

# Indiana Medicaid Statewide Uniform Preferred Drug List (SUPDL)

## OptumRx Call Center

For prior authorization requests, claims processing issues or questions about the SUPDL, please contact OptumRx at 855-577-6317

Or fax the prior authorization requests to 855-577-6384

## Indiana Health Coverage Programs (IHCP) Drug Coverage

In accordance with 405 IAC 5-24, the IHCP covers all FDA-approved legend drugs with the exception of the following:

- Drugs designated by Centers for Medicare and Medicaid Services (CMS, formerly HCFA) as “less than effective” (DESI), or identical, related, or similar to a DESI drug
- Anorectics or any agent used to promote weight loss
- Topical minoxidil preparations
- Fertility enhancement drugs
- Drugs used primarily or solely for cosmetic purposes

**Note:** Inclusion of, or reference to, any given drug does not indicate market availability of the drug. Drugs that will be or have been withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

## Nomenclature

- **Statewide Uniform Preferred Drug List (SUPDL)** - a list of drugs within select therapeutic drug classes, developed and maintained by the Drug Utilization Review (DUR) Board, designated as *preferred* or *non-preferred* based upon clinical and financial considerations.
  - **Preferred Drug** – Covered drug designated by the DUR Board as a principle agent for use within a therapeutic class.
    - Mental health drugs are considered *preferred* (see Mental Health Drugs section below).
  - **Non-preferred Drug** – Covered drug designated by the DUR Board as secondary agent for use within a therapeutic class. Non-preferred drugs typically require prior authorization and history of trial and failure of (each of) the preferred agent(s), as confirmed by claims history, chart documentation, or provider attestation including dates of trial for each preferred agent (unless otherwise specified on the SUPDL).
    - Legacy continuation of therapy - The process whereby criteria are established exempting a drug from prior authorization, under specific conditions, when it would otherwise require prior authorization.
    - Brand name drugs, with an available substitutable generic, are *non-preferred* unless otherwise specified on the SUPDL. All preferred brand products on the SUPDL with a new available generic will list the generic agent added as non-preferred until it is financially advantageous to move to preferred. Once the generic agent is financially advantageous, it will replace the brand product as preferred. All non-preferred brand products on the SUPDL with a new available generic will list the generic agent added as non-preferred until it is reviewed by the Therapeutics Committee in the product’s regularly scheduled review cycle.

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

- Prior authorization is typically required for a prescriber's specification of "brand medically necessary".
  - Certain drugs, sometimes referred to as "narrow therapeutic index" drugs, are exempt from the requirement of prior authorization for "brand medically necessary"; see information in the Pharmacy Services Module found at this link: <https://www.in.gov/medicaid/files/pharmacy%20services.pdf>
- **Neutral Drug** – Covered drug that is in a therapeutic class not included on the SUPDL. As such, the drug has neither *preferred* nor *non-preferred* status.
  - **Line Extension Drug** – A new strength, formulation, or dosage form of a particular chemical entity for a given manufacturer that has been approved by the FDA. The SUPDL status of a line extension drug is the same as the status of the chemical entity to which it pertains unless otherwise determined by the DUR Board.
  - **Point of Sale Quick Check (PSQC)** – real-time automated prior authorization system that utilizes clinical prior authorization edits supported by a member's medical and pharmacy claims data. This process results in quicker PA determination for pharmacy claims processed by the fee-for-service (FFS) pharmacy benefit, with less intervention on the part of the pharmacy and prescribing providers.
  - **Status Pending Drug** – Covered drug that is subject to the SUPDL, but for which *preferred* or *non-preferred* status has yet to be assigned.

### Prior Authorization (PA)

This term is defined at 405 IAC 5-2-20. Any IHCP covered legend drug (including drugs that are or are not listed on the SUPDL) may require PA. Prior authorization is generally required in order to ensure appropriate drug utilization, conformance to established therapeutic guidelines, and fiscal reasonability.

Prior authorization request forms are located at <https://www.in.gov/medicaid/providers/index.html> under Pharmacy Services. Select "[PA Criteria and Administrative Forms](#)" under the "Quick Links" column on the right-hand margin. Drug specific PA criteria are attached to each associated drug class within the SUPDL document. Non-specific criteria are located at the end of the SUPDL document.

### Mental Health Drugs

In accordance with Indiana law, all antianxiety, antidepressant, antipsychotic, and "cross indicated" drugs are considered as being preferred. Drugs that are (1) classified in a central nervous system drug category or classification (according to *Drug Facts and Comparisons*) created after March 12, 2002, and (2) prescribed for the treatment of a mental illness (as defined by the most recent publication of the American Psychiatric Association's *Diagnostic and Statistical Manual of Mental Disorders*) are also considered as *preferred*. Please note that since these drugs/classes are *preferred*, they are not shown on the SUPDL document. **Lack of inclusion on the SUPDL does not mean these drugs are non-covered by the IHCP.** Click the following link for a list of utilization edits on mental health medications: [Utilization Edits for Mental Health Medications](#).

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

## Indiana Medicaid Statewide Uniform Preferred Drug List Table of Contents

Indiana Health Coverage Programs (IHCP) Drug Coverage.....1	Angiotensin Receptor Blockers ..... 12	DPP4 Inhibitors and Combination Agents ..... 28
Nomenclature.....1	Angiotensin Receptor Blocker Combinations ..... 13	GLP-1 Receptor Agonists and Combinations ..... 29
Prior Authorization (PA).....2	Beta Adrenergic Blockers ..... 13	Glucagon Agents ..... 29
Mental Health Drugs.....2	Beta Adrenergic Blockers with Diuretics ..... 13	Growth Hormones ..... 29
<b>ANTI-INFECTIVES ..... 5</b>	Calcium Channel Blockers ..... 14	Insulins – Intermediate Acting ..... 30
Antivirals – Anti-Herpetic ..... 5	Miscellaneous Cardiac Agents ..... 14	Insulins – Rapid Acting..... 30
Antivirals – Influenza ..... 5	<b>CNS AND OTHERS..... 15</b>	Insulins – Short Acting ..... 31
Cephalosporins – 3 <sup>rd</sup> Generation ..... 5	Agents for the Treatment of Opioid Use Disorder or Overdose ..... 15	Insulins – Long Acting ..... 31
Fluoroquinolones ..... 5	Antiemetic/Antivertigo Agents ..... 16	Miscellaneous Oral Antidiabetic Agents ..... 31
Hepatitis C Agents..... 5	Antiseizure Agents ..... 18	SGLT Inhibitors and Combinations..... 32
Macrolides ..... 6	Gastroprotective Agents ..... 20	Testosterones ..... 33
Ophthalmic Antibiotics ..... 6	Movement Disorder Agents ..... 20	Urea Cycle Disorders..... 33
Ophthalmic Antibiotics/ Corticosteroid Combinations ..... 7	Narcotic Antitussives and Combinations..... 20	<b>ESTROGEN AND RELATED AGENTS..... 34</b>
Otic Antibiotics..... 7	Narcotics..... 21	Estrogen and Related Agents ..... 34
Topical Antifungals..... 8	Skeletal Muscle Relaxants ..... 22	Contraceptives ..... 35
Topical Antivirals..... 8	Smoking Deterrent Agents ..... 23	<b>GASTROINTESTINAL AGENTS ..... 36</b>
Topical Antiviral and Anti-inflammatory Steroid Combinations ..... 8	<b>DERMATOLOGIC ..... 24</b>	Anti-ulcer Agents ..... 36
Vaginal Antimicrobials..... 9	Acne Agents..... 24	<i>H. Pylori</i> Agents ..... 36
<b>ANTIMIGRAINE ..... 10</b>	Antipsoriatics..... 25	H2 Receptor Antagonists..... 36
Antimigraine Preparations ..... 10	<b>ELECTROLYTE DEPLETERS ..... 26</b>	Laxatives and Cathartics ..... 37
<b>CARDIOVASCULAR..... 12</b>	Electrolyte Depleters ..... 26	Pancreatic Enzymes ..... 38
ACE Inhibitors ..... 12	<b>ENDOCRINE..... 27</b>	Proton Pump Inhibitors ..... 38
ACE Inhibitor Combinations ..... 12	Anaphylaxis Agents..... 27	Ulcerative Colitis Agents..... 39
	Bone Formation Stimulating Agents..... 27	<b>GENITOURINARY ..... 40</b>
	Bone Resorption Inhibitors ..... 27	BPH Agents ..... 40

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

Urinary Tract Antispasmodic/Anti-Incontinence Agents .....	41	Beta Agonists – Short Acting.....	49	Topical Post-Herpetic Neuralgia Agents.....	55
<b>HEMATOLOGIC.....</b>	<b>42</b>	Bronchodilator Agents-Beta Adrenergic and Anticholinergic Combinations .....	49	<b>MISCELLANEOUS INFORMATION .....</b>	<b>56</b>
Direct Oral Anticoagulants .....	42	Nasal Antihistamines/Nasal Anti-Inflammatory Steroids.....	50		
Hematinics.....	42	Oral Inhaled Glucocorticoids.....	51		
Leukocyte Stimulants.....	43	Pulmonary Antihypertensives .....	51		
Platelet Aggregation Inhibitors .....	43	Respiratory and Allergy Biologics .....	51		
<b>LIPOTROPICS.....</b>	<b>44</b>	<b>TARGETED IMMUNOMODULATORS .....</b>	<b>52</b>		
Bile Acid Sequestrants .....	44	Targeted Immunomodulators .....	52		
Fibric Acid Derivatives .....	44	<b>TOPICAL AGENTS .....</b>	<b>53</b>		
HMG CoA Reductase Inhibitors.....	44	Dry Eye Disease or Keratoconjunctivitis .....	53		
Lipotropics .....	45	Miotics-Intraocular Pressure Reducers .....	53		
<b>MULTIPLE SCLEROSIS AGENTS.....</b>	<b>46</b>	Ophthalmic Antihistamines.....	54		
Multiple Sclerosis Agents .....	46	Ophthalmic Anti-Inflammatory Agents .....	54		
<b>RESPIRATORY.....</b>	<b>47</b>	Ophthalmic Mast Cell Stabilizers .....	54		
Antihistamine-Decongestant Combinations/2 <sup>nd</sup> Generation Antihistamines .....	47	Otic Preparations .....	54		
Antiviral Monoclonal Antibody.....	48	Topical Anti-Inflammatory Agents – NSAIDS .....	55		
Beta Adrenergics and Corticosteroids.....	48	Topical Antiparasitics.....	55		
Beta Agonists – Long Acting.....	48	Topical Immunomodulators .....	55		

