  QuilliChew ER<sup>™</sup>  Quillivant<sup>™</sup> XR susp</p>	
<b>Miscellaneous Drugs</b>			
<p>atomoxetine (generic for  Strattera<sup>®</sup>)  clonidine ER  guanfacine ER</p>	<p>***<i>armodafinil</i> (generic Nuvigil<sup>™</sup>)  ***<i>modafinil</i>  ***<i>Nuvigil<sup>™</sup></i> (AG)  ***<i>Provigil<sup>®</sup></i> (AG)  Intuniv<sup>®</sup>  Sunosi<sup>™</sup>  Strattera<sup>®</sup>  Wakix<sup>®</sup></p>	<p>***<u>Nuvigil<sup>™</sup>/Provigil<sup>®</sup>/armodafinil/modafinil:</u>  Refer to <a href="#">Narcolepsy Medications SA Form</a></p>	
<b>Dermatologic</b>			
<b>*Acne Agents, Topical (AG)</b>			
<b>Combo Benzoyl Peroxide , Clindamycin, Erythromycin Topical</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<p>Acne Medication gel, lot  benzoyl peroxide wash, cr, gel,  lot (OTC)  clindacin ETZ 1% pledget  clindamycin ph 1% solution  clindamycin Phos 1% pledget,  swab</p>	<p>Acanya<sup>™</sup> w/pump  Acne Clearing System<sup>®</sup> (OTC)  Aczone<sup>®</sup> Gel and Gel Pump  Aklied<sup>®</sup>  Amzeeq<sup>™</sup>  Arazlo<sup>™</sup>  Avar Cleanser, Medicated Pad</p>	<p><b>Routine PDL edits plus</b>  <b>*Clinical Criteria for Dermatologic Acne Agents</b></p> <ul style="list-style-type: none"> <li>• Prescriptions for members over the age of 18 years will require the submission of a SA to evaluate treatment diagnosis; <b>AND</b></li> <li>• Drugs are intended for acne <b>only</b>. <b>SA for a cosmetic indication cannot be approved.</b></li> </ul>	





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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
	<p><b>clindamycin/benzoyl peroxide (Duac®)</b>  <b>erythromycin solution</b>  <b>Panoxyl-4 Acne Cr Wash (OTC)</b>  <b>Panoxyl 10 cleansing bar, foaming wash (OTC)</b></p>	<p><i>Avar-E</i>  <i>Avar-E LS</i>  <i>Avar LS Cleanser, Medicated Pad</i>  <i>Azelex®</i>  <i>Benzaclin® &amp; Benzaclin® Pump</i>  <i>BP 10-1</i>  <i>Benzefoam™ regular &amp; Ultra™</i>  <i>Benzepro</i>  <i>benzoyl peroxide wash/cr/gel/lotion/foam/towelette (RX)</i>  <i>benzoyl peroxide 6%, 9% cleanser (OTC)</i>  <i>BPO Kit</i>  <i>Cleocin T®</i>  <i>Clindacin™ Pac Kit</i>  <i>Clindagel®</i>  <i>clindamycin phosphate(generic for Clindagel®)</i>  <i>clindamycin/benzoyl peroxide (generic for Acanya® Pump)</i>  <i>clindamycin/benzoyl peroxide (generics for Benzaclin®)</i>  <i>clindamycin phosphate foam, el, lotion, med swab</i>  <i>clindamycin/tretinoin (generic Veltin®)</i>  <i>Delos™ Lotion</i>  <i>Duac® gel</i>  <i>erythromycin gel/med. swab</i>  <i>Evoclin™</i>  <i>Inova™</i>  <i>Lavoclen™ Cleanser &amp; Kit</i>  <i>Neuac™ topical/kit</i>  <i>Onexton™ gel &amp; w/Pump</i>  <i>Ovace® Wash</i></p>	



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<p><i>Ovace® Plus shampoo/cr/lotion/foam</i>  <i>Pacnex®HP &amp; LP</i>  <i>Panoxyl® 3% cr (OTC)</i>  <i>Promiseb® Complete</i>  <i>Rosula Cleanser</i>  <i>Se BPO® Wash Kit &amp; cleanser</i>  <i>Sulfacetamide Cleanser ER</i>  <i>Sulfacetamide Cleanser, Shampoo, Susp</i>  <i>Sulfacetamide Sodium/Sulfur Cr, Susp, Sunscreen</i>  <i>SSS 10-5 Foam</i>  <i>Sulfacetamide/Sulfur/Cleanser, Cleanser Kit, Lotion Med. Pad</i>  <i>Sulfacetamide/Sulfur/Urea Cleanser</i>  <i>Sumadan Wash, Kit</i>  <i>Sumadan XLT</i>  <i>Sumaxin CP Kit</i>  <i>Veltin®</i></p>	
	<b>Retinoids/Combinations, Topical</b>		
	<p><b>Differin 0.1% gel (OTC)</b>  <b>Retin® A 0.025., 0.05, 0.1 % cr &amp; 0.01, 0.025% gel</b></p>	<p><i>Acnefree® Severe Kit (OTC)</i>  <i>adapalene 0.1% cr/gel/lot</i>  <i>adapalene 0.3% gel/gel w/pump</i>  <i>adapalene-benzoyl peroxide (generic Epiduo®)</i>  <i>Altreno™</i>  <i>Atralin® 0.05% gel</i>  <i>Avage® 0.1% cr</i>  <i>Avita® 0.025% cr/gel</i>  <i>Differin® 0.1% cr/gel/lot RX</i>  <i>Differin® 0.3% gel pump</i>  <i>Epiduo® &amp; Epiduo® Forte Gel</i>  <i>*Fabior™01% Foam (AG)</i>  <i>Renova® 0.02% cr/cr pump</i></p>	<p><b>*Age Edit for Fabior™ Foam</b></p> <ul style="list-style-type: none"> <li>Member must be between the ages of 12 and 18 years of age</li> </ul>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		Retin <sup>®</sup> -A Micro 0.04%, 0.1% gel Retin <sup>®</sup> -A Micro 0.08%, 0.04%, 0.1% pump Tazorac <sup>®</sup> cr/gel tazarotene 0.1% cr <b>tretinoin 0.025, 0.1% cr &amp; 0.01,            0.025, 0.05% gel</b> tretinoin microsphere 0.04% & 0.1% gel Ziana <sup>®</sup> gel	
<b>Antifungal Topical</b>			
	antifungal 1% cr, powder antifungal 2% cr ciclopirox soln ciclodan 8% soln clotrimazole cr (OTC & RX) clotrimazole soln (OTC) clotrimazole-betamethasone cr ketoconazole shampoo ketoconazole cr miconazole cr/spray (OTC) nystatin cr/oint/ powder terbinafine cr (OTC) tolnaftate cr/powder/soln (OTC)	Alevazol <sup>®</sup> OTC Azolen <sup>®</sup> Tincture OTC Bensal HP <sup>®</sup> Ciclodan <sup>®</sup> Kit ciclopirox cr/shampoo/gel ciclopirox kit ciclopirox suspension clotrimazole soln (RX) clotrimazole-betamethasone lot *CNL 8 <sup>™</sup> Kit Desenex <sup>®</sup> Aero Powder (OTC) econazole Ertaczo <sup>®</sup> Exelderm <sup>®</sup> cr & soln Extina <sup>®</sup> Fungi-Nail <sup>®</sup> (OTC) Fungoid <sup>®</sup> Kit (OTC) Fungoid <sup>®</sup> (OTC) *Jublia <sup>®</sup> ketoconazole foam *Kerydin <sup>®</sup> Lamisil AT <sup>®</sup> cr/gel (OTC) Lamisil <sup>®</sup> Spray (OTC)	<b>LENGTH OF AUTHORIZATIONS: 6 months</b>  <b>Routine PDL edits plus</b>  <b>Select non-preferred topical Antifungals (CNL-8<sup>™</sup>, Jublia<sup>®</sup>, Kerydin<sup>™</sup>, Luzu<sup>®</sup>, Penlac<sup>®</sup>) require the submission of a Clinical SA. Refer to <a href="#">Antifungal Topical SA Form</a></b>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<p><i>Loprox® Kit/ Shampoo/susp</i>  <i>Lotrimin AF® cr (OTC)</i>  <i>Lotrimin Ultra® (OTC)</i>  <i>Lotrisone® cr</i>  <i>luliconazole (generic for Luzu)</i>  <i>**Luzu®</i>  <i>miconazole nitrate (OTC)</i>  <i>miconazole Oint/ powder (OTC)</i>  <i>Mentax®</i>  <i>Naftin® cr/gel</i>  <i>Naftifine CR</i>  <i>Nyata Kit®</i>  <i>Nizoral A-D® Shampoo (OTC)</i>  <i>nystatin-triamcinolone cr/oint</i>  <i>oxiconazole cr (generic Oxistat®)</i>  <i>Oxistat® cr</i>  <i>Oxistat® Lotion</i>  <i>Pediaderm AF®</i>  <i>PediPak®</i>  <i>*Penlac®</i>  <i>tavaborole (generic Kerydin®)</i>  <i>Tinactin® Aero powder/spray(OTC)</i>  <i>tolnaftate aero powde/spray (OTC)</i>  <i>Vusion®</i></p>	
<b>Immunomodulators Atopic Dermatitis</b>			
	<p><b>*Elidel®</b></p>	<p><i>*Eucrisa™</i>  <i>**Dupixent® (QL, AG)</i>  <i>pimecrolimus (new generic for Elidel)</i>  <i>*Protopic®</i>  <i>*tacrolimus</i></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year; EXCEPT Dupixent® 6 months</p> <p><b>ATOPIC DERMATITIS - REAUTHORIZATION:</b></p> <ul style="list-style-type: none"> <li>Documentation (i.e., progress note) of positive clinical response will be required</li> </ul> <p>Length of authorization: Initial approval = 6 months; renewal = 1 year</p> <p><b>Routine PDL edits plus</b></p> <p><b>*Clinical Criteria for Elidel®, Eucrisa™, Protopic® &amp; tacrolimus</b></p> <ul style="list-style-type: none"> <li>Member must have a FDA approved diagnosis:</li> </ul>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
			<ul style="list-style-type: none"> <li>○ Atopic dermatitis</li> <li>○ Elidel® mild to moderate for ages &gt; 2 years</li> <li>○ Eucisa™: mild to moderate for ages equal to or &gt; 3months</li> <li>○ Protopic® 0.03%: moderate to severe for ages &gt; 2 years.</li> <li>○ Protopic® 0.1%: moderate to severe for ages &gt; 18 years; <b>AND</b></li> <li>● Failure to topical corticosteroids (i.e., desonide, fluticasone propionate, hydrocortisone butyrate, etc.)</li> </ul> <p><b>** See Cytokine and CAM Antagonists Appendix A for Clinical Criteria**</b></p> <p><b>**Clinical Criteria for Dupixent®</b></p> <ul style="list-style-type: none"> <li>❖ <b><u>Atopic Dermatitis</u></b></li> <li>● ≥ 6 years of age; <b>AND</b></li> <li>● Diagnosis of moderate to severe atopic dermatitis; <b>AND</b></li> <li>● Prior documented trial and failure (or contraindication) of 1 topical corticosteroids of medium to high potency (e.g., mometasone, fluocinolone) and 1 topical calcineurin inhibitors (tacrolimus or pimecrolimus); <b>AND</b></li> <li>● Inadequate response to a 3 month minimum trial of at least 1 immunosuppressive systemic agent (e.g., cyclosporine, azathioprine, methotrexate, mycophenolate mofetil, etc.); <b>AND</b></li> <li>● Inadequate response (or is not a candidate) to a 3 month minimum trial of phototherapy (e.g., psoralens with UVA light [PUVA], UVB, etc) provided member has reasonable access to photo treatment; <b>AND</b></li> </ul> <p>Is not pregnant; <b>AND</b></p> <p><b><u>Quantity limit Dupixent®</u></b>  <i>2 prefilled syringes for the initial dose, then 1 single-dose syringe every 14 days</i></p> <ul style="list-style-type: none"> <li>❖ <b><u>Chronic Rhinosinusitis with Nasal Polyposis</u></b></li> <li>● ≥ 18 years of age; <b>AND</b></li> <li>● Diagnosis of In adequately controlled Chronic Rhinosinusitis with Nasal Polyposis (CRSwNP)</li> <li>● Is Add to current maintenance treatment</li> </ul>
<b>Psoriasis, Topical</b>			
	calcipotriene cr/oint/soln	betamethasone/ calcipotriene	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year