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTI-INFECTIVES</b>			
<b>Antivirals – Anti-Herpetic</b>	<ul style="list-style-type: none"> <li>acyclovir</li> <li>valacyclovir               <ul style="list-style-type: none"> <li>ST – must have diagnosis of HIV or trial and failure of acyclovir or medical justification for use over acyclovir</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>famciclovir</li> <li>Sitavig</li> </ul>	
<b>Antivirals – Influenza</b>	<ul style="list-style-type: none"> <li>amantadine</li> <li>oseltamivir</li> <li>Relenza</li> <li>rimantadine               <ul style="list-style-type: none"> <li>AGE – 60 years and older</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>rimantadine               <ul style="list-style-type: none"> <li>AGE – under 60 years old</li> </ul> </li> <li>Rapivab</li> <li>Xofluza</li> </ul>	
<b>Cephalosporins – 3<sup>rd</sup> Generation</b>	<ul style="list-style-type: none"> <li>cefdinir</li> <li>cefepodoxime</li> </ul>	<ul style="list-style-type: none"> <li>cefixime caps, susp</li> <li>Suprax chew, susp</li> </ul>	
<b>Fluoroquinolones</b> <b>*Note: All fluoroquinolones will be limited to 14 days per claim*</b>	<ul style="list-style-type: none"> <li>ciprofloxacin</li> <li>levofloxacin</li> <li>moxifloxacin</li> </ul>	<ul style="list-style-type: none"> <li>Baxdela</li> <li>ofloxacin</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>Cipro suspension</li> <li>ciprofloxacin suspension</li> <li>levofloxacin solution</li> </ul>	<a href="#">PA Criteria for ciprofloxacin and levofloxacin solution</a>
<b>Hepatitis C Agents</b>	<ul style="list-style-type: none"> <li>Pegasys</li> <li>ribavirin</li> </ul> <p>PA criteria must be met for the following (<b>note: treatment naïve patients must only meet age and quantity limits</b>):</p> <ul style="list-style-type: none"> <li>Epclusa 200-50mg</li> <li>Epclusa 150-37.5mg</li> <li>Mavyret</li> <li>sofosbuvir/velpatasvir 400-100mg</li> <li>Zepatier</li> </ul>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>Epclusa 400-100mg</li> <li>Harvoni</li> <li>ledipasvir/sofosbuvir</li> <li>Sovaldi</li> <li>Viekira</li> <li>Vosevi</li> </ul>	<a href="#">Hepatitis C Agents PA Criteria</a>  <a href="#">Hepatitis C Agents PA Form</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTI-INFECTIVES - Continued</b>			
<b>Macrolides</b>	<ul style="list-style-type: none"> <li>• azithromycin suspension</li> <li>• azithromycin 600 mg oral tablets               <ul style="list-style-type: none"> <li>○ QL – 1 tablet/day</li> </ul> </li> <li>• azithromycin 500 mg oral tablets               <ul style="list-style-type: none"> <li>○ QL – 7 tablets/30 days</li> </ul> </li> <li>• azithromycin 250 mg oral tablets               <ul style="list-style-type: none"> <li>○ QL – 6 tablets/30 days</li> </ul> </li> <li>• clarithromycin</li> <li>• erythromycin capsules</li> <li>• erythromycin ethylsuccinate susp               <ul style="list-style-type: none"> <li>○ ST – must be under 12 years of age or unable to swallow tablets/capsules</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• E.E.S. Granules               <ul style="list-style-type: none"> <li>○ ST – must have tried and failed erythromycin ethylsuccinate suspension in the past 90 days OR member must be under 12 years of age or unable to swallow tablets/capsules and prescriber has provided valid medical justification for the use of E.E.S. Granules over preferred agents</li> </ul> </li> <li>• E.E.S. tablets</li> <li>• erythrocin stearate</li> <li>• erythromycin tablets</li> <li>• erythromycin tablets EC</li> <li>• Zmax</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Dificid</li> </ul>	<a href="#">Dificid PA Criteria</a>  <a href="#">Dificid PA Form</a>
<b>Ophthalmic Antibiotics</b>	<ul style="list-style-type: none"> <li>• Besivance susp</li> <li>• Ciloxan oint</li> <li>• ciprofloxacin soln</li> <li>• erythromycin oint</li> <li>• Gentak ointment</li> <li>• gentamicin soln</li> <li>• moxifloxacin soln               <ul style="list-style-type: none"> <li>○ AGE – 30 years of age or older; ST – patients under 30 years of age must have tried at least one preferred agent other than moxifloxacin within the past 30 days</li> </ul> </li> <li>• neomycin/polymyxin B/gramicidin soln</li> <li>• ofloxacin soln</li> <li>• polymyxin B/bacitracin oint</li> <li>• polymyxin B/trimethoprim soln</li> <li>• sulfacet sod oint</li> <li>• tobramycin soln</li> </ul>	<ul style="list-style-type: none"> <li>• Azasite soln</li> <li>• bacitracin oint</li> <li>• gatifloxacin soln</li> <li>• levofloxacin soln</li> <li>• Natacyn</li> <li>• neomycin/bacitracin/polymyxin oint</li> <li>• Tobrex oint</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTI-INFECTIVES - Continued</b>			
<b>Ophthalmic Antibiotics/ Corticosteroid Combinations</b>	<ul style="list-style-type: none"> <li>• neomycin/polymyxin B/dexamethasone oint</li> <li>• neomycin/polymyxin B/dexamethasone susp</li> <li>• sulfacetamide sodium/prednisolone soln</li> <li>• Tobradex oint</li> <li>• Tobradex ST susp</li> <li>• tobramycin/dexamethasone susp</li> <li>• Zylet susp</li> </ul>	<ul style="list-style-type: none"> <li>• Blephamide S.O.P. oint</li> <li>• neomycin/polymyxin/bacitracin/hc oint</li> <li>• neomycin/polymyxin/hc susp</li> <li>• Pred-G susp</li> <li>• Pred-G S.O.P. oint</li> </ul>	
<b>Otic Antibiotics</b>	<ul style="list-style-type: none"> <li>• ofloxacin otic soln</li> </ul> <p><b>Antibiotic/Steroid Combinations</b></p> <ul style="list-style-type: none"> <li>• ciprofloxacin-dexamethasone otic susp</li> <li>• Cipro HC susp</li> <li>• Cortisporin TC otic susp</li> <li>• neomycin/polymyxin B/hydrocortisone otic soln</li> <li>• neomycin/polymyxin B/hydrocortisone otic susp</li> </ul>	<ul style="list-style-type: none"> <li>• ciprofloxacin soln</li> <li>• Otiprio</li> </ul> <p><b>Antibiotic/Steroid Combinations</b></p> <ul style="list-style-type: none"> <li>• ciprofloxacin-fluocinolone PF otic susp</li> </ul>	
<b>Systemic Antifungals</b>	<ul style="list-style-type: none"> <li>• fluconazole 50 mg tabs <ul style="list-style-type: none"> <li>○ QL – 3 tabs/30 days</li> </ul> </li> <li>• fluconazole 150 mg tabs <ul style="list-style-type: none"> <li>○ QL – 4 tabs/30 days</li> </ul> </li> <li>• fluconazole suspension</li> <li>• itraconazole</li> <li>• ketoconazole</li> <li>• terbinafine</li> </ul>	<ul style="list-style-type: none"> <li>• Cresemba</li> <li>• itraconazole solution <ul style="list-style-type: none"> <li>○ ST – must be 12 years of age and under or unable to swallow capsules/tablets</li> </ul> </li> <li>• Noxafil PAK <ul style="list-style-type: none"> <li>○ ST – must be 2 years of age or older and less than 13 years of age</li> </ul> </li> <li>• posaconazole tablet &amp; 200 mg/5 mL suspension <ul style="list-style-type: none"> <li>○ ST – must have tried fluconazole for treatment of oropharyngeal candidiasis or must be severely immunocompromised and need prophylaxis against invasive Aspergillus or Candida infections</li> </ul> </li> <li>• Tolsura</li> <li>• voriconazole suspension <ul style="list-style-type: none"> <li>○ ST – must be 12 years of age and under or unable to swallow capsules/tablets</li> </ul> </li> <li>• voriconazole tabs</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Brexafemme</li> <li>• Vivjoa</li> </ul>	<p><a href="#">Antimicrobials for Treatment of Vaginal Infections PA Criteria</a></p> <p><a href="#">Antimicrobials for Treatment of Vaginal Infections PA form</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTI-INFECTIVES - Continued</b>			
<b>Topical Antifungals</b>	<p><i>All generics unless otherwise specified</i></p> <ul style="list-style-type: none"> <li>• ciclopirox (cream &amp; topical solution)</li> <li>• clotrimazole</li> <li>• Jublia</li> <li>• miconazole</li> <li>• terbinafine 1% cream</li> <li>• tolnaftate 1% cream, powder, spray</li> </ul>	<ul style="list-style-type: none"> <li>• ciclopirox gel, kit, topical shampoo, topical suspension</li> <li>• econazole</li> <li>• Ertaczo</li> <li>• Extina</li> <li>• ketoconazole topical foam</li> <li>• Loprox kit</li> <li>• luliconazole</li> <li>• Luzu</li> <li>• Mentax</li> <li>• miconazole/zinc/pet oint</li> <li>• naftifine 1% cream</li> <li>• naftifine 2% cream, gel</li> <li>• Naftin 1% gel</li> <li>• Oxistat</li> <li>• tavaborole solution</li> <li>• Vusion</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• sulconazole cream, solution</li> </ul>	
<b>Topical Antivirals</b>	<ul style="list-style-type: none"> <li>• Zovirax cream</li> </ul>	<ul style="list-style-type: none"> <li>• acyclovir ointment</li> <li>• Denavir cream</li> <li>• docosanol OTC cream</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• acyclovir cream</li> </ul>	
<b>Topical Antiviral and Anti-inflammatory Steroid Combinations</b>	<ul style="list-style-type: none"> <li>• Xerese <ul style="list-style-type: none"> <li>○ QL – 1 tube per claim per 90 days</li> </ul> </li> </ul>	N/A	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTI-INFECTIVES - Continued</b>			
<b>Vaginal Antimicrobials</b>	<p><b>Antibacterials</b></p> <ul style="list-style-type: none"> <li>• Cleocin 2% cream</li> <li>• metronidazole vaginal gel</li> <li>• Nuversa</li> <li>• Solosec</li> </ul> <p><b>Antifungals</b></p> <ul style="list-style-type: none"> <li>• clotrimazole OTC <ul style="list-style-type: none"> <li>○ QL – 2 treatment courses/month</li> </ul> </li> <li>• miconazole cream OTC <ul style="list-style-type: none"> <li>○ QL – 2 treatment courses/month</li> </ul> </li> <li>• tioconazole OTC <ul style="list-style-type: none"> <li>○ QL – 2 treatment courses/month</li> </ul> </li> <li>• terconazole cream</li> </ul>	<p><b>Antibacterials</b></p> <ul style="list-style-type: none"> <li>• Cleocin Ovules</li> <li>• Clindesse</li> <li>• Vandazole</li> <li>• Xaciato</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• clindamycin 2% cream</li> </ul> <p><b>Antifungals</b></p> <ul style="list-style-type: none"> <li>• Gynazole-1</li> <li>• miconazole combination pack OTC <ul style="list-style-type: none"> <li>○ QL – 2 treatment courses/month</li> </ul> </li> <li>• miconazole suppositories OTC <ul style="list-style-type: none"> <li>○ QL – 2 treatment courses/month</li> </ul> </li> <li>• miconazole suppositories (Rx)</li> <li>• terconazole suppositories</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTIMIGRAINE</b>			