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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		Calcitrene <sup>®</sup> calcitriol calcipotriene/betamethasone (generic for Taclonex <sup>®</sup> ) Dovonex <sup>®</sup> Duobrii <sup>™</sup> *Enstilar <sup>®</sup> Foam (AG) Micanol <sup>®</sup> Sorilux <sup>™</sup> Taclonex <sup>®</sup> & Taclonex <sup>®</sup> Scalp Vectical	Routine PDL edits plus  <u>*Clinical Criteria for Enstilar<sup>®</sup> Foam</u> <ul style="list-style-type: none"> <li>Length of Authorization: 4 weeks</li> <li>Diagnosis of plaque psoriasis; AND</li> <li>Minimum age of 18 years</li> </ul>
<b>Rosacea Agents, Topical</b>			
	Metrocream <sup>®</sup> Metrogel <sup>®</sup> Metro lotion <sup>®</sup>	azelaic acid (generic for Finacea <sup>®</sup> ) Finacea <sup>®</sup> foam/gel ivermectin (generic Soolantra) metronidazole cr/gel/lot Mirvaso <sup>®</sup> Noritate <sup>®</sup> Rosadan <sup>™</sup> Kit Soolantra <sup>®</sup> Zilxi <sup>™</sup>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year  Routine PDL edits
<b>Steroids</b>			
	<b>Steroids, Topical Low Potency</b> hydrocortisone cr/gel/lot/oint	anusol-hc 2.5% cr alclometasone cr/oint aqua glycolic HC Capex <sup>®</sup> shampoo Derma-smoothe-FS desonate gel/cr/lot/oint Desowen <sup>®</sup> lot fluocinolone 0.01% oil Micort <sup>™</sup> -HC Pediderm <sup>®</sup> HC & Pediderm <sup>®</sup> TA	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year  Routine PDL edits



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Preferred Agents	Non-Preferred Agents	SA Criteria
	<p><i>Texacort<sup>®</sup></i></p>	
<b>Steroids, Topical Medium Potency</b>		
<p><b>fluticasone propionate cr/oint</b>  <b>mometasone furoate cr/oint/soln</b></p>	<p><i>betamethasone valerate foam</i>  <i>clocortolone cr</i>  <i>Cloderm<sup>®</sup></i>  <i>Cordran<sup>®</sup> tape</i>  <i>Cutivate<sup>®</sup> cr/lot</i>  <i>Dermatop<sup>®</sup> cr/oint</i>  <i>Elocon<sup>®</sup> cr/oint/soln</i>  <i>fluocinolone acetonide cr/oint/soln</i>  <i>flurandrenolide cr/oint/tape</i>  <i>fluticasone propionate lot</i>  <i>hydrocortisone valerate cr/oint</i>  <i>hydrocortisone butyrate (generic for locoid lotion)</i>  <i>hydrocortisone butyrate cr/oint/soln/ emollient</i>  <i>Luxiq<sup>®</sup></i>  <i>Momexin<sup>®</sup></i>  <i>Pandel<sup>®</sup></i>  <i>prednicarbate cr/oint</i>  <i>Synalar<sup>®</sup></i>  <i>Synalar TS<sup>®</sup></i>  <i>Ticanase kit<sup>®</sup></i></p>	
<b>Steroids, Topical High Potency</b>		
<p><b>betamethasone valerate cr/lot/oint</b>  <b>triamcinolone acetonide cr/lot/oint</b></p>	<p><i>amcinonide cr/lot/oint</i>  <i>betamet diprop &amp; prop gly cr/lot/oint</i>  <i>betamet diprop cr/foam/gel/lot/oint</i>  <i>DermacinRx<sup>®</sup> SilaPak<sup>™</sup></i>  <i>DermacinRx<sup>®</sup> Silazone</i>  <i>DermacinRx<sup>®</sup> Therazole Pak</i></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p><b>Routine PDL edits plus</b></p> <p><b>*Clinical Criteria for Sernivo<sup>™</sup></b></p> <ul style="list-style-type: none"> <li>Length of Authorization: 4 weeks (treatment beyond 4 weeks is not recommended).</li> <li>Member must have diagnosis of mild to moderate plaque psoriasis: <b>AND</b></li> </ul>