<b>Antimigraine Preparations</b>	<ul style="list-style-type: none"> <li>• rizatriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 12 tabs/30 days</li> </ul> </li> <li>• rizatriptan ODT <ul style="list-style-type: none"> <li>○ QL – 1 box – 12 tabs/30 days</li> </ul> </li> <li>• sumatriptan nasal spray <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 inhalers/30 days</li> </ul> </li> <li>• sumatriptan tablets <ul style="list-style-type: none"> <li>○ QL – 1 box – 9 tabs/30 days</li> </ul> </li> <li>• sumatriptan stat dose or stat dose refill package <ul style="list-style-type: none"> <li>○ QL – 1 box – 2 injections/30 days</li> </ul> </li> <li>• sumatriptan vial <ul style="list-style-type: none"> <li>○ QL – 2 vials – 2 injections/30 days</li> </ul> </li> </ul> <p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Elyxyb <ul style="list-style-type: none"> <li>○ QL – 6 bottles/30 days</li> </ul> </li> <li>• Nurtec ODT <ul style="list-style-type: none"> <li>○ QL – 8 tabs/30 days for acute treatment;</li> <li>○ QL – 16 tabs/30 days for preventative treatment</li> </ul> </li> <li>• Ubrelvy <ul style="list-style-type: none"> <li>○ QL – 10 tabs/20 days</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• almotriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 tabs/30 days</li> </ul> </li> <li>• frovatriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 9 tabs/30 days</li> </ul> </li> <li>• naratriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 9 tabs/30 days</li> </ul> </li> <li>• Onzetra Xsail <ul style="list-style-type: none"> <li>○ QL – 1 box (8 pouches)/30 days</li> </ul> </li> <li>• Relpax <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 tabs/30 days</li> </ul> </li> <li>• sumatriptan/naproxen <ul style="list-style-type: none"> <li>○ QL – 1 box – 9 tabs/30 days</li> </ul> </li> <li>• Tosymra Solution</li> <li>• Treximet <ul style="list-style-type: none"> <li>○ QL – 1 box – 9 tabs/30 days</li> </ul> </li> <li>• Zembrace SymTouch <ul style="list-style-type: none"> <li>○ QL – 1 box (4 injections)/30 days</li> </ul> </li> <li>• zolmitriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 tabs/30 days</li> </ul> </li> <li>• zolmitriptan nasal spray <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 inhalers/30 days</li> </ul> </li> <li>• zolmitriptan ODT <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 tabs/30 days</li> </ul> </li> </ul> <p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Reyvow <ul style="list-style-type: none"> <li>○ QL – 50 mg dose – 4 (50 mg) tabs/30 days</li> <li>○ QL – 100 mg dose – 4 (100 mg) tabs/30 days</li> <li>○ QL – 200 mg dose – 8 (100 mg) tabs/30 days</li> </ul> </li> <li>• Zavzpret <ul style="list-style-type: none"> <li>○ QL – 6 devices/22 days</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• eletriptan <ul style="list-style-type: none"> <li>○ QL – 1 box – 6 tabs/30 days</li> </ul> </li> </ul>	<a href="#">Antimigraine PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ANTIMIGRAINE - Continued</b>			
<b>Antimigraine Preparations - Continued</b>	<b>Prophylaxis</b> PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Ajovy               <ul style="list-style-type: none"> <li>○ QL – 225 mg/month or 675 mg/3 months</li> </ul> </li> <li>• Emgality               <ul style="list-style-type: none"> <li>○ QL migraine – 240 mg loading dose; then 120 mg/month</li> <li>○ QL cluster headache – 300mg at start of headache and once monthly thereafter until end of headache</li> </ul> </li> <li>• Qulipta               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> </ul>	<b>Prophylaxis</b> PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Aimovig               <ul style="list-style-type: none"> <li>○ QL – 140 mg/month</li> </ul> </li> <li>• Vyepti               <ul style="list-style-type: none"> <li>○ QL – 3 mL/90 days</li> </ul> </li> </ul>	<a href="#">Antimigraine PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CARDIOVASCULAR</b>			
<b>ACE Inhibitors</b>	<ul style="list-style-type: none"> <li>• benazepril</li> <li>• enalapril</li> <li>• fosinopril</li> <li>• lisinopril</li> <li>• quinapril</li> <li>• ramipril</li> </ul>	<ul style="list-style-type: none"> <li>• captopril</li> <li>• enalapril 1 mg/mL solution               <ul style="list-style-type: none"> <li>○ ST – must be under 12 years of age or unable to swallow tablets</li> </ul> </li> <li>• moexipril</li> <li>• perindopril</li> <li>• Qbrelis               <ul style="list-style-type: none"> <li>○ ST – must be 6 years of age or older and less than 12 years of age OR 12 years of age and older AND unable to swallow tablets</li> </ul> </li> <li>• trandolapril</li> </ul>	
<b>ACE Inhibitor Combinations</b>	<p><b><i>ACE Inhibitors with Calcium Channel Blockers</i></b></p> <ul style="list-style-type: none"> <li>• amlodipine/benazepril               <ul style="list-style-type: none"> <li>○ QL – 30 caps/30 days</li> </ul> </li> </ul> <p><b><i>ACE Inhibitors with Diuretics</i></b></p> <ul style="list-style-type: none"> <li>• benazepril/HCTZ</li> <li>• enalapril/HCTZ</li> <li>• lisinopril/HCTZ</li> <li>• quinapril/HCTZ</li> </ul>	<p><b><i>ACE Inhibitors with Calcium Channel Blockers</i></b></p> <ul style="list-style-type: none"> <li>• trandolapril/verapamil               <ul style="list-style-type: none"> <li>○ QL – 30 caps/30 days</li> </ul> </li> </ul> <p><b><i>ACE Inhibitors with Diuretics</i></b></p> <ul style="list-style-type: none"> <li>• fosinopril/HCTZ</li> </ul>	
<b>Angiotensin Receptor Blockers</b>	<ul style="list-style-type: none"> <li>• Edarbi               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• irbesartan               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• losartan 25 mg, 50 mg               <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day</li> </ul> </li> <li>• losartan 100 mg               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• olmesartan 5 mg               <ul style="list-style-type: none"> <li>○ QL – 3 tabs/day</li> </ul> </li> <li>• olmesartan 20 mg, 40 mg               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• telmisartan               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• valsartan 40 mg, 80 mg, 160 mg               <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day</li> </ul> </li> <li>• valsartan 320 mg               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• candesartan 4 mg, 8 mg, 16 mg               <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day</li> </ul> </li> <li>• candesartan 32 mg               <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• valsartan solution               <ul style="list-style-type: none"> <li>○ ST – must be unable to swallow tablets</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CARDIOVASCULAR - Continued</b>			
<b>Angiotensin Receptor Blocker Combinations</b>	<p><b>Angiotensin Receptor Blockers with Diuretics</b></p> <ul style="list-style-type: none"> <li>• Edarbyclor</li> <li>• losartan/HCTZ</li> <li>• valsartan/HCTZ</li> </ul> <p><b>Angiotensin Receptor Blockers with Calcium Channel Blockers</b> N/A</p> <p><b>Angiotensin Receptor Blockers with Calcium Channel Blockers and Diuretics</b> N/A</p>	<p><b>Angiotensin Receptor Blockers with Diuretics</b></p> <ul style="list-style-type: none"> <li>• candesartan/HCTZ</li> <li>• irbesartan/HCTZ</li> <li>• olmesartan/HCTZ</li> <li>• telmisartan/HCTZ</li> </ul> <p><b>Angiotensin Receptor Blockers with Calcium Channel Blockers</b></p> <ul style="list-style-type: none"> <li>• olmesartan/amlodipine <ul style="list-style-type: none"> <li>○ ST – trial and failure of individual components</li> </ul> </li> <li>• telmisartan/amlodipine <ul style="list-style-type: none"> <li>○ ST – trial and failure of individual components</li> </ul> </li> <li>• valsartan/amlodipine <ul style="list-style-type: none"> <li>○ ST – trial and failure of individual components</li> </ul> </li> </ul> <p><b>Angiotensin Receptor Blockers with Calcium Channel Blockers and Diuretics</b></p> <ul style="list-style-type: none"> <li>• amlodipine/olmesartan/HCTZ <ul style="list-style-type: none"> <li>○ ST – trial and failure of individual components</li> </ul> </li> <li>• amlodipine/valsartan/HCTZ <ul style="list-style-type: none"> <li>○ ST – trial and failure of individual components</li> </ul> </li> </ul>	
<b>Beta Adrenergic Blockers</b>	<ul style="list-style-type: none"> <li>• acebutolol</li> <li>• atenolol</li> <li>• bisoprolol</li> <li>• carvedilol</li> <li>• labetalol</li> <li>• metoprolol</li> <li>• metoprolol succinate ER</li> <li>• nebivolol</li> <li>• propranolol</li> <li>• propranolol ER caps</li> <li>• sotalol</li> </ul>	<ul style="list-style-type: none"> <li>• betaxolol</li> <li>• carvedilol ER cap <ul style="list-style-type: none"> <li>○ QL – 1 cap/day</li> </ul> </li> <li>• Hemangeol solution <ul style="list-style-type: none"> <li>○ ST – must be 5 weeks of age or older and less than or equal to 1 year of age</li> </ul> </li> <li>• Kapspargo</li> <li>• nadolol</li> <li>• pindolol</li> <li>• Sotylize oral solution <ul style="list-style-type: none"> <li>○ ST – must be under 12 years of age or unable to swallow capsules/tablets</li> </ul> </li> <li>• timolol</li> </ul>	
<b>Beta Adrenergic Blockers with Diuretics</b>	<ul style="list-style-type: none"> <li>• atenolol/chlorthalidone</li> <li>• bisoprolol/HCTZ</li> </ul>	<ul style="list-style-type: none"> <li>• metoprolol/HCTZ</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CARDIOVASCULAR - Continued</b>			
<b>Calcium Channel Blockers</b>	<p><b><i>Dihydropyridine</i></b></p> <ul style="list-style-type: none"> <li>• amlodipine</li> <li>• felodipine ER</li> <li>• nifedipine (short-acting)</li> <li>• nifedipine ER</li> </ul> <p><b><i>Non-Dihydropyridine</i></b></p> <ul style="list-style-type: none"> <li>• Calan SR</li> <li>• Cardizem LA</li> <li>• diltiazem (long-acting formulations)</li> <li>• diltiazem (non-time released)</li> <li>• nimodipine</li> <li>• verapamil (long-acting formulations)</li> <li>• verapamil (non-time released)</li> </ul> <p><b><i>Liquid Formulation</i></b></p> <ul style="list-style-type: none"> <li>• Norliqva <ul style="list-style-type: none"> <li>○ ST – must be unable to swallow tablets</li> </ul> </li> </ul> <p><b><i>Combinations</i></b> N/A</p>	<p><b><i>Dihydropyridine</i></b></p> <ul style="list-style-type: none"> <li>• isradipine (non-time released)</li> <li>• levamlodipine</li> <li>• nicardipine (non-time released)</li> <li>• nisoldipine</li> </ul> <p><b><i>Non-Dihydropyridine</i></b></p> <ul style="list-style-type: none"> <li>• Cardizem CD</li> <li>• Matzim LA</li> <li>• verapamil ER PM</li> <li>• Verelan PM</li> </ul> <p><b><i>Liquid Formulation</i></b></p> <ul style="list-style-type: none"> <li>• Katerzia <ul style="list-style-type: none"> <li>○ ST – must be 6 years of age or older and less than 12 years of age OR unable to swallow tablets AND previous trial and failure of Norliqva OR medical rationale for use</li> </ul> </li> <li>• Nymalize <ul style="list-style-type: none"> <li>○ ST – must be 18 years of age or older AND unable to swallow capsule formulation</li> </ul> </li> </ul> <p><b><i>Combinations</i></b></p> <ul style="list-style-type: none"> <li>• amlodipine/atorvastatin <ul style="list-style-type: none"> <li>○ ST – prescriber must provide documentation that separate components are not suitable for use</li> </ul> </li> </ul>	
<b>Miscellaneous Cardiac Agents</b>	PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• ivabradine</li> <li>• Entresto</li> </ul>	PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Camzyos</li> <li>• Verquvo</li> </ul>	<a href="#">Cardiac Agents PA Criteria</a>  <a href="#">Cardiac Agents PA Form</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS</b>			