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>desoximetasone cr/gel/oint/spray</i> <i>desoximetasone (generic Topicort® spray)</i> <i>diflorasone diacetate cr/oint</i> <i>Diprolene® lot/oint</i> <i>DiproleneAF® cr</i> <i>Ellzia™ Pak Kit</i> <i>fluocinonide cr/ emollient/ gel/oint/soln</i> <i>Halog® cr/oint</i> <i>Kenalog® aerosol</i> <i>Loprox® Suspension Kit</i> <i>*Sernivo™</i> <i>Silazone® II Kit</i> <i>Topicort® cr/gel/oint/spray</i> <i>Trianex® oint</i> <i>triamcinolone spray</i> <i>triamcinolone/dimethicone</i> <i>Vanos® cr</i> <i>Whytederm® Tdpak</i>	<ul style="list-style-type: none"> <li>At least 18 years of age</li> </ul>
	<b>steroids, Topical Very High Potency</b> <b>clobetasol emollient</b> <b>clobetasol propionate cr/gel/oint/soln</b> <b>halobetasol propionate cr</b>	<i>Apexicon™ E</i> <i>Bryhali™ (halobetasol propionate)</i> <i>clobetasol lot/shampoo</i> <i>clobetasol propionate foam/spray</i> <i>Clobex® lot/shampoo/spray</i> <i>Clodan® kit</i> <i>Halonate®</i> <i>halobetasol propionate oint (generic for Lexette®)</i> <b>Impeklo™</b> <i>Olux®</i> <i>Olux®-E</i> <i>Temovate® oint</i>	





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	Preferred Agents	Non-Preferred Agents	SA Criteria
		Ultravate® cr/lotion/oint Ultravate® PAC & Ultravate® X	
<b>Endocrine and Metabolic Agents</b>			
<b>Androgenic Agents (Testosterone – Topical)</b>			
	<b>AndroGel® Pump</b> <b>Androderm® Patch</b>	AndroGel® packet, patch Axiron® soln Fortesta® Natesto Nasal Gel® Testim® testosterone pump (generic for AndroGel® Pump) testosterone (generic for Axiron®) testosterone gel/packet/pump (generic for Vogelxo™) testosterone (generic for Fortesta®) Vogelxo™ gel/packet/pump Xyosted™	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL edits Plus</b>
<b>Antihyperuricemics</b>			
	Allopurinol <b>colchicine tabs</b> Probenecid® probenecid & colchicine	<b>colchicine caps</b> Colcris® febuxostat (generic Uloric®) Gloperba® Mitigare® Uloric®	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL edits plus</b>
<b>Contraceptives*(long-acting IUDs &amp; injectable)</b>			
	Kyleena™ Liletta® medroxyprogesterone 150mg Mirena® Nexplanon® Paragard®	Depo-Provera® 104 mg and 150 mg	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year  <b>Routine PDL edits</b>



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Preferred Agents	Non-Preferred Agents	SA Criteria
Skyla®		
<b>Diabetes Hypoglycemics: Injectable Amylin Analogs</b>		
	*SymLin® *SymLin® Pens	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year  <u>*Clinical Criteria for Injectable Amylin Analogs</u> <ul style="list-style-type: none"> <li>• Member must have a history of at least a 90day trial of insulin.</li> <li>• SymLin® is only indicated as adjunct therapy with insulin.</li> <li>• Member meeting ALL of the following criteria may be approved:               <ul style="list-style-type: none"> <li>○ Diagnosis of Type 1 or 2 diabetes; AND</li> <li>○ On insulin therapy, AND</li> <li>○ Failure to achieve adequate glycemic control (HbA1c ≤ 6.5%)</li> </ul> </li> </ul>
<b>Diabetes Hypoglycemics: Injectable and Oral Incretin Mimetics <span style="color: red;">CLOSED CLASS</span></b>		
Byetta® (exenatide) Bydureon™ (exenatide ER) <b>Trulicity™ (lixisenatide)</b> Victoza® (liraglutide)	Adlyxin™ (lixisenatide) Bydureon™ Bcise SQ Soliqua® 100/33 (insulin glargine & lixisenatide inj) Ozempic® Rybelsus Tab® Tanzeum™ (albiglutide) Xultophy® 100/3.6 (insulin glargine & lixisenatide inj)	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year  <b>Routine PDL edits</b>
<b>Diabetes Hypoglycemics: Injectable Insulins</b>		
<b>Insulin Mix</b>		<u>LENGTH OF AUTHORIZATIONS:</u> 1 year
Humalog® Mix 50/50 vial Humalog® Mix 75/25 vial Humulin® 70/30 Pen OTC, vial Novolog® Mix 70/30 pen/vial	Humalog® Mix 50/50 Kwikpen Humalog® Mix 75/25 Kwikpen <b>insulin lispro protamine mix pen(generic for Humalog mix pen)</b> Novolin® 70/30 vial (OTC)	<b>Routine PDL edits</b>



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Preferred Agents		Non-Preferred Agents	SA Criteria
<b>Insulin N</b>			
<b>Humulin® N pen/vial (OTC)</b>		<i>Humulin® N pen</i> <i>Novolin® N vial (OTC)</i>	
<b>Insulin R</b>			
<b>Humulin® R pen/vial</b>		<i>Novolin® R vial (OTC)</i>	
<b>Long-Acting Insulins</b>			
<b>Lantus® Solostar® &amp; vial</b> (insulin glargine)		<i>Basaglar® KwikPen® (insulin glargine)</i>	
<b>Levemir® pen/vial</b> (insulin detemir)		<i>Semglee™ (insulin glargine)</i> <i>Toujeo® Solostar® (insulin glargine)</i> 300 Units/mL <i>Tresiba® FlexTouch® Pen (insulin degludec)</i> 100 U/ml, 200 U/ml	
<b>Rapid-Acting Insulins</b>			
<b>Humulin 500 U/M pen/vial</b>		<i>Admelog® Solostar Pen/vial</i>	
<b>Humalog® vial</b>		<i>Afrezza® cartridge (inhalation)</i>	
<b>Humalog® Cartridge</b>		<i>Apidra® cartridge/Solostar/vial</i>	
<b>Humalog Kwikpen 100 unit/ml</b>		<i>Fiasp®/FlexTouch® Pen/PenFill®</i>	
<b>Humalog Jr. Kwikpen</b>		<i>Humalog Kwikpen 200 unit/ml</i>	
<b>Novolog® cart/vial/Flexpen</b>		<i>insulin lispro Jr. Kwikpen (generic for Humalog Jr. Kwikpen)</i>	
<b>insulin lispro vial</b>		<i>Lyumjev™</i>	
<b>Diabetes Oral Hypoglycemics</b>			
<b>Oral Hypoglycemics Alpha-Glucosidase Inhibitors</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<b>acarbose</b>		<i>Glyset®</i> <i>miglitol (generic Glyset®)</i>	<b>Routine PDL edits plus</b>
<b>Oral Hypoglycemics Biguanides</b>			<b>Metformin Step Edit for <u>all Oral Hypoglycemics</u> (excluding metformin)</b>
<b>metformin</b>		<i>Fortamet®</i>	<ul style="list-style-type: none"> <li>Member is 18 or older</li> <li>Patients with a <b>hemoglobin A1C &lt; 7.5%</b> must have a minimum 90-day trial of metformin (unless contraindicated*)</li> </ul>
<b>metformin ER</b> (generic for <i>Glucophage® XR</i> )		<i>Glucophage® IR &amp; XR</i> <i>Glutmetza®</i> <i>Riomet® susp</i>	

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		<i>metformin ER (generic Fortamet®)</i> <i>metformin ER (generic Glumetza®)</i> <i>metformin (generic Riomet®)</i>	<ul style="list-style-type: none"> <li>Patients with a hemoglobin A1C <math>\geq</math> 7.5% should be started on metformin (unless contraindicated) <b>plus a second agent</b> (e.g., DPP-IV, SGLT2, GLP-1 receptor agonists, TZDs, sulfonylureas). <b>A 90-day trial of metformin is NOT required.</b></li> </ul> <p><b>*Contraindications include:</b></p> <ul style="list-style-type: none"> <li>severe renal impairment (eGFR below 30mL/min/1.73m2)</li> <li>known <b>intolerance</b></li> <li>acute or chronic metabolic acidosis including diabetic ketoacidosis</li> </ul> <p>Age edit for all Oral Hypoglycemics is 18 years of age or older, except Metformin which is 10 years of age.</p>
<b>Oral Hypoglycemics Biguanide Combination Drugs</b>			
glyburide/metformin	<i>glipizide/metformin</i> <i>Glucovance®</i>		
<b>Oral Hypoglycemics DPP-IV Inhibitors &amp; Combination <b>CLOSED</b></b>			
Janumet® Janumet XR® Januvia® Jentadueto™ Tradjenta™	<i>alogliptin (generic Nesina™)</i> <i>alogliptin/metformin (generic Kazano™)</i> <i>alogliptin/pioglitazone (generic Oseni™)</i> <i>Jentadueto XR™</i> <i>Kazano™</i> <i>Kombiglyze XR™</i> <i>Nesina™</i> <i>Onglyza™</i> <i>Oseni™</i> <i>Trijardy™ XR</i> <i>(empagliflozin/linagliptin/metformin)</i>		
<b>Oral Hypoglycemics Meglitinides</b>			
repaglinide nateglinide	<i>Prandin® &amp; PrandiMet™</i> <i>repaglinide/metformin</i> <i>Starlix</i>		
<b>Oral Hypoglycemics Second Generation Sulfonylureas</b>			
glimepiride glipizide glipizide ER glyburide glyburide micronized	<i>Amaryl®</i> <i>Diabeta®</i> <i>Glucotrol®</i> <i>Glucotrol XL®</i> <i>Glynase®</i>		



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Preferred Agents	Non-Preferred Agents	SA Criteria
<b>*Oral Hypoglycemics Sodium Glucose Co-Transporter 2 Inhibitor</b> <b>CLOSED CLASS</b>		
<b>Farxiga™ (AG)</b> <b>Glyxambi® (AG)</b> <b>Invokana™ (AG)</b> <b>Invokamet™ (AG)</b> <b>Invokamet™ XR (AG)</b> <b>Jardiance®(AG)</b> <b>Synjardy® (AG)</b> <b>Xigduo™ XR (AG)</b>	<b>Segluromet™</b> <i>(ertugliflozin/metformin)</i> <b>Steglatro™</b> <b>Steglujan™</b> <b>Synjardy® XR</b>	
<b>Oral Hypoglycemics Thiazolidinediones</b>		
<b>pioglitazone</b>	<b>Avandia®</b> <b>Actoplus Met® IR &amp; XR</b> <b>Actos®</b> <b>Avandaryl®</b> <b>Avandamet®</b> <b>Duetact®</b> <i>pioglitazone/metformin</i> <i>pioglitazone/glimepiride</i>	
<b>Erythropoiesis Stimulating Proteins</b>		
<b>Epogen®</b> <b>Retacrit™</b>	<b>Aranesp® vial/syringe</b> <b>Procrit®</b> <b>Mircera®</b>	<b>LENGTH OF AUTHORIZATIONS:</b> for duration of the prescription up to 6 months <b>Routine PDL edits</b> <i>Omontys® is not PDL eligible, may be covered under medical benefit</i>
<b>Glucocorticoids, Oral</b>		
<b>budesonide EC</b> <b>dexamethasone soln/tab</b> <b>hydrocortisone</b> <b>methylprednisolone dose pk</b> <b>methylprednisolone 4 mg tab</b> <b>prednisolone sodium phosphate soln</b> <b>prednisolone soln</b> <b>prednisone soln/tab/dose pk</b>	<b>Alkindi® Sprinkle</b> <b>Cortef®</b> <i>cortisone acetate</i> <i>dexamethasone elixir/intensol</i> <b>Dexpak®</b> <b>*Emflaza™ (AG)</b> <b>Hemady®</b> <b>*Emflaza™ (AG)</b> <b>Entocort® EC</b>	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year <b>Routine PDL edits plus</b> <b>*Clinical Criteria for Emflaza™</b> <ul style="list-style-type: none"> <li>• Trial and failure of all drugs does not apply to Emflaza™</li> <li>• Diagnosis for treatment of Duchenne muscular dystrophy (DMD)</li> <li>• Minimum Age Limit = 2 years of age</li> </ul>



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Preferred Agents	Non-Preferred Agents	SA Criteria
	<p><i>Flo-Pred<sup>®</sup></i>  <i>Medrol<sup>®</sup> dose pk/tab</i>  <i>methylprednisolone 8,16 &amp; 32mg tab</i>  <i>Millipred DP<sup>®</sup> tab Does Pk</i>  <i>Millipred<sup>®</sup> soln/tab</i>  <i>Orapred<sup>®</sup> ODT</i>  <b>Ortikos<sup>™</sup></b>  <i>prednisolone sod phosphate ODT/soln</i>  <i>prednisone intensol</i>  <i>Rayos<sup>®</sup> DR tab</i>  <i>TaperDex<sup>®</sup></i>  <i>Veripred<sup>®</sup></i></p>	
<p><b>*Growth Hormone</b> <span style="float: right;"><b>CLOSED CLASS</b></span></p>		
<p><b>Genotropin<sup>®</sup> Cartridge,</b>  <b>Miniquick</b>  <b>Norditropin FlexPro<sup>®</sup></b></p>	<p><i>Humatrope<sup>®</sup> cartridge/vial</i>  <i>Nutropin AQ<sup>®</sup> NuSpin<sup>™</sup></i>  <i>Omnitrope<sup>®</sup> cartridge/vial</i>  <i>Saizen<sup>®</sup> cartridge/vial</i>  <i>Serostim<sup>®</sup> vial</i>  <i>Zomacton<sup>®</sup> vial</i>  <i>Zorbtive<sup>®</sup> vial</i></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p><b>ALL Growth Hormone drugs (preferred and non-preferred) require the submission of a Clinical SA. Refer to <a href="#">(Growth Hormone SA Fax Form)</a></b></p>
<p><b>*Hereditary Angioedema (HAE) Agents</b></p>		
<p><b>Berinert<sup>®</sup></b>  <b>Cinryze<sup>™</sup></b>  <b>Kalbitor<sup>®</sup></b></p>	<p><i>Firazyr<sup>®</sup></i>  <i>Haegarda<sup>®</sup></i>  <i>Icatibant (generic Firazyr<sup>®</sup>)</i>  <b>Orladeyo<sup>™</sup></b>  <i>Ruconest<sup>®</sup></i>  <i>Takzyro<sup>™</sup></i></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p><b>Routine PDL edits plus</b></p> <p><b>*_ALL Hereditary Angioedema drugs (preferred and non-preferred) require the submission of a Clinical SA. Refer to <a href="#">Hereditary Angioedema (HAE) SA Form</a></b></p> <p><b>**Black box warning KALBITOR<sup>®</sup></b></p> <p><b>Anaphylaxis has been reported after administration of KALBITOR<sup>®</sup>. Because of the risk of anaphylaxis, KALBITOR should only be administered by a healthcare professional with appropriate medical support to manage anaphylaxis and hereditary angioedema. Healthcare</b></p>