<b>Agents for the Treatment of Opioid Use Disorder or Overdose</b>	<p><b>Agents for Opioid Use Disorder – oral</b></p> <ul style="list-style-type: none"> <li>• buprenorphine sublingual tablets <ul style="list-style-type: none"> <li>○ AGE – 16 years of age and older</li> <li>○ QL – 24mg/day</li> </ul> </li> <li>• buprenorphine/naloxone sublingual tablets <ul style="list-style-type: none"> <li>○ AGE – 16 years of age and older</li> <li>○ QL – 24mg/day</li> </ul> </li> <li>• Suboxone Film <ul style="list-style-type: none"> <li>○ AGE – 16 years of age and older</li> <li>○ QL – 24mg/day</li> </ul> </li> <li>• Zubsolv <ul style="list-style-type: none"> <li>○ AGE – 16 years of age and older</li> <li>○ QL – 17.2mg/day</li> </ul> </li> </ul> <p><b>Agents for Opioid Use Disorder – injectable</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Sublocade</li> </ul> <p><b>Agents for Opioid Overdose</b></p> <ul style="list-style-type: none"> <li>• Kloxxado</li> <li>• nalmefene</li> <li>• naloxone injection</li> <li>• naloxone nasal spray</li> <li>• Narcan Nasal</li> <li>• Opvee</li> <li>• Zimhi</li> </ul>	<p><b>Agents for Opioid Use Disorder – oral</b> Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• buprenorphine/naloxone sublingual films <ul style="list-style-type: none"> <li>○ AGE – 16 years of age and older</li> <li>○ QL – 24mg/day</li> </ul> </li> </ul> <p><b>Agents for Opioid Use Disorder – injectable</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Brixadi</li> </ul> <p><b>Agents for Opioid Overdose</b> N/A</p>	<p><a href="#">Opioid Use Disorder Treatments</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - continued</b>			
<b>Antiemetic/Antivertigo Agents</b>	<p><b><i>Appetite Stimulant</i></b> N/A</p> <p><b><i>H1 Antagonist/Vitamin</i></b></p> <ul style="list-style-type: none"> <li>• Diclegis <ul style="list-style-type: none"> <li>○ QL – 4 tabs/day; Max 270/365 days</li> </ul> </li> </ul> <p><b><i>Selective 5-HT3 Receptor Antagonist</i></b></p> <ul style="list-style-type: none"> <li>• ondansetron oral tablets &amp; disintegrating tablets <ul style="list-style-type: none"> <li>○ QL – 90 tabs/30 days</li> </ul> </li> <li>• ondansetron oral solution <ul style="list-style-type: none"> <li>○ QL – 1 bottle/Rx</li> </ul> </li> <li>• ondansetron solution for injection</li> </ul>	<p><b><i>Appetite Stimulant</i></b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• dronabinol</li> </ul> <p><b><i>H1 Antagonist/Vitamin</i></b></p> <ul style="list-style-type: none"> <li>• Bonjesta <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day; Max 270/365 days</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• doxylamine/pyridoxine oral tabs <ul style="list-style-type: none"> <li>○ QL – 4 tabs/day; Max 270/365 days</li> </ul> </li> </ul> <p><b><i>Selective 5-HT3 Receptor Antagonist</i></b></p> <ul style="list-style-type: none"> <li>• Anzemet oral tabs <ul style="list-style-type: none"> <li>○ QL – 10 units/Rx</li> </ul> </li> <li>• granisetron oral tablets</li> <li>• granisetron solution for injection</li> <li>• palonosetron injection <ul style="list-style-type: none"> <li>○ QL – 1 vial/Rx</li> </ul> </li> <li>• Sancuso transdermal system <ul style="list-style-type: none"> <li>○ ST – physician documentation required indicating oral medications are unsuitable for patient use</li> </ul> </li> <li>• Sustol</li> </ul>	<a href="#">Dronabinol PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - continued</b>			
<b>Antiemetic/Antivertigo Agents - continued</b>	<p><b><i>Substance P-Neurokinin 1 Receptor Antagonist</i></b></p> <ul style="list-style-type: none"> <li>• aprepitant 40 mg and 80 mg oral capsules <ul style="list-style-type: none"> <li>○ QL – 6 caps/Rx</li> </ul> </li> <li>• Emend Tripack <ul style="list-style-type: none"> <li>○ QL – 2 packs (6 caps)/Rx</li> </ul> </li> <li>• fosaprepitant vials <ul style="list-style-type: none"> <li>○ QL – 2 vials/Rx</li> </ul> </li> </ul> <p><b><i>Substance P-NK 1 Antagonist/Selective 5-HT3 Antagonist</i></b></p> <p>N/A</p>	<p><b><i>Substance P-Neurokinin 1 Receptor Antagonist</i></b></p> <ul style="list-style-type: none"> <li>• aprepitant 125 mg oral capsules <ul style="list-style-type: none"> <li>○ QL – 6 caps/Rx</li> </ul> </li> <li>• Cinvanti injection <ul style="list-style-type: none"> <li>○ QL – 2 vials/Rx</li> </ul> </li> <li>• Emend IV solution <ul style="list-style-type: none"> <li>○ QL – 2 vials/Rx</li> </ul> </li> <li>• Emend suspension <ul style="list-style-type: none"> <li>○ ST – must have tried Emend oral capsules or have inability to swallow or tolerate the capsule formulation</li> <li>○ QL – 3 packets /Rx</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• aprepitant 80 mg/125 mg tripack <ul style="list-style-type: none"> <li>○ QL – 2 packs (6 caps)/Rx</li> </ul> </li> </ul> <p><b><i>Substance P-NK 1 Antagonist/Selective 5-HT3 Antagonist</i></b></p> <ul style="list-style-type: none"> <li>• Akynzeo <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes or provide medical justification for use of the combination product</li> </ul> </li> </ul>	<a href="#">Dronabinol PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED		NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS – continued</b>				
<b>Antiseizure Agents</b>  <i>Note: Utilization Edits may apply for mental health medications; see Utilization Edits for Mental Health Medications for associated quantity limits</i>	<ul style="list-style-type: none"> <li>• carbamazepine IR, ER cap, chew</li> <li>• Carbatrol</li> <li>• Celontin</li> <li>• clobazam tab, susp</li> <li>• Depakote Sprinkle</li> <li>• Diastat rectal</li> <li>• diazepam rectal gel</li> <li>• Dilantin cap, chew, susp</li> <li>• divalproex DR, ER, sprinkle cap</li> <li>• Epitol</li> <li>• ethosuximide cap, soln</li> <li>• felbamate susp</li> <li>• Felbatol tablets</li> <li>• fosphenytoin</li> <li>• gabapentin cap, tab, soln</li> <li>• lacosamide tab</li> <li>• Lamictal chew</li> <li>• Lamictal XR Kit</li> <li>• lamotrigine tab, chew, ODT, ER tab</li> <li>• lamotrigine starter kit</li> <li>• lamotrigine ODT starter kit</li> <li>• levetiracetam IR, ER, soln, inj</li> </ul>	<ul style="list-style-type: none"> <li>• pregabalin</li> <li>• Nayzilam</li> <li>• Neurontin cap, tab</li> <li>• oxcarbazepine tab, susp</li> <li>• Oxtellar XR</li> <li>• phenobarbital tab, soln, inj</li> <li>• phenytoin cap, chew, susp, inj</li> <li>• primidone</li> <li>• Qudexy XR</li> <li>• Roweepra</li> <li>• Subvenite</li> <li>• Subvenite starter kit</li> <li>• Sympazan</li> <li>• Tegretol IR, XR tab, susp</li> <li>• tiagabine</li> <li>• topiramate tab, IR sprinkle cap</li> <li>• Trileptal Susp</li> <li>• Trokendi XR</li> <li>• valproate inj</li> <li>• valproic acid cap, solution</li> <li>• Valtoco</li> <li>• zonisamide cap</li> </ul>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Aptiom</li> <li>• Banzel tablet</li> <li>• Briviact inj, sol, tab</li> <li>• Diacomit</li> <li>• Elepsia XR</li> <li>• Epidiolex</li> <li>• Fintepla</li> <li>• Fycompa tab, susp</li> <li>• lacosamide inj, sol</li> <li>• Motpoly XR</li> <li>• rufinamide susp</li> <li>• Spritam</li> <li>• vigabatrin</li> <li>• vigadrone</li> <li>• Xcopri tab</li> <li>• Xcopri Titration Pak <ul style="list-style-type: none"> <li>○ QL – 1 Pak/90 days</li> </ul> </li> <li>• Zonisade</li> <li>• Ztalmy</li> </ul> <p>PA criteria AND Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• felbamate</li> <li>• methsuximide</li> <li>• rufinamide tablet</li> </ul>	<p><a href="#">Antiseizure Agents Prior Authorization Criteria</a></p> <p><a href="#">Utilization Edits for Mental Health Medications</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - Continued</b>			
<p><b>Antiseizure Agents - continued</b></p> <p><i>Note: Utilization Edits may apply for mental health medications; see Utilization Edits for Mental Health Medications for associated quantity limits</i></p>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Eprontia</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• carbamazepine ER tab</li> <li>• carbamazepine suspension</li> <li>• topiramate ER capsule</li> <li>• topiramate ER sprinkle capsule</li> </ul> <p>Brand Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Depakote DR, ER</li> <li>• Lamictal IR, ODT, XR</li> <li>• Lamictal IR Starter Kit</li> <li>• Lamictal ODT Starter Kit</li> <li>• Lyrica</li> <li>• Neurontin sol</li> <li>• Onfi tab, susp</li> <li>• Topamax IR, Sprinkle</li> <li>• Trileptal IR tab</li> </ul>		<p><a href="#">Antiseizure Agents Prior Authorization Criteria</a></p> <p><a href="#">Utilization Edits for Mental Health Medications</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - Continued</b>			
<b>Gastroprotective Agents</b>	<ul style="list-style-type: none"> <li>Celebrex</li> <li>naproxen-esomeprazole magnesium</li> </ul>	<ul style="list-style-type: none"> <li>diclofenac-misoprostol delayed release tablets</li> <li>ibuprofen-famotidine</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>celecoxib</li> </ul>	
<b>Movement Disorder Agents</b>	<ul style="list-style-type: none"> <li>benztropine tablet, injection</li> <li>trihexyphenidyl tablet, solution</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>Austedo</li> <li>Austedo XR</li> <li>Austedo Titration Kit</li> <li>Austedo XR Titration Kit</li> <li>Ingrezza</li> <li>Ingrezza Therapy Pack</li> <li>tetrabenazine</li> </ul>	N/A	<a href="#">Movement Disorder Agents PA Criteria</a>
<b>Narcotic Antitussives and Combinations</b>  <i>See Opioid Overutilization with Age and Quantity Limits PA Criteria for product-specific age and quantity limits</i>  <i>*Note: All narcotic antitussives will require PA for members under 18 years of age *</i>	<p>PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>guaifenesin/codeine 100-10mg/5mL solution</li> <li>hydrocodone/ homatropine syrup</li> <li>hydrocodone/homatropine tab</li> <li>Hydromet syrup</li> <li>promethazine VC/codeine syrup</li> <li>promethazine with codeine</li> </ul>	<p>PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>hydrocodone polst/chlorpheniramine polst ER</li> <li>Tuxarin ER</li> </ul>	<a href="#">Opioid Overutilization with Age and Quantity Limits PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - Continued</b>			