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Preferred Agents	Non-Preferred Agents	SA Criteria
		<p>professionals should be aware of the similarity of symptoms between hypersensitivity reactions and hereditary angioedema and patients should be monitored closely.</p>
<b>*Pancreatic Enzymes</b>		
<p>Creon<sup>®</sup> Zenpep<sup>®</sup></p>	<p>Pancreaze<sup>®</sup> Pertzye<sup>®</sup> Ultresa<sup>®</sup> Viokace<sup>®</sup></p>	<p><b>LENGTH OF AUTHORIZATION:</b> 1 year</p> <p>Routine PDL edits plus</p> <p><b>*Clinical Criteria for Pancreatic Enzymes</b></p> <ul style="list-style-type: none"> <li>All drugs preferred and non preferred require a diagnosis of pancreatic insufficiency due to cystic fibrosis or chronic pancreatitis or pancreatectomy.               <ul style="list-style-type: none"> <li>If a member has a diagnosis of Cystic Fibrosis they do not have to try and fail of a preferred.</li> <li>If member has a feeding tube, then two different pancreatic enzymes can be approved for use together.</li> </ul> </li> </ul>
<b>Progestational Agents</b>		
<p>Makena<sup>®</sup> Auto-injector medroxyprogesterone acetate (tab only) norethindrone acetate progesterone cap/inj</p>	<p>Aygestin<sup>®</sup> Crinone (Vaginal) Depo-Provera 400 MG/ML hydroxyprogesterone caproate SDV hydroxyprogesterone caproate (generic for Makena MDV) Makena<sup>®</sup> Multi Dose Vial (MDV) &amp; Single Dose Vial (SDV) Prometrium<sup>®</sup> Provera<sup>®</sup></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p>Routine PDL edits</p>
<b>Progestins Used For Cachexia</b>		
<p>megestrol acetate</p>	<p>Megace<sup>®</sup> Megace<sup>®</sup> ES megestrol suspension ES</p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p>Routine PDL edits</p>
<b>Vaginal Estrogens</b>		
<p>Premarin<sup>®</sup> Vaginal cr</p>	<p>Estrace<sup>®</sup> Vaginal cr</p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 6 months</p>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
	Vagifem® Vaginal tab	estradiol cream (generic Estrace®) Estring® Vaginal ring Femring® Vaginal ring Invexxy® Intrarosa™ Osphena® tab Yuvafem®	Routine PDL edits
<b>Gastrointestinal</b>			
<b>G I Antibiotics</b>			
	Firvanq™ sol metronidazole tab vancomycin cap	Aemcolo™ Alinia® Difucid® Flagyl® cap/tab/ER metronidazole cap neomycin nitazoxanide (generic Alinia®) paromomycin Solosec™ Tindamax® tinidazole Xifaxan® vancomycin compounded oral soln kit Vancocin®	Length of authorization: 1 year  Routine PDL edits plus
<b>Antiemetic/Antivertigo Agents</b>			
<b>Cannabinoids (delta-9THC derivatives)</b>			
	*dronabinol cap	*Cesamet™ *Marinol® *Syndros®	<b>LENGTH OF AUTHORIZATIONS:</b> 6 months  *Dronabinol plus all <b>non-preferred</b> Antiemetic/Antivertigo agents require submission of a Clinical SA. Refer to <a href="#">Antiemetic/Antivertigo SA form</a>
<b>5HT3 Receptor Blockers</b>			
	ondansetron ODT/soln/tab	Aloxi® Anzemet® Akynzeo®	<b>LENGTH OF AUTHORIZATIONS:</b> 3 months, unless otherwise noted  Routine PDL





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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>granisetron Granisol<sup>®</sup> soln/tab</i> <i>Kytril<sup>®</sup></i> <i>palonosetron (generic Aloxi<sup>®</sup>)</i> <i>Sancuso<sup>®</sup> patch</i> <i>Zofran<sup>®</sup> ODT/soln/tab</i> <i>Zuplenz<sup>®</sup> film</i>	
	<b>NK-1 Receptor Antagonist</b>		
		<i>aprepitant capsule/pack</i> <i>Cinvanti<sup>™</sup> (Intraven)</i> <i>Emend<sup>®</sup> Bi Pak/ cap</i> <i>Emend<sup>®</sup> Tri-fold pack/susp</i> <i>Varubi<sup>™</sup> IV, Tab</i>	<b>LENGTH OF AUTHORIZATIONS:</b> Length of chemotherapy regimen or a maximum of 6 months
	<b>Other</b>		<b>LENGTH OF AUTHORIZATIONS:</b> 1 year, unless otherwise noted
	<b>HM motion relief 25 mg tab</b> <b>meclizine cap, tab chew, tab</b> <b>metoclopramide sol, syr, tab,</b> <b>vial</b> <b>motion sickness rlf 25 mg tab</b> <b>motion-time tab chew</b> <b>**phenadoz sup</b> <b>Prochlorperazine tab</b> <b>**promethazine sol, syrup, tab</b> <b>(AG)</b> <b>SM motion sickness 25 mg tab</b> <b>travel sickness 25 mg tab chew</b>	<i>Antivert<sup>®</sup></i> <b><i>Barhemsys<sup>®</sup></i></b> <i>Bonjesta<sup>™</sup></i> <i>Compazine<sup>®</sup> supp/tab</i> <i>Compro<sup>®</sup></i> <i>Diclegis<sup>®</sup></i> <i>dimenhydrinate</i> <i>doxylamine succinate/ vit B6</i> <i>Metozolv<sup>®</sup> ODT</i> <i>metoclopramide ODT</i> <i>**Phenergan<sup>®</sup> (AG)</i> <i>prochlorperazine supp</i> <i>**promethazine 50mg supp (AG)</i> <i>Reglan<sup>®</sup></i> <i>scopolamine (generic Transderm-Scop<sup>®</sup>)</i> <i>Tigan<sup>®</sup></i> <i>Transderm-Scop<sup>®</sup></i> <i>trimethobenzamide</i> <i>Vistaril<sup>®</sup></i>	<b>**Promethazine products approved for members <math>\geq</math> 2 years</b>
	<b>*GI Motility, Chronic</b>		



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<p>Amitiza<sup>®</sup>                      Linzess<sup>™</sup>                      Movantik<sup>®</sup></p>	<p>Alosetron                      lubiprostone (generic Amitiza<sup>®</sup>)                      Lotronex<sup>®</sup>                      Motegrity<sup>™</sup>                      Relistor<sup>®</sup>                      Symproic<sup>®</sup>                      Trulance<sup>™</sup>                      Viberzi<sup>™</sup></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 6 months</p> <p>Routine PDL edits plus</p> <p>*All GI Motility drugs (<b>preferred and non-preferred</b>) require the submission of a Clinical SA. Refer to <a href="#">Chronic GI Motility SA form</a></p>
<b>H. Pylori Treatment</b>		
<p>Pylera<sup>®</sup></p>	<p>Helidac<sup>®</sup>                      Omeclamox<sup>®</sup>-Pak                      lansoprazole/amoxicillin/                      clarithromycin                      Prevpac<sup>®</sup>                      Talicia<sup>®</sup></p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 14 days</p> <p>Routine PDL edits</p>
<b>Histamine-2 Receptor Antagonists (H-2 RA)</b>		
<p>famotidine (OTC &amp; RX)                      *famotidine oral susp</p>	<p>cimetidine tab/syrup (OTC/RX)                      nizatidine cap/susp                      Pepcid<sup>®</sup> susp/tab (OTC/RX)                      ranitidine cap (OTC/RX)                      Zantac<sup>®</sup> syrup/tab (OTC/RX)</p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 1 year</p> <p>Routine PDL edits plus</p> <p>*AutoPA for famotidine oral susp</p> <ul style="list-style-type: none"> <li>• Is under 12 years of age</li> <li>• Continuation of therapy, has a claim in past 90 days</li> <li>• Is using a Tube feeding</li> </ul>
<b>Proton Pump Inhibitors</b>		
<p>omeprazole RX                      pantoprazole</p>	<p>Aciphex<sup>®</sup> DR tab/sprinkle                      Dexilant<sup>®</sup>                      esomeprazole magnesium cap/susp                      esomeprazole strontium                      lansoprazole cap                      Nexium<sup>®</sup>                      omeprazole OTC                      omeprazole magnesium OTC                      omeprazole/sodium bicarbonate</p>	<p><b>LENGTH OF AUTHORIZATIONS:</b> 12 weeks; unless member meets an exception; then 1 year</p> <p>Routine PDL edits plus</p> <p>*All <b>non-preferred</b> Proton Pump Inhibitors require submission of a Clinical SA. Refer to <a href="#">Proton Pump Inhibitor SA form</a></p> <p>Preferred agents require a SA for use over 90 days</p>



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		<p><i>pantoprazole susp (generic Protonix® susp)</i>  <i>Prevacid® RX, OTC&amp; Solutab</i>  <i>Prilosec® Rx &amp; Susp</i>  <i>Protonix®</i>  <i>rabeprazole DR tab</i>  <i>Zegerid® cap/OTC/susp packet</i></p>	
<b>Ulcerative Colitis Oral and Rectal Preparations (5-ASA DERIVATIVES)</b>			
<b>Ulcerative Colitis – Oral</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<p><b>Apriso®</b>  <b>balsalazide disodium</b>  <b>Pentasa®</b>  <b>sulfasalazine DR &amp; IR</b></p>	<p><i>Asacol®HD</i>  <i>Azulfidine® IR &amp;DR</i>  <i>budesonide ER (generic Uceris™)</i>  <i>Colazal®</i>  <i>Delzicol™</i>  <i>Dipentum</i>  <i>*Giazo™ (QL)</i>  <i>Lialda®</i>  <i>mesalamine (generic Asacol® HD)</i>  <i>mesalamine (generic Lialda®)</i>  <i>Uceris™</i></p>	<p><i>Routine PDL edits</i></p> <p><i>*Giazo is limited to an 8week supply</i></p>	
<b>Ulcerative Colitis – Rectal</b>			
<p><b>mesalamine rectal supp</b>  <b>mesalamine enema</b></p>	<p><i>Canasa® rectal supp</i>  <i>mesalamine kit</i>  <i>Rowasa® enema/kit</i>  <i>SFRowasa®</i>  <i>Uceris®</i></p>		
<b>Genitourinary</b>			
<b>Alpha-Blockers and Androgen Hormone Inhibitors For Benign Prostatic Hypertrophy (BPH)</b>			
<b>Alpha-Blockers for BPH</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<p><b>alfuzosin</b>  <b>tamsulosin HCL</b></p>	<p><i>Flomax®</i>  <i>Rapaflo®</i>  <i>Silodosin (generic Rapaflo)</i>  <i>Uroxatral®</i></p>	<p><b>Routine PDL edits plus</b></p>	



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Preferred Agents		Non-Preferred Agents	SA Criteria
<b>Androgen Hormone Inhibitors for BPH</b>			
dutasteride finasteride	Avodart® Dutasteride/tamsulosin Jalyn® Proscar®		
<b>Phosphodiesterase (PDE) 5 Inhibitor for BPH</b>			
	*Cialis® (ST) tadalafil 5mg (ST)		*Step edit for Cialis® & tadalafil 5mg - trial and failure of Alpha Blockers and Androgen Inhibitors for BPH. Prescriber must attest that the member is not on the state's sex offenders list. Consult or evaluation by Urologist.
<b>Urinary Antispasmodics (Bladder Relaxant)</b>			
oxybutynin tab/syrup oxybutynin ER solifenacin Toviaz™	darifenacin ER (generic Enablex®) Detrol® & Detrol® LA Ditropan® & *Ditropan® XL flavoxate Gelnique™ gel/gel Pump Myrbetriq™ Oxytrol® transdermal includes OTC Sanctura XR trospium IR & ER tolterodine IR & ER VESIcare®		
<b>Immunological Agents</b>			
<b>*Multiple Sclerosis</b>			
Avonex® Avonex® Adm Pack Betaseron® Copaxone 20 mg syringe® **Kesimpta® (ST) Tecfidera®	***Ampyra® Aubagio® Bafiertam™ (monomethyl fumarate) Copaxone® 40 mg syringe® dimethyl fumarate (generic Tecfidera®) dalfampridine ER (generic Ampyra®) Extavia® Kit *Gilenya® (ST) Glatopa™ Mavenclad®	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year <b>Routine PDL edits plus.</b> <b>*All agents require adherence to the documented PI age and diagnosis.</b> <b>**Kesimpta® requires step trial of brand Tecfidera® or a preferred Injectable agent, to be determined by an AutoPA.</b> <b><u>Mavzent, Mavenclad or Zeposia SA Form</u></b>	

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	Mayzent® Plegridy® Rebif® SQ Rebif® Rebidose Pen® ****Vumerity™ Zeposia®	<p><b>Step Edit for Gilenya®</b>—a trial and failure of a preferred injectable drug. In order to receive a non-preferred oral drug both an injectable preferred and Gilenya® must have been tried and failed.</p> <p><b>Vumerity™</b>                      Approved for the treatment of relapsing forms of multiple sclerosis, to include clinically isolated syndrome, relapsing-remitting disease, and active secondary progressive disease, in adults.</p>
<b>Drugs to improve walking in patients with (MS)</b>	***Ampyra® ***dalfampridine ER (generic Ampyra®)	<p>***Select <b>non-preferred</b> MS drugs (Ampyra®) require the submission of a Clinical SA. Refer to <a href="#">MS - Ampyra® SA form</a></p>
<b>Cytokine and CAM Antagonists And Related Agents</b>		<p><b>CLOSED CLASS</b></p>
Enbrel® Pen, Sureclick, Syringe, Vial Humira® Pen, Syringe methotrexate tab/PF vial/MDV Renflexis®	Actemra® SQ & ACTPEN Arcalyst® Avsola™ Cimzia® & Cimzia® Syringe Kit Cosentyx™ Enspryng™ Entyvio® **Dupixent® Ilaris® Ilumya™ Inflectra® Kevzara® inj, pen Kineret® Olumiant® Otezla® Otrexup® Orencia® Rasuvo™ Remicade® Rinvoq™ Skyrizi™ Siliq®	<p><b>LENGTH OF AUTHORIZATION:</b> 1 year</p> <p>Routine PDL edits plus</p> <p>*All <b>non-preferred</b> Cytokine and CAM Antagonists require submission of a Clinical SA. Refer to <a href="#">Cytokine and CAM Antagonists and Related Agents SA Form</a></p> <p>** See Cytokine and CAM Antagonists Appendix A for Clinical Criteria**</p> <p>For a list of Cytokine and CAM Antagonists and criteria for approval see <a href="#">Appendix A</a></p>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>Simponi<sup>®</sup></i> <i>Stelara<sup>®</sup> vial/syringe</i> <i>Taltz<sup>®</sup></i> <i>Tremfya<sup>™</sup></i> <i>Trexall<sup>®</sup></i> <i>Xatmep<sup>™</sup></i> <i>Xeljanz<sup>™</sup> &amp; Xeljanz<sup>™</sup> XR</i>	
<b>Ophthalmic</b>			
<b>Antibiotics</b>			
	<b>bacitracin/polymyxin b sulfate oint</b> <b>ciprofloxacin drops</b> <b>erythromycin</b> <b>gentamicin drops/ointment</b> <b>moxifloxacin drops (Vigamox generic)</b> <b>ofloxacin drops</b> <b>polymyxin/trimethoprim</b> <b>tobramycin</b>	<i>AzaSite<sup>™</sup> drops</i> <i>bacitracin</i> <i>Besivance<sup>®</sup> drops</i> <i>Bleph<sup>®</sup>-10</i> <i>Ciloxan<sup>®</sup> drops/ointment</i> <i>Garamycin<sup>®</sup> drops/ointment</i> <i>gatifloxacin 0.5% soln</i> <i>Ilotycin<sup>®</sup></i> <i>levofloxacin drops</i> <i>moxifloxacin drops (generic)</i> <i>Moxeza<sup>®</sup></i> <i>Moxeza<sup>®</sup> drops</i> <i>Natacyn<sup>®</sup></i> <i>neomycin/polymix/gramicidin</i> <i>neomycin/bacitracin/polymyxin oint</i> <i>Neosporin<sup>®</sup></i> <i>Ocuflox<sup>®</sup> s</i> <i>Polytrim<sup>®</sup></i> <i>sulfacetamide oint/ soln</i> <i>Tobrex<sup>®</sup> drops/ointment</i> <i>Vigamox<sup>®</sup></i> <i>Zymaxid<sup>®</sup></i>	<b><u>LENGTH OF AUTHORIZATIONS:</u></b> Date of service only; no refills  <b>Routine PDL edits</b>



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Preferred Agents		Non-Preferred Agents	SA Criteria
<b>Antibiotic/Steroid Combinations</b>			
neomycin/polymyxin/dexamethasone oint/susp sulfacetamide/prednisolone <b>Tobradex® oint/susp</b>	<i>Blephamide®</i> <i>Blephamide® S.O.P.</i> <i>Maxitrol® oint/susp</i> <i>neomycin/bacitracin/poly/HC</i> <i>neomycin/polymyxin/HC</i> <i>Pred-G® oint/susp</i> <i>Tobradex® ST</i> <b><i>tobramycin/dexamethasone susp</i></b> <i>Zylet®</i>	<b>LENGTH OF AUTHORIZATION:</b> Routine PDL edits	Date of service only; no refills
<b>Antihistamines/Mast Cell Stabilizers</b>			
<b>Antihistamines</b>			<b>LENGTH OF AUTHORIZATIONS:</b>
Alaway OTC® ketotifen fumarate olopatadine (generic Patanol & Pataday) Pazeo® Zaditor® OTC	<i>Bepreve®</i> <i>Elestat®</i> <i>epinastine 0.05% eye drops</i> <i>*Ilevro™ 0.3% (QL)</i> <i>Lastacaft®</i> <i>Optivar®</i> <i>Patanol® Rx and OTC</i> <i>Pataday® Rx and OTC</i> <i>Zerviate™</i>	Routine PDL edits *Ilevro™ is limited to 1 bottle plus 1 refill	1 year
<b>Mast Cell Stabilizers</b>			
cromolyn sodium	<i>Alocril®</i> <i>Alomide®</i>		
<b>Anti-inflammatory Agents</b>			
<b>NSAIDS</b>			<b>LENGTH OF AUTHORIZATIONS:</b>
diclofenac sodium flurbiprofen sodium ketorolac 0.4%& 0.5%	<i>Acular® 0.5% &amp; LS® 0.4%</i> <i>Acuvail®</i> <i>bromfenac 0.09%</i> <i>BromSite™</i> <i>*Ilevro™0.3% (QL)</i> <i>Inveltys™ (loteprednol etabonate)</i>	Routine PDL edits *Ilevro™ is limited to 1 bottle plus 1 refill	Date of service only; no refills