<p><b>Narcotics</b></p> <p><i>See Opioid Overutilization with Age and Quantity Limits PA Criteria for product-specific age and quantity limits</i></p> <p><i>Note: All codeine products will require PA for members under 18 years of age</i></p>	<p><b>Short Acting</b> PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• apap/codeine</li> <li>• buprenorphine inj</li> <li>• butorphanol injection <ul style="list-style-type: none"> <li>○ AGE – 18 years of age and older</li> </ul> </li> <li>• butorphanol 10 mg/mL nasal spray</li> <li>• codeine sulfate</li> <li>• codeine/butalbital/apap/caffeine</li> <li>• codeine/butalbital/asa/caffeine</li> <li>• hydrocodone/apap</li> <li>• hydrocodone/ibu</li> <li>• hydromorphone</li> <li>• levorphanol</li> <li>• meperidine</li> <li>• morphine</li> <li>• nalbuphine</li> <li>• Nucynta</li> <li>• opium tincture</li> <li>• oxycodone</li> <li>• oxycodone/apap</li> <li>• pentazocine/naloxone</li> <li>• tramadol</li> <li>• tramadol/APAP</li> </ul> <p><b>Long Acting</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Butrans</li> <li>• fentanyl patches</li> <li>• morphine ER tab (MS Contin)</li> <li>• Nucynta ER</li> </ul>	<p><b>Short Acting</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• fentanyl citrate lozenges</li> <li>• fentanyl citrate buccal tablets</li> <li>• Fentora buccal tablets</li> </ul> <p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Apadaz</li> <li>• apap/caffeine/dihydrocodeine</li> <li>• benzhydrocodone/APAP</li> <li>• belladonna and opium suppositories</li> <li>• Nalocet</li> <li>• oxycodone AD</li> <li>• oxymorphone IR</li> <li>• Prolate</li> <li>• RoxyBond</li> <li>• Seglentis</li> <li>• tramadol 5 mg/mL solution</li> <li>• Trezix</li> </ul> <p><b>Long Acting</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Belbuca</li> <li>• hydrocodone ER cap (Zohydro)</li> <li>• Hysingla ER</li> <li>• hydromorphone ER tab (Exalgo)</li> <li>• methadone</li> <li>• morphine ER cap (Avinza, Kadian)</li> <li>• oxycodone ER tab</li> <li>• Oxycontin</li> <li>• oxymorphone ER tab (Opana)</li> <li>• Tramadol ER (Conzip, Ryzolt, Ultram ER)</li> <li>• Xtampza ER</li> </ul> <p>PA criteria AND Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• buprenorphine patches</li> <li>• hydrocodone ER tab (Hysingla ER)</li> </ul>	<p><a href="#">APAP High Dose PA Criteria</a></p> <p><a href="#">Fentanyl Citrate PA Criteria</a></p> <p><a href="#">Opioid Overutilization with Age and Quantity Limits PA Criteria</a></p> <p><a href="#">Opioid PA Form – Request to Exceed MME Limit</a></p> <p><a href="#">Opioid with Concurrent Buprenorphine/Naloxone PA Form</a></p> <p><a href="#">Benzodiazepine and Opioid Concurrent Therapy PA Form</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - Continued</b>			
<b>Skeletal Muscle Relaxants</b>	<ul style="list-style-type: none"> <li>• baclofen</li> <li>• chlorzoxazone</li> <li>• cyclobenzaprine IR (tabs)</li> <li>• methocarbamol</li> <li>• orphenadrine citrate</li> <li>• tizanidine tablets</li> </ul> <p><b>Granules/Liquid Formulation</b></p> <ul style="list-style-type: none"> <li>• Lyvispah granules               <ul style="list-style-type: none"> <li>○ ST – must be unable to swallow tablets</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• Amrix               <ul style="list-style-type: none"> <li>○ ST – must try cyclobenzaprine tablets within the past 30 days</li> </ul> </li> <li>• dantrolene</li> <li>• Fexmid</li> <li>• Lorzone</li> <li>• metaxalone</li> <li>• Norgesic</li> <li>• Norgesic Forte</li> <li>• Orphengesic Forte</li> <li>• orphenadrine/aspirin/caffeine</li> <li>• tizanidine capsules</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• carisoprodol               <ul style="list-style-type: none"> <li>○ QL – 4 tabs/day</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• cyclobenzaprine ER (caps)               <ul style="list-style-type: none"> <li>○ ST – must try cyclobenzaprine tablets within the past 30 days</li> </ul> </li> </ul> <p><b>Granules/Liquid Formulation</b></p> <ul style="list-style-type: none"> <li>• baclofen 5 mg/5 mL sol; baclofen 10 mg/5 mL sol; baclofen 25 mg/5mL susp; Fleqsuvy susp               <ul style="list-style-type: none"> <li>○ ST – trial and failure of Lyvispah (baclofen) or medical rationale for use</li> </ul> </li> </ul>	<p><a href="#">Carisoprodol Agents PA Criteria</a></p> <p><a href="#">Carisoprodol Agents PA Form</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>CNS AND OTHERS - Continued</b>			
<b>Smoking Deterrent Agents</b>	<p><b><i>Nicotine Replacement</i></b></p> <ul style="list-style-type: none"> <li>• nicotine gum <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 24 pieces/day</li> </ul> </li> <li>• nicotine lozenge <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 20 pieces/day</li> </ul> </li> <li>• nicotine patch <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 1 patch/day</li> </ul> </li> <li>• nicotine patch kit <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 1 kit/90 days</li> </ul> </li> </ul> <p><b><i>Other Smoking Deterrents</i></b></p> <ul style="list-style-type: none"> <li>• bupropion SR 150</li> <li>• varenicline <ul style="list-style-type: none"> <li>○ AGE – 18 years of age or older</li> </ul> </li> </ul>	<p><b><i>Nicotine Replacement</i></b></p> <ul style="list-style-type: none"> <li>• Nicotrol NS <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 12 bottles/30 days</li> </ul> </li> <li>• Nicotrol Inhaler <ul style="list-style-type: none"> <li>○ AGE – 10 years of age or older</li> <li>○ QL – 3 inhalers/31 days</li> </ul> </li> </ul> <p><b><i>Other Smoking Deterrents</i></b> N/A</p>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>DERMATOLOGIC</b>			
<p><b>Acne Agents</b></p> <p><i>Note: All acne agents for members over the age of 25 years require step therapy with one covered OTC acne product</i></p> <p><i>Note: A 14-day trial each of at least 2 preferred agents is required prior to receiving a non-preferred agent.</i></p>	<p><i>All legend generic products are preferred unless otherwise specified</i></p> <ul style="list-style-type: none"> <li>• Adapalene (cream, gel) <ul style="list-style-type: none"> <li>○ AGE – 25 years and under – ST – must have tried a preferred topical tretinoin product</li> <li>○ AGE – &gt;25 years – ST – must have tried one covered OTC acne product AND a preferred topical tretinoin product</li> </ul> </li> <li>• benzoyl peroxide cream, liquid, gel</li> <li>• Finacea foam</li> <li>• Retin-A (all formulations except micro)</li> <li>• Ziana</li> </ul> <p><b>Oral Formulations</b></p> <ul style="list-style-type: none"> <li>• Amnesteem</li> <li>• Claravis</li> <li>• Myorisan</li> <li>• Zenatane</li> </ul>	<p><i>All legend brand products are non-preferred unless otherwise specified</i></p> <ul style="list-style-type: none"> <li>• adapalene/benzoyl peroxide gel</li> <li>• Avar cleanser</li> <li>• Benzepro</li> <li>• BP cleanser</li> <li>• BP 10-1 wash</li> <li>• Cabtreo</li> <li>• clindamycin foam</li> <li>• clindamycin 1.2%/benzoyl peroxide 2.5%</li> <li>• clindamycin 1.2%/benzoyl peroxide 3.75%</li> <li>• dapsone gel</li> <li>• Erygel</li> <li>• PR benzoyl peroxide wash</li> <li>• Retin-A Micro</li> <li>• sodium sulfacetamide med pads</li> <li>• sulfacetamide sod top susp</li> <li>• sodium sulfacetamide-sulfur cleanser, cream, lotion, wash</li> <li>• sulfacetamide topical lotion</li> <li>• tretinoin microsphere</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Avita</li> <li>• clindamycin phosphate-tretinoin gel</li> <li>• tretinoin cream, gel</li> </ul> <p><b>Oral Formulations</b></p> <ul style="list-style-type: none"> <li>• isotretinoin</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>DERMATOLOGIC - Continued</b>			
<b>Antipsoriatics</b>	<ul style="list-style-type: none"> <li>• calcipotriene cream</li> <li>• calcipotriene topical solution</li> <li>• Enstilar</li> <li>• Taclonex scalp suspension</li> <li>• tazarotene 0.1% cream</li> <li>• Vectical ointment</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• acitretin</li> </ul>	<ul style="list-style-type: none"> <li>• calcipotriene 0.005% foam</li> <li>• calcipotriene ointment</li> <li>• calcipotriene/betamethasone ointment</li> <li>• calcitriol ointment</li> <li>• Duobrii</li> <li>• methoxsalen</li> <li>• Sorilux foam</li> <li>• tazarotene 0.05% gel</li> <li>• tazarotene 0.1% gel</li> <li>• Vtama</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• calcipotriene/betamethasone suspension</li> </ul>	<a href="#">Soriatane PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ELECTROLYTE DEPLETERS</b>			
Electrolyte Depleters	<p><b>Miscellaneous Agents</b> N/A</p> <p><b>Phosphate Binders</b></p> <ul style="list-style-type: none"> <li>• calcium acetate capsules</li> <li>• calcium acetate tabs</li> <li>• calcium carbonate 500 mg</li> <li>• calcium carbonate 500 mg, 750 mg, 1000 mg chew</li> <li>• calcium carbonate 1.25 gm (500 mg elemental calcium) chew</li> <li>• calcium carbonate 1.25 gm (500 mg elemental calcium) tab</li> <li>• calcium carbonate 1250 mg/5 mL susp <ul style="list-style-type: none"> <li>○ QL – 30 mL/day</li> </ul> </li> <li>• Magnebind 300 mg tab <ul style="list-style-type: none"> <li>○ QL – 2 bottles (300 tabs)/30 days</li> </ul> </li> <li>• Magnebind Rx</li> <li>• sevelamer HCl 800 mg tabs</li> <li>• sevelamer carbonate powder</li> <li>• sevelamer carbonate tabs</li> </ul> <p><b>Potassium Binders</b></p> <ul style="list-style-type: none"> <li>• Lokelma</li> <li>• SPS (sodium polystyrene sulfonate)</li> <li>• Veltassa</li> </ul>	<p><b>Miscellaneous Agents</b></p> <ul style="list-style-type: none"> <li>• Xphozah <ul style="list-style-type: none"> <li>○ ST – must have tried and failed preferred phosphate binders OR submit medical rationale for use over ALL preferred phosphate binders</li> </ul> </li> </ul> <p><b>Phosphate Binders</b></p> <ul style="list-style-type: none"> <li>• Auryxia</li> <li>• Fosrenol powder packet <ul style="list-style-type: none"> <li>○ ST – member must be under 18 years of age or unable to swallow tablets</li> </ul> </li> <li>• lanthanum carbonate chew</li> <li>• sevelamer HCl 400 mg tabs</li> <li>• Velphoro</li> </ul> <p><b>Potassium Binders</b> N/A</p>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE</b>			
<b>Anaphylaxis Agents</b>	<ul style="list-style-type: none"> <li>epinephrine auto-injector</li> </ul>	<ul style="list-style-type: none"> <li>Auvi-Q</li> <li>Epipen</li> <li>Symjepi</li> </ul>	
<b>Bone Formation Stimulating Agents</b>	PA criteria must be met for the following: <ul style="list-style-type: none"> <li>Forteo</li> </ul>	PA criteria must be met for the following: <ul style="list-style-type: none"> <li>Evenity</li> <li>teriparatide 620 mcg/2.48 mL</li> <li>Tymlos</li> </ul> PA criteria AND Generic Medically Necessary PA criteria must be met for the following: <ul style="list-style-type: none"> <li>teriparatide 600 mcg/2.4 mL</li> </ul>	<a href="#">Bone Formation Stimulating Agents PA Criteria</a>  <a href="#">Bone Formation Stimulating Agents PA Form</a>
<b>Bone Resorption Inhibitors</b>	<p><b><i>Bisphosphonates</i></b></p> <ul style="list-style-type: none"> <li>alendronate</li> <li>risedronate tablets               <ul style="list-style-type: none"> <li>ST – must try alendronate within the past 90 days</li> </ul> </li> </ul> <p><b><i>Bone Modifying Monoclonal Antibodies</i></b> N/A</p> <p><b><i>Calcitonin</i></b></p> <ul style="list-style-type: none"> <li>calcitonin-salmon nasal</li> </ul> <p><b><i>SERMs</i></b></p> <ul style="list-style-type: none"> <li>raloxifene</li> </ul>	<p><b><i>Bisphosphonates</i></b></p> <ul style="list-style-type: none"> <li>alendronate oral solution 70mg/75mL               <ul style="list-style-type: none"> <li>ST – must be 5 years of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> <li>Fosamax Plus D</li> <li>ibandronate</li> <li>ibandronate pre-filled syringe               <ul style="list-style-type: none"> <li>QL – one single-use, pre-filled syringe per 90 days</li> </ul> </li> <li>risedronate DR (generic Atelvia)</li> </ul> <p><b><i>Bone Modifying Monoclonal Antibodies</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>Prolia injection</li> <li>Xgeva</li> </ul> <p><b><i>Calcitonin</i></b></p> <ul style="list-style-type: none"> <li>calcitonin (salmon) injection               <ul style="list-style-type: none"> <li>ST – trial and failure of calcitonin-salmon nasal or medical justification for use over the preferred calcitonin agent</li> </ul> </li> </ul> <p><b><i>SERMs</i></b> N/A</p>	<a href="#">Bone Resorption Inhibitors PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<b>DPP4 Inhibitors and Combination Agents</b>	<p><b><i>DPP4-I</i></b></p> <ul style="list-style-type: none"> <li>• Januvia <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> <li>• Tradjenta <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> </ul> <p><b><i>DPP4-I &amp; metformin combination</i></b></p> <ul style="list-style-type: none"> <li>• Janumet <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> <li>• Janumet XR <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> <li>• Jentadueto <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> <li>• Jentadueto XR <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> </ul> <p><b><i>DPP4-I &amp; thiazolidinedione combination</i></b> N/A</p>	<p><b><i>DPP4-I</i></b></p> <ul style="list-style-type: none"> <li>• alogliptin <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred agent for 60 of the past 100 days</li> </ul> </li> <li>• saxagliptin <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred agent for 60 of the past 100 days</li> </ul> </li> <li>• Zituvio <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred agent for 60 of the past 100 days</li> </ul> </li> </ul> <p><b><i>DPP4-I &amp; metformin combination</i></b></p> <ul style="list-style-type: none"> <li>• alogliptin/metformin <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred combination agent for 60 of the past 100 days</li> </ul> </li> <li>• saxagliptin/metformin ER <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred combination agent for 60 of the past 100 days</li> </ul> </li> <li>• Zituvimet <ul style="list-style-type: none"> <li>○ ST – must have tried a preferred combination agent for 60 of the past 100 days</li> </ul> </li> </ul> <p><b><i>DPP4-I &amp; thiazolidinedione combination</i></b></p> <ul style="list-style-type: none"> <li>• alogliptin/pioglitazone <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes for 60 of the past 100 days</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<b>GLP-1 Receptor Agonists and Combinations</b>	<p><b>GLP-1 RA</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Byetta</li> <li>• Ozempic</li> <li>• Trulicity</li> <li>• Victoza</li> </ul> <p><b>GIP/GLP-1 RA</b> N/A</p> <p><b>Combination Agents</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Soliqua</li> </ul>	<p><b>GLP-1 RA</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Bydureon BCise</li> <li>• Rybelsus</li> </ul> <p><b>GIP/GLP-1 RA</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Mounjaro</li> </ul> <p><b>Combination Agents</b> PSQC criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Xultophy</li> </ul>	<a href="#">GLP-1 RA/GIP RA/Combinations PA Criteria</a>
<b>Glucagon Agents</b>	<ul style="list-style-type: none"> <li>• Baqsimi nasal spray</li> <li>• Gvoke injection</li> <li>• Zegalogue injection</li> </ul>	<ul style="list-style-type: none"> <li>• Glucagon Kit</li> </ul>	
<b>Growth Hormones</b>	<p><b>Somatropin products</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Genotropin</li> <li>• Norditropin</li> <li>• Serostim</li> </ul> <p><b>Long-acting products</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Skytrofa</li> </ul> <p><b>Miscellaneous growth hormone products</b> N/A</p>	<p><b>Somatropin products</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Humatrope</li> <li>• Nutropin AQ</li> <li>• Omnitrope</li> <li>• Saizen</li> <li>• Zomacton</li> </ul> <p><b>Long-acting products</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Ngenla</li> <li>• Sogroya</li> </ul> <p><b>Miscellaneous growth hormone products</b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Increlex</li> <li>• Voxzogo</li> </ul>	<a href="#">Growth Hormone PA Criteria</a>  <a href="#">Growth Hormone for Adults PA Form</a>  <a href="#">Growth Hormone for Children PA Form</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<b>Insulins – Intermediate Acting</b>	<ul style="list-style-type: none"> <li>• insulin aspart (70/30) (Novolog mix ABA)</li> <li>• Humalog Mix 50/50</li> <li>• Humalog Mix 75/25</li> <li>• Humulin N</li> <li>• Humulin 50/50</li> <li>• Humulin 70/30 (all formulations)</li> <li>• Novolin N</li> <li>• Novolin N ReliOn (vials only)</li> <li>• Novolin 70/30</li> <li>• Novolin 70/30 ReliOn (vials only)</li> <li>• Novolog Mix 70/30 (all formulations)</li> <li>• Novolog Mix 70/30 ReliOn (all formulations)</li> </ul>	<ul style="list-style-type: none"> <li>• insulin lispro protamine/insulin lispro Kwikpen</li> <li>• Novolin N ReliOn (prefilled pen, innolets, syringes and cartridges)</li> <li>• Novolin 70/30 ReliOn (prefilled pen, innolets, syringes and cartridges)</li> </ul>	
<b>Insulins – Rapid Acting</b>	<ul style="list-style-type: none"> <li>• Apidra</li> <li>• Apidra SoloStar</li> <li>• Humalog (all formulations)</li> <li>• insulin aspart (all formulations)</li> </ul>	<ul style="list-style-type: none"> <li>• Admelog</li> <li>• Admelog Solostar</li> <li>• Fiasp</li> <li>• Humalog Tempo Pen</li> <li>• insulin lispro (all formulations)</li> <li>• Lyumjev</li> <li>• Lyumjev Tempo Pen</li> <li>• Novolog (all formulations)</li> <li>• Novolog ReliOn (all formulations)</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<b>Insulins – Short Acting</b>	<ul style="list-style-type: none"> <li>• Humulin R (all formulations)</li> <li>• Novolin R (all formulations)</li> <li>• Novolin R ReliOn (vials only)</li> </ul>	<ul style="list-style-type: none"> <li>• Afrezza</li> <li>• Novolin R ReliOn (prefilled pen, innolets, syringes and cartridges)</li> </ul>	
<b>Insulins – Long Acting</b>	<ul style="list-style-type: none"> <li>• insulin degludec Flex &amp; vials</li> <li>• Lantus (cartridges, pens, &amp; vials)</li> </ul>	<ul style="list-style-type: none"> <li>• Basaglar</li> <li>• Basaglar Tempo Pen</li> <li>• insulin glargine (all manufacturers)</li> <li>• Levemir (Flextouch, &amp; vials)</li> <li>• Rezvoglar</li> <li>• Semglee</li> <li>• Toujeo Solostar</li> <li>• Tresiba Flex and vials</li> </ul>	
<b>Miscellaneous Oral Antidiabetic Agents</b>	<p><b><i>Alpha glucosidase inhibitors</i></b></p> <ul style="list-style-type: none"> <li>• acarbose</li> </ul> <p><b><i>Biguanides</i></b></p> <ul style="list-style-type: none"> <li>• Glumetza</li> <li>• metformin</li> <li>• metformin ER (all strengths except 500 mg &amp; 1 gram ER tabs, generics of Fortamet)</li> </ul> <p><b><i>Meglitinide</i></b></p> <ul style="list-style-type: none"> <li>• repaglinide</li> </ul> <p><b><i>Sulfonylureas and Combinations</i></b></p> <ul style="list-style-type: none"> <li>• glimepiride</li> <li>• glipizide</li> <li>• glipizide ER</li> <li>• glipizide/metformin <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> <li>• glyburide</li> <li>• glyburide/metformin <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> </ul> </li> </ul> <p><b><i>Thiazolidinediones and Combinations</i></b></p> <ul style="list-style-type: none"> <li>• pioglitazone <ul style="list-style-type: none"> <li>○ ST – must have tried metformin</li> <li>○ QL – 34 tabs/30 days</li> </ul> </li> </ul>	<p><b><i>Alpha glucosidase inhibitors</i></b></p> <ul style="list-style-type: none"> <li>• miglitol</li> </ul> <p><b><i>Biguanides</i></b></p> <ul style="list-style-type: none"> <li>• metformin 500 mg &amp; 1 gm ER (generics of Fortamet)</li> <li>• metformin HCl solution <ul style="list-style-type: none"> <li>○ ST – must be 10 years of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• metformin ER (generics of Glumetza)</li> </ul> <p><b><i>Meglitinide</i></b></p> <ul style="list-style-type: none"> <li>• nateglinide</li> </ul> <p><b><i>Sulfonylureas and Combinations</i></b> N/A</p> <p><b><i>Thiazolidinediones and Combinations</i></b></p> <ul style="list-style-type: none"> <li>• pioglitazone/glimepiride <ul style="list-style-type: none"> <li>○ ST – prescriber must provide documentation that separate components are unsuitable for use</li> </ul> </li> <li>• pioglitazone/metformin <ul style="list-style-type: none"> <li>○ ST – prescriber must provide documentation that separate components are unsuitable for use</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<b>SGLT Inhibitors and Combinations</b>	<p><b>SGLT1-I/SGLT2-I</b> N/A</p> <p><b>SGLT2-I</b></p> <ul style="list-style-type: none"> <li>• Farxiga</li> <li>• Invokana</li> <li>• Jardiance</li> </ul> <p><b>SGLT2-I &amp; metformin combination</b></p> <ul style="list-style-type: none"> <li>• Invokamet</li> <li>• Synjardy</li> <li>• Xigduo XR</li> </ul> <p><b>SGLT2-I &amp; DPP4-I combination</b> N/A</p> <p><b>SGLT2-I, DPP4-I, &amp; metformin combination</b> N/A</p>	<p><b>SGLT1-I/SGLT2-I</b></p> <ul style="list-style-type: none"> <li>• Inpefa <ul style="list-style-type: none"> <li>○ ST – must try and fail each of the following active ingredients as monotherapy or combination product: canagliflozin, dapagliflozin, empagliflozin OR medical justification for use</li> </ul> </li> </ul> <p><b>SGLT2-I</b></p> <ul style="list-style-type: none"> <li>• Brenzavvy</li> <li>• dapagliflozin (Farxiga ABA)</li> <li>• Steglatro</li> </ul> <p><b>SGLT2-I &amp; metformin combination</b></p> <ul style="list-style-type: none"> <li>• dapagliflozin/metformin (Xigduo ABA)</li> <li>• Invokamet XR</li> <li>• Segluromet</li> <li>• Synjardy XR</li> </ul> <p><b>SGLT2-I &amp; DPP4-I combination</b></p> <ul style="list-style-type: none"> <li>• Glyxambi <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes or provide medical justification for use over preferred agents</li> </ul> </li> <li>• Qtern <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes or provide medical justification for use over preferred agents</li> </ul> </li> <li>• Steglujan <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes or provide medical justification for use over preferred agents</li> </ul> </li> </ul> <p><b>SGLT2-I, DPP4-I, &amp; metformin combination</b></p> <ul style="list-style-type: none"> <li>• Trijardy XR <ul style="list-style-type: none"> <li>○ ST – must have tried and failed combination therapy with preferred agents of the same classes or provide medical justification for use over preferred agents</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ENDOCRINE - Continued</b>			