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>Nevanac<sup>®</sup></i> <i>Ocufen<sup>®</sup></i> <i>Prolensa<sup>™</sup></i>	
	<b>Corticosteroids</b>		
	<b>Durezol<sup>®</sup></b> <b>fluorometholone</b> <b>prednisolone acetate</b>	<i>Alrex<sup>™</sup></i> <i>Dexamethasone</i> <i>Flarex<sup>®</sup></i> <i>FML<sup>®</sup>, FML Forte<sup>®</sup> &amp; FML<sup>®</sup> S.O.P.</i> <i>loteprednol etabonate (generic for Lotemax<sup>™</sup>)</i> <i>Lotemax<sup>™</sup> drops/gel/oint</i> <i>Maxidex<sup>®</sup></i> <i>Omnipred<sup>®</sup></i> <i>Pred Forte<sup>®</sup> &amp; Pred Mild<sup>®</sup></i> <i>prednisolone sod phosphate</i> <i>Vexol<sup>®</sup></i> <i>Yutiq<sup>®</sup> (fluocinolone acetonide intravitreal implant)</i>	
<b>Glaucoma Agents</b>			
	<b>Alpha 2 Adrenergic Agents</b>		<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
	<b>Alphagan P<sup>®</sup> 0.1 &amp; 0.15%</b> <b>brimonidine 0.2%</b>	<i>apraclonidine 0.5% drops</i> <i>brimonidine tartrate 0.15%</i> <i>Iopidine<sup>®</sup> 0.5% &amp; 1%</i>	<b>Routine PDL edits</b>
	<b>Beta Blockers</b>		
	<b>carteolol 1%</b> <b>Combigan<sup>®</sup></b> <b>levobunolol 0.5%</b> <b>metipranolol 0.3%</b> <b>timolol maleate</b>	<i>Betagan<sup>®</sup> 0.5%</i> <i>betaxolol 0.5%</i> <i>Betoptic-S<sup>®</sup> 0.25%</i> <i>Istalol<sup>®</sup> 0.5%</i> <i>timolol (generic for Timoptic Ocudose)</i> <i>Timoptic<sup>®</sup> drops 0.25% &amp; 0.5%</i> <i>Timoptic<sup>®</sup> XE 0.25% &amp; 0.5% sol-gel</i>	





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<b>Carbonic Anhydrase Inhibitors</b>			
<b>Azopt® 1% dorzolamide dorzolamide/timolol</b>	<i>Cosopt® 0.5%-2% Cosopt® PF Simbrinza™ Trusopt® 2% dorzolamide/timolol PF</i>		
<b>Rho Kinase Inhibitor</b>			
<b>Rhopressa® Rocklatan</b>			
<b>Prostaglandin Analogs</b>			
<b>latanoprost Travatan Z®</b>	<i>bimatoprost Lumigan® 0.03% &amp; 0.01% Rescula® travoprost 0.004% Vyzulta™ Xalatan® 0.005% Xelpros® (latanoprost) Zioptan™</i>		
<b>Respiratory</b>			
<b>*Anti-Allergens, Oral</b>			
<b>Grass Pollen</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
	<i>*Oralair® SL</i>		
<b>Peanut</b>			<b>*All non-preferred Anti-Allergen drugs require the submission of a Clinical SA. Refer to <a href="#">Oralair SA Form</a> or <a href="#">Palforzia SA Form</a></b>
	<i>*Palforzia™</i>		
<b>Antihistamines: First and Second Generation</b>			
<b>First Generation Antihistamines</b>			<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<b>Generic only class</b>	<i>All Brands require a SA</i>		<b>Routine PDL edits</b>



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Preferred Agents	Non-Preferred Agents	SA Criteria																											
<b>Second Generation Antihistamines and Combinations</b>																													
cetirizine liquid 1mg/1mL (RX/OTC) cetirizine tabs OTC levocetirizine tab/OTC loratadine tab/syrup OTC	cetirizine chew tab (OTC) cetirizine liquid 5mg/5mL (OTC) cetirizine D tab (OTC) Clarinex® Clarinex-D® desloratadine ODT fexofenadine fexofenadine/PSE ER fexofenadine suspension loratadine ODT loratadine D 12 & 24 hr Semprex-D®	Endo has made a business decision to permanently discontinue Semprex-D. There are no generic version																											
<b>Beta-Adrenergic Agents</b>																													
<b>Long Acting Beta Adrenergic s (LABA) MDIs or Nebulizers</b>		<b>LENGTH OF AUTHORIZATIONS:</b> 1 year																											
Foradil® (AG) Serevent Diskus® (AG)	Arcapta DS(AG) Brovana® (AG) Perforomist® (AG) Striverdi® Respimat (AG)	<b>Routine PDL edits plus</b>  <b>Clinical Criteria for LABAs for Children</b> <b>LENGTH OF AUTHORIZATION:</b> 3 months Each drug listed below will require a SA for ages less than the FDA/PI indicated age. <table border="1" data-bbox="1075 1105 1999 1421"> <thead> <tr> <th>Brand Name</th> <th>Age where SA is required</th> <th>FDA Indications</th> </tr> </thead> <tbody> <tr> <td>*Advair® HFA</td> <td>Children &lt; 12</td> <td>Asthma &amp; COPD</td> </tr> <tr> <td>Advair® Diskus &amp; Wixela® 100/50</td> <td>Children &lt; 4</td> <td>Asthma &amp; COPD</td> </tr> <tr> <td>*Advair® Diskus &amp; Wixela® 250/50</td> <td>Children &lt; 12</td> <td>Asthma &amp; COPD</td> </tr> <tr> <td>Advair® Diskus &amp; Wixela® 500/50</td> <td>Children &lt; 12</td> <td>Asthma &amp; COPD</td> </tr> <tr> <td>Airduo™ Respieliek®</td> <td>Children &lt; 12</td> <td>Asthma only</td> </tr> <tr> <td>Anoro™ Ellipta</td> <td>Children &amp; Adolescents &lt; 18</td> <td>COPD only</td> </tr> <tr> <td>Arcapta® Neohaler</td> <td>Children &amp; Adolescents &lt; 18</td> <td>COPD only</td> </tr> <tr> <td>ArmonAir® Digihaler™</td> <td>Children &lt; 12</td> <td>Asthma only</td> </tr> </tbody> </table>	Brand Name	Age where SA is required	FDA Indications	*Advair® HFA	Children < 12	Asthma & COPD	Advair® Diskus & Wixela® 100/50	Children < 4	Asthma & COPD	*Advair® Diskus & Wixela® 250/50	Children < 12	Asthma & COPD	Advair® Diskus & Wixela® 500/50	Children < 12	Asthma & COPD	Airduo™ Respieliek®	Children < 12	Asthma only	Anoro™ Ellipta	Children & Adolescents < 18	COPD only	Arcapta® Neohaler	Children & Adolescents < 18	COPD only	ArmonAir® Digihaler™	Children < 12	Asthma only
Brand Name	Age where SA is required	FDA Indications																											
*Advair® HFA	Children < 12	Asthma & COPD																											
Advair® Diskus & Wixela® 100/50	Children < 4	Asthma & COPD																											
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Anoro™ Ellipta	Children & Adolescents < 18	COPD only																											
Arcapta® Neohaler	Children & Adolescents < 18	COPD only																											
ArmonAir® Digihaler™	Children < 12	Asthma only																											