<p><b>Testosterones</b></p> <p><i>See Testosterone PA Criteria for product-specific age and quantity limits</i></p>	<p><b><i>Injectable Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Depo-Testosterone</li> <li>• testosterone cypionate</li> </ul> <p><b><i>Oral Agents</i></b> N/A</p> <p><b><i>Topical Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Androderm</li> <li>• Testim 1% (50 mg)/5 gm gel tubes</li> <li>• testosterone 1% (25 mg)/2.5 gm gel packets</li> <li>• testosterone 1% (12.5 mg)/act gel pump</li> <li>• testosterone 1.62% (20.25 mg)/act metered pump gel</li> </ul>	<p><b><i>Injectable Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Aveed</li> <li>• Testopel pellet</li> <li>• testosterone enanthate</li> <li>• Xyosted</li> </ul> <p><b><i>Oral Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Danazol</li> <li>• Jatenzo</li> <li>• Methitest</li> <li>• methyltestosterone</li> <li>• Tlando</li> </ul> <p><b><i>Topical Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Natesto</li> <li>• testosterone 1% (50 mg)/5 gm gel packets/tubes</li> <li>• testosterone 1.62% (40.5 mg)/2.5 gm gel packets</li> <li>• testosterone 1.62% (20.25 mg)/1.25 gm gel packets</li> <li>• testosterone 2% (10 mg)/act metered pump</li> <li>• testosterone 30 mg/act solution</li> <li>• Vogelxo 1% (50 mg)/5 gm gel packets</li> <li>• Vogelxo 1% (12.5 mg)/act gel pump</li> </ul>	<p><a href="#">Testosterones PA Criteria</a></p> <p><a href="#">Testosterones PA Form</a></p>
<p><b>Urea Cycle Disorders (Hyperammonemia Treatments)</b></p>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Carbaglu</li> <li>• Pheburane</li> <li>• sodium phenylbutyrate powder</li> <li>• sodium phenylbutyrate tab</li> </ul>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Olpruva</li> <li>• Ravicti</li> </ul> <p>PA criteria AND Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• carglumic acid</li> </ul>	<p><a href="#">Urea Cycle Disorder Agents</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ESTROGEN AND RELATED AGENTS</b>			
<b>Estrogen and Related Agents</b>	<p><i>All legend generic products are preferred unless otherwise specified</i></p> <ul style="list-style-type: none"> <li>• Depo-estradiol</li> <li>• Evamist mist</li> <li>• Menest</li> <li>• Minivelle</li> <li>• Premarin</li> <li>• Prempro</li> <li>• Provera</li> <li>• Vivelle Dot</li> </ul> <p><b><i>Vaginal Preparations</i></b></p> <ul style="list-style-type: none"> <li>• Estring</li> <li>• Premarin Vaginal Cream</li> <li>• Vagifem</li> </ul> <p><b><i>Uterine disorder agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Oriahnn</li> <li>• Orilissa</li> </ul>	<p><i>All legend brand products are non-preferred unless otherwise specified</i></p> <ul style="list-style-type: none"> <li>• estradiol TD gel 0.1%</li> <li>• ethinyl estradiol and norethindrone tabs</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Veozah</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• estradiol TD patch (generic formulations of Minivelle and Vivelle Dot)</li> </ul> <p><b><i>Vaginal Preparations</i></b></p> <ul style="list-style-type: none"> <li>• estradiol vaginal cream</li> <li>• Femring</li> <li>• Yuvaferm</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• estradiol vaginal tablets</li> </ul> <p><b><i>Uterine disorder agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Myfembree</li> </ul>	<p><a href="#">Uterine Disorder Agents PA Criteria</a></p> <p><a href="#">Uterine Disorder Agents PA Form</a></p> <p><a href="#">Veozah PA Criteria</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>ESTROGEN AND RELATED AGENTS - Continued</b>			
<p><b>Contraceptives</b></p> <p><i>Note: All contraceptive agents participating in the Medicaid Drug Rebate Program are preferred unless otherwise specified; Brand Medically Necessary PA criteria will apply to brands with available generics</i></p>	<p><b>Injectable Contraception</b></p> <ul style="list-style-type: none"> <li>• Depo-SubQ Provera</li> <li>• medroxyprogesterone contraceptive 150mg/mL suspension for injection <ul style="list-style-type: none"> <li>○ QL – 1mL/84 days for contraception</li> </ul> </li> </ul> <p><b>Oral/Topical Contraception</b></p> <ul style="list-style-type: none"> <li>• drospirenone</li> <li>• norethindrone</li> <li>• Phexxi <ul style="list-style-type: none"> <li>○ QL – 1 box/month</li> </ul> </li> <li>• progestin/estrogen combinations</li> <li>• Twirla</li> <li>• Xulane</li> </ul> <p><b>Long-Acting Reversible Contraception</b></p> <ul style="list-style-type: none"> <li>• Kyleena <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> <li>• Liletta <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> <li>• Mirena <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> <li>• Nexplanon <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> <li>• Paragard <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> <li>• Skyla <ul style="list-style-type: none"> <li>○ QL – 1 device/365 days</li> </ul> </li> </ul> <p><b>Emergency Contraception</b></p> <ul style="list-style-type: none"> <li>• levonorgestrel 1.5mg</li> <li>• ulipristal</li> </ul>	<p><b>Injectable Contraception</b> N/A</p> <p><b>Oral/Topical Contraception</b></p> <ul style="list-style-type: none"> <li>• Zafemy <ul style="list-style-type: none"> <li>○ ST – must have previous trial of all preferred patch formulations of contraception OR medical justification for use</li> </ul> </li> </ul> <p><b>Long-Acting Reversible Contraception</b> N/A</p> <p><b>Emergency Contraception</b> N/A</p>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GASTROINTESTINAL AGENTS</b>			
<b>Anti-ulcer Agents</b>	<ul style="list-style-type: none"> <li>• misoprostol tablets</li> <li>• sucralfate suspension               <ul style="list-style-type: none"> <li>○ ST – must be 1 year of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> <li>• sucralfate tablets</li> </ul>		
<b>H. Pylori Agents</b>	<ul style="list-style-type: none"> <li>• Pylera</li> </ul>	<ul style="list-style-type: none"> <li>• Helidac</li> <li>• lansoprazole/amoxicillin/clarithromycin caps</li> <li>• Talicia</li> <li>• Voquezna Dual Pak</li> <li>• Voquezna Triple Pak</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• bismuth subcitrate/metronidazole/tetracycline</li> </ul>	
<b>H2 Receptor Antagonists</b>	<ul style="list-style-type: none"> <li>• cimetidine tabs               <ul style="list-style-type: none"> <li>○ QL – 60/30 days</li> </ul> </li> <li>• famotidine tabs               <ul style="list-style-type: none"> <li>○ QL – 60/30 days</li> </ul> </li> <li>• nizatidine caps               <ul style="list-style-type: none"> <li>○ QL – 60/30 days</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• famotidine oral suspension               <ul style="list-style-type: none"> <li>○ ST – member must be under 12 years of age or unable to swallow tablets</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GASTROINTESTINAL AGENTS - Continued</b>			
<b>Laxatives and Cathartics</b>	<ul style="list-style-type: none"> <li>• Linzess; lubiprostone               <ul style="list-style-type: none"> <li>○ ST – requires trial of lactulose, sorbitol, or polyethylene glycol</li> </ul> </li> <li>• Relistor injection               <ul style="list-style-type: none"> <li>○ ST – requires trial of lactulose, sorbitol or polyethylene glycol AND diagnosis of opioid-induced constipation</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• Ibsrela               <ul style="list-style-type: none"> <li>○ ST – requires trial of lubiprostone and Linzess OR trial of lactulose, sorbitol or polyethylene glycol AND medical justification for use over preferred agents</li> </ul> </li> <li>• Motegrity               <ul style="list-style-type: none"> <li>○ ST – requires trial of lubiprostone and Linzess OR trial of lactulose, sorbitol or polyethylene glycol AND medical justification for use over preferred agents</li> </ul> </li> <li>• Movantik               <ul style="list-style-type: none"> <li>○ ST – requires trial of lactulose, sorbitol or polyethylene glycol AND diagnosis of opioid-induced constipation AND medical justification for use over preferred agents</li> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• Relistor tabs               <ul style="list-style-type: none"> <li>○ ST – requires trial of lactulose, sorbitol or polyethylene glycol AND diagnosis of opioid-induced constipation AND medical justification for use over preferred agents</li> <li>○ QL – 3 tabs (450 mg)/day</li> </ul> </li> <li>• Symproic               <ul style="list-style-type: none"> <li>○ ST – requires trial of lactulose, sorbitol or polyethylene glycol AND diagnosis of opioid-induced constipation AND medical justification for use over preferred agents</li> <li>○ QL – 1 tab (0.2mg)/day</li> </ul> </li> <li>• Trulance               <ul style="list-style-type: none"> <li>○ ST – requires trial of lubiprostone and Linzess OR trial of lactulose, sorbitol or polyethylene glycol AND medical justification for use over preferred agents</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GASTROINTESTINAL AGENTS - Continued</b>			
<p><b>Pancreatic Enzymes</b></p> <p><i>Note: Access will be granted to non-preferred agents after cumulatively utilizing 30 days of preferred agent therapy in the past 180 days</i></p>	<ul style="list-style-type: none"> <li>• Creon</li> <li>• Zenpep</li> </ul>	<ul style="list-style-type: none"> <li>• Pertzye</li> <li>• Viokace</li> </ul>	
<p><b>Proton Pump Inhibitors</b></p> <p><i>Note: PA is required for members utilizing therapy for greater than 90 days in a 180-day period.</i></p>	<p><i>See Proton Pump Inhibitors PA Criteria for product-specific quantity limits</i></p> <ul style="list-style-type: none"> <li>• Dexilant</li> <li>• esomeprazole capsules</li> <li>• lansoprazole capsules</li> <li>• omeprazole capsules</li> <li>• pantoprazole tablets</li> </ul> <p><b>IV Solutions</b> N/A</p> <p><b>Oral Solutions</b></p> <ul style="list-style-type: none"> <li>• Nexium packets</li> <li>• Protonix packets</li> </ul>	<p><i>See Proton Pump Inhibitors PA Criteria for product-specific quantity limits, step therapy, and criteria to access non-preferred agents</i></p> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• omeprazole magnesium/sodium bicarbonate caps</li> <li>• rabeprazole</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• dexlansoprazole</li> </ul> <p><b>IV Solutions</b></p> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Nexium IV</li> <li>• pantoprazole IV</li> </ul> <p><b>Oral Solutions</b></p> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Konvomep oral suspension</li> <li>• lansoprazole ODT</li> <li>• omeprazole/sodium bicarb powder</li> <li>• Prilosec packets</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• esomeprazole packets</li> <li>• pantoprazole packets</li> </ul>	<p><a href="#">Proton Pump Inhibitor PA Criteria</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GASTROINTESTINAL AGENTS - Continued</b>			
<b>Ulcerative Colitis Agents</b>	<p><b>Oral Formulations</b></p> <ul style="list-style-type: none"> <li>• Apriso</li> <li>• balsalazide</li> <li>• budesonide DR caps</li> <li>• Dipentum</li> <li>• mesalamine DR (Delzicol) cap</li> <li>• mesalamine DR (Lialda) tab</li> <li>• Pentasa</li> <li>• sulfasalazine IR</li> <li>• sulfasalazine ER</li> </ul> <p><b>Rectal Formulations</b></p> <ul style="list-style-type: none"> <li>• mesalamine enema</li> <li>• mesalamine suppositories</li> <li>• sfRowasa</li> </ul>	<p><b>Oral Formulations</b></p> <ul style="list-style-type: none"> <li>• budesonide ER tabs</li> <li>• Ortikos ER caps</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• mesalamine ER (Apriso) cap</li> <li>• mesalamine ER (Pentasa) cap</li> </ul> <p><b>Rectal Formulations</b></p> <ul style="list-style-type: none"> <li>• Uceris rectal foam</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• budesonide rectal foam</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GENITOURINARY</b>			