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	Preferred Agents	Non-Preferred Agents	SA Criteria	
			Bevespi Aerosphere™	Children & Adolescents < 18 COPD only
			Breo® Ellipta™	Children & Adolescents < 18 Asthma & COPD
			Brovana®	Children & Adolescents < 18 COPD only
			*Dulera®	Children < 12 Asthma only
			Dupixent®	Children < 12 Asthma only
			fluticasone/salmeterol pow	Children < 12 Asthma only
			Foradil® Aerolizer	Children < 5 Asthma & COPD
			Perforomist®	Children & Adolescents < 18 COPD only
			Serevent® Diskus	Children < 4 Asthma & COPD
			Stiolto™ Respimat®	Children < 18 years COPD only
			Striverdi®-Respimat	Children < 18 years COPD only
			*Symbicort® 80/4.5	Children < 6 Asthma & COPD
			*Symbicort® 160/4.5	Children < 12 Asthma & COPD
	<b>Short Acting Metered Dose Inhalers or Devices</b>			
	Proair® HFA Ventolin® HFA	albuterol HFA (Proair®) albuterol HFA (Ventolin®) albuterol HFA (Proventil®) levalbuterol tartrate HFA ProAir® Digihaler™ ProAir® RespiClick Proventil® HFA Xopenex® HFA		
	<b>Short Acting Nebulizers</b>			
	albuterol sulfate (premixed)	levalbuterol soln Xopenex®		
	<b>Biologic: Human Monoclonal IgG4 Antibody Inhibits Interleukin-4 (IL-4) and Interleukin-13 (IL-13)</b>			
		Dupixent®	<b>LENGTH OF AUTHORIZATION:</b> 1 year Routine PDL edits plus	



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Preferred Agents	Non-Preferred Agents	SA Criteria
		<ul style="list-style-type: none"> <li>• Routine PDL</li> <li>• ≥ 12 years of age; AND</li> <li>• Diagnosis Moderate to severe Asthma with               <ul style="list-style-type: none"> <li>○ eosinophilic phenotype; OR</li> <li>○ Oral corticosteroid dependent; AND</li> </ul> </li> <li>• Prescribing provider is a pulmonologist or</li> <li>• An allergy/asthma specialist; AND</li> <li>• Have a diagnosis of step 5 or higher (moderate to severe) asthma; AND</li> <li>• Inadequately controlled asthma despite treatment with high dose inhaled or oral corticosteroid daily for at least 3 consecutive months; AND</li> <li>• Long-acting beta agonist (unless is not a candidate) daily for at least 3 consecutive months; AND</li> <li>• Is Add to current maintenance treatment; AND</li> <li>• Is not pregnant</li> </ul>
<b>COPD: Bronchodilators and Phosphodiesterase 4 (PDE4) Inhibitors <span style="color: red;">CLOSED CLASS</span></b>		
<b>Atrovent HFA®</b> <b>Anoro™ Ellipta® (AG)</b> <b>Bevespi Aerosphere™</b> <b>Combivent® Respimat</b> <b>ipratropium bromide soln</b> <b>ipratropium/albuterol nebs</b> <b>Spiriva®</b> <b>Stiolto Respimat™ (AG)</b>	<i>*Daliresp®</i> <i>Duaklir Pressair</i> <i>Incruse™ Ellipta®</i> <i>Lonhala™ Magnair™</i> <i>Spiriva® Respimat</i> <i>Tudorza™</i> <i>Yupelri™ (revefenacin)</i>	<u><b>LENGTH OF AUTHORIZATION:</b></u> 1 year  <b>Routine PDL edits plus</b>  <u><b>*Clinical Criteria for Daliresp®</b></u> <ul style="list-style-type: none"> <li>• If the member has a diagnosis of severe COPD associated with chronic bronchitis and a history of exacerbations; <b>AND</b></li> <li>• Trial/failure on at least one first-line or second-line agent (inhaled anticholinergics, long acting beta agonists or inhaled corticosteroids); <b>AND</b></li> <li>• Adjunctive therapy (Daliresp® must be used in conjunction with first-line or second-line agent).</li> </ul>
<b>Corticosteroids: Inhaled and Nasal Steroids</b>		
<b>Inhaled Corticosteroids: Combination Drugs (Glucocorticoid and Long Acting Beta Adrenergic) <span style="color: red;">CLOSED CLASS</span></b>		<u><b>LENGTH OF AUTHORIZATIONS:</b></u> 1 year
<b>Advair® Diskus (AG)</b> <b>Advair® HFA(AG)</b> <b>Asmanex Twisthaler</b> <b>Dulera® (AG)</b>	<b>AirDuo® Digihaler®</b> <i>Airduo™ Resplick® (AG)</i> <i>Breo® Ellipta™ (AG)</i> <i>Breztri Aerosphere™</i>	<b>Routine PDL edits</b>



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Preferred Agents		Non-Preferred Agents	SA Criteria
Symbicort® (AG)		<i>fluticasone/salmeterol (generic Airduo)</i> <i>fluticasone/salmeterol powder (generic Advair) (AG)</i> <i>Trelegy® Ellipta</i> <i>Wixela® (fluticasone/salmeterol) (AG /</i>	
<b>Inhaled Corticosteroids: Metered Dose Inhalers <span style="color: red;">CLOSED CLASS</span></b>			
<b>Flovent® Diskus &amp; HFA</b> <b>Pulmicort Flexhaler®</b>		<i>Alvesco®</i> <i>Aerospan™</i> <i>ArmonAir® Digihaler™(AG)</i> <i>Arnuity™ Ellipta®</i> <i>Asmanex HFA®</i> <i>QVAR® &amp; QVAR® Redihaler</i>	
<b>Inhaled Corticosteroids: Nebulizer Solution <span style="color: red;">CLOSED CLASS</span></b>			
<b>budesonide respules</b>		<i>Pulmicort® Respules</i>	<b>LENGTH OF AUTHORIZATIONS:</b> 1 year
<b>Nasal Steroids</b>			<b>Routine PDL edits</b>
<b>fluticasone Rx</b>		<i>Azelastine/fluticasone nasal spray (generic for Dymista®)</i> <i>Beconase AQ®</i> <i>budesonide (generic for Rhinocort® Aqua)</i> <i>budesonide (generic Rhinocort® Allergy OTC)</i> <i>Children's Qnasl™</i> <i>Clarispray OTC</i> <i>Dymista®</i> <i>Flonase®</i> <i>Flonase Sensimist (OTC)</i> <i>flunisolide</i> <i>fluticasone OTC</i>	



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>mometasone (generic Nasonex®)</i> <i>Nasonex®</i> <i>Omnaris®</i> <i>Qnasl™</i> <i>Rhinocort Aqua®</i> <i>Rhinocort® Allergy OTC</i> <i>Sinuva®</i> <i>Ticanase®</i> <i>triamcinolone OTC</i> <i>triamcinolone acetonide</i> <i>Veramyst®</i> <i>Xhance™</i> <i>Zetonna™</i>	
<b>*Cough and Cold Drug</b>			
	<b>Ala-Hist DM</b> <b>benzonatate cap</b> <b>codeine/ promethazine</b> <b>guaifenesin/codeine phosphate</b> <b>hydrocodone/ homatropine</b> <b>Iophen-C NR</b> <b>phenylephrine</b> <b>HCl/promethazine HCl</b> <b>promethazine DM syrup</b> <b>Tusnel® Pediatric Drops</b>	<i>lohist-DM syrup</i> <i>All other Legend cough and cold drugs are non-preferred</i> <i>Tessalon® perle</i>	<b><u>LENGTH OF AUTHORIZATION:</u></b> Date of Service Only  <b>Routine PDL edits</b>  <b>* Children under the age of 6 years are not eligible for cough and cold drugs.</b>
<b>Epinephrine, Self-Injected</b>			
	<b>epinephrine 0.15 mg &amp; 0.3 mg (authorized generic EpiPen® &amp; EpiPen® Jr)</b>	<i>Auvi-Q®</i> <i>Epipen®</i> <i>Epipen® Jr</i> <i>epinephrine 0.15mg &amp; 0.3mg (generic Adrenaclick)</i> <i>Symjepi™ (epinephrine)</i>	<b><u>LENGTH OF AUTHORIZATIONS:</u></b> 1 year  <b>Routine PDL edits</b>



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	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
<b>Intranasal Antihistamines</b>			
	azelastine 0.1%	<i>Astepro® 0.15%</i> <i>olopatadine</i> <i>Patanase®</i>	<b><u>LENGTH OF AUTHORIZATIONS:</u></b> 1 year  <b>Routine PDL edits</b>
<b>Leukotriene Receptor Antagonists</b>			
	montelukast tabs/chewable tabs	<i>Accolate®</i> <i>Singulair® tabs/chew tabs/granules</i> <i>montelukast granules</i> <i>zafirlukast</i> <i>Zyflo™</i> <i>Zyflo CR™</i> <i>zileuton ER</i>	<b><u>LENGTH OF AUTHORIZATIONS:</u></b> 1 year  <b>Routine PDL edits</b>