<b>BPH Agents</b>	<ul style="list-style-type: none"> <li>• alfuzosin ER</li> <li>• dutasteride</li> <li>• finasteride</li> <li>• tamsulosin</li> </ul>	<ul style="list-style-type: none"> <li>• dutasteride/tamsulosin               <ul style="list-style-type: none"> <li>○ ST – must provide documentation that separate components are not suitable for use</li> </ul> </li> <li>• Entadfi               <ul style="list-style-type: none"> <li>○ ST – prescriber must provide documentation of trial and failure of nonselective alpha-blocker, a selective alpha-blocker, a 5-alpha reductase inhibitor (must include finasteride), and a combination product for the treatment of BPH or a medically justifiable reason that the agents are not suitable for use; therapy duration must not exceed 26 weeks</li> </ul> </li> <li>• silodosin               <ul style="list-style-type: none"> <li>○ ST – requires trial of alfuzosin ER and tamsulosin OR medical justification for use of silodosin over alfuzosin ER and tamsulosin</li> </ul> </li> <li>• tadalafil 2.5mg and 5mg               <ul style="list-style-type: none"> <li>○ ST – prescriber must provide documentation of trial and failure of nonselective alpha-blocker, a selective alpha-blocker, a 5-alpha reductase inhibitor, and a combination product for the treatment of BPH or a medically justifiable reason that the agents are not suitable for use; therapy duration must not exceed 26 weeks if using concurrently with finasteride</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>GENITOURINARY - Continued</b>			
<b>Urinary Tract Antispasmodic/Anti-Incontinence Agents</b>	<ul style="list-style-type: none"> <li>• bethanechol</li> <li>• fesoterodine ER</li> <li>• Gelnique</li> <li>• Myrbetriq tablets</li> <li>• oxybutynin IR</li> <li>• oxybutynin ER</li> <li>• Oxytrol</li> <li>• solifenacin</li> </ul>	<ul style="list-style-type: none"> <li>• darifenacin</li> <li>• flavoxate</li> <li>• Gemtesa <ul style="list-style-type: none"> <li>○ ST – member must have trialed and failed Myrbetriq or have intolerance or contraindication to Myrbetriq</li> </ul> </li> <li>• Myrbetriq granules <ul style="list-style-type: none"> <li>○ ST – must be 3 years of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> <li>• tolterodine</li> <li>• tolterodine SR</li> <li>• trospium</li> <li>• trospium ER</li> <li>• Vesicare LS <ul style="list-style-type: none"> <li>○ ST – must be 2 years of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• mirabegron tablets</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>HEMATOLOGIC</b>			
<b>Direct Oral Anticoagulants</b>	<ul style="list-style-type: none"> <li>• Eliquis <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day of 2.5mg; 4 tabs/day for 7 days, then 2 tabs/day for 5mg</li> </ul> </li> <li>• Eliquis Starter Pack <ul style="list-style-type: none"> <li>○ QL – 1 pack/90 days</li> </ul> </li> <li>• Pradaxa capsules</li> <li>• Xarelto 2.5mg tablets <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day</li> </ul> </li> <li>• Xarelto 10mg tablets <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• Xarelto 15 mg tablets <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day for max 21 consecutive days every 90 days; no duration restriction for once-daily dosing</li> </ul> </li> <li>• Xarelto 20 mg tablets <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> </ul> </li> <li>• Xarelto Starter Kit <ul style="list-style-type: none"> <li>○ QL – 1 starter kit/90 days</li> </ul> </li> <li>• Xarelto suspension <ul style="list-style-type: none"> <li>○ ST – must be under 12-years of age or unable to swallow tablets</li> <li>○ QL – 20 mg/day (20 mL/day)</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• Pradaxa Pak <ul style="list-style-type: none"> <li>○ ST – must be under 8 years of age or unable to swallow capsules OR have medical rationale for use of pellet formulation</li> </ul> </li> <li>• Savaysa <ul style="list-style-type: none"> <li>○ QL – 1 tab/day</li> <li>○ ST – must have trialed Eliquis and Xarelto OR medical justification for use of Savaysa over Eliquis and Xarelto</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• dabigatran capsules</li> </ul>	
<b>Hematinics</b>	<p><b><i>Erythropoiesis-Stimulating Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Aranesp</li> <li>• Epogen</li> <li>• Retacrit</li> </ul> <p><b><i>Miscellaneous Hematinics</i></b></p> <ul style="list-style-type: none"> <li>• N/A</li> </ul>	<p><b><i>Erythropoiesis-Stimulating Agents</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Mircera</li> <li>• Procrit</li> </ul> <p><b><i>Miscellaneous Hematinics</i></b> PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Jesduvroq</li> <li>• Reblozyl</li> </ul>	<p><a href="#">Hematinic Agents PA Criteria</a></p> <p><a href="#">Jesduvroq PA Criteria</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>HEMATOLOGIC - Continued</b>			
<b>Leukocyte Stimulants</b>	<p><b>Short-Acting</b></p> <ul style="list-style-type: none"> <li>• Neupogen</li> <li>• Releuko</li> </ul> <p><b>Long-Acting</b></p> <ul style="list-style-type: none"> <li>• Fylnetra</li> <li>• Nyvepria</li> </ul>	<p><b>Short-Acting</b></p> <ul style="list-style-type: none"> <li>• Granix</li> <li>• Leukine</li> <li>• Nivestym</li> <li>• Zarxio</li> </ul> <p><b>Long-Acting</b></p> <ul style="list-style-type: none"> <li>• Fulphila</li> <li>• Neulasta</li> <li>• Neulasta Onpro</li> <li>• Rolvedon</li> <li>• Stimufend</li> <li>• Udenyca</li> <li>• Udenyca Onbody</li> <li>• Ziextenzo</li> </ul>	
<b>Platelet Aggregation Inhibitors</b>	<ul style="list-style-type: none"> <li>• aspirin/dipyridamole</li> <li>• Brilinta <ul style="list-style-type: none"> <li>○ QL – 2 tabs/day</li> </ul> </li> <li>• cilostazol</li> <li>• clopidogrel 75 mg</li> <li>• clopidogrel 300 mg tablets <ul style="list-style-type: none"> <li>○ QL – 1 tab/Rx</li> </ul> </li> <li>• Prasugrel</li> </ul>	<ul style="list-style-type: none"> <li>• Durlaza</li> <li>• Zontivity</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>LIPOTROPICS</b>			
<b>Bile Acid Sequestrants</b>	<ul style="list-style-type: none"> <li>cholestyramine multi-dose containers</li> <li>colesevelam tablets and suspension</li> <li>Prevalite powder/packets</li> </ul>	<ul style="list-style-type: none"> <li>cholestyramine packets</li> <li>colestipol (granules/tablets)</li> </ul>	
<b>Fibric Acid Derivatives</b>	<ul style="list-style-type: none"> <li>fenofibrate micronized cap (generic Antara)</li> <li>fenofibrate tab (generic Tricor)</li> <li>gemfibrozil</li> </ul>	<ul style="list-style-type: none"> <li>Antara</li> <li>fenofibrate cap</li> <li>fenofibrate micronized cap (generic Lofibra)</li> <li>fenofibric acid cap (generic Trilipix)</li> <li>fenofibric acid tab</li> <li>fenofibrate tab (generic Fenoglide)</li> <li>Lipofen</li> </ul>	
<b>HMG CoA Reductase Inhibitors</b>	<ul style="list-style-type: none"> <li>atorvastatin</li> <li>lovastatin</li> <li>pravastatin</li> <li>rosuvastatin</li> <li>simvastatin</li> </ul>	<ul style="list-style-type: none"> <li>Altoprev</li> <li>Atorvaliq <ul style="list-style-type: none"> <li>ST – must be 10 years of age or older and less than 12 years of age OR unable to swallow tablets</li> </ul> </li> <li>Ezallor</li> <li>fluvastatin</li> <li>fluvastatin ER</li> <li>pitavastatin</li> <li>Zypitamag</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>LIPOTROPICS – Continued</b>			
<b>Lipotropics</b>	<ul style="list-style-type: none"> <li>• ezetimibe</li> <li>• ezetimibe/simvastatin               <ul style="list-style-type: none"> <li>○ ST – must have trial history of a single-agent HMG CoA reductase inhibitor for 90 of the past 120 days</li> </ul> </li> <li>• omega-3-acid ethyl esters</li> <li>• Vascepa               <ul style="list-style-type: none"> <li>○ Age – 18 years of age or older</li> <li>○ QL – 4 capsules/day</li> </ul> </li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Praluent</li> <li>• Repatha</li> </ul>	<ul style="list-style-type: none"> <li>• Nexletol               <ul style="list-style-type: none"> <li>○ ST – must have trialed and failed two statin agents OR a statin in combination with ezetimibe OR medical justification for use over preferred statins and ezetimibe</li> </ul> </li> <li>• Nexlizet               <ul style="list-style-type: none"> <li>○ ST – must have trialed and failed a statin in combination with ezetimibe OR medical justification for use over preferred statins and ezetimibe</li> </ul> </li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Evkeeza</li> <li>• Juxtapid</li> <li>• Leqvio</li> <li>• niacin ER</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• icosapent ethyl               <ul style="list-style-type: none"> <li>○ Age – 18 years of age or older</li> <li>○ QL – 4 capsules/day</li> </ul> </li> </ul>	<p><a href="#">PCSK9 Inhibitors and Select Lipotropics PA Criteria</a></p> <p><a href="#">PCSK9 Inhibitors and Select Lipotropics PA Form</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>MULTIPLE SCLEROSIS AGENTS</b>			
<b>Multiple Sclerosis Agents</b>	PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Avonex</li> <li>• Bafiertam</li> <li>• Betaseron</li> <li>• Copaxone</li> <li>• dalfampridine</li> <li>• dimethyl fumarate</li> <li>• fingolimod 0.5 mg</li> <li>• Gilenya 0.25 mg</li> <li>• Kesimpta</li> <li>• Ocrevus</li> <li>• Plegridy</li> <li>• Rebif</li> <li>• teriflunomide</li> <li>• Tascenso ODT</li> <li>• Zeposia</li> </ul>	PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Briumvi</li> <li>• Extavia</li> <li>• Lemtrada</li> <li>• Mavenclad</li> <li>• Mayzent</li> <li>• Ponvory</li> <li>• Tysabri</li> <li>• Vumerity</li> </ul> PSQC/PA criteria AND Generic Medically Necessary PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• glatiramer</li> <li>• Glatopa</li> </ul>	<a href="#">Multiple Sclerosis PA with Quantity Limits Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>RESPIRATORY</b>			
<b>Antihistamine-Decongestant Combinations/2<sup>nd</sup> Generation Antihistamines</b>	<ul style="list-style-type: none"> <li>• cetirizine 5 mg OTC tabs               <ul style="list-style-type: none"> <li>○ AGE – under 18 years</li> </ul> </li> <li>• cetirizine 10 mg OTC tabs</li> <li>• fexofenadine OTC tabs</li> <li>• levocetirizine Rx tabs</li> <li>• loratadine 10 mg OTC tabs</li> <li>• loratadine 10 mg OTC RDT tabs</li> </ul> <p><b>Combinations</b></p> <ul style="list-style-type: none"> <li>• loratadine/pseudoephedrine 12-hour OTC tabs               <ul style="list-style-type: none"> <li>○ QL – 2 tablets/day</li> <li>○ ST – previous trial and failure of a preferred single-agent 2nd generation antihistamine</li> </ul> </li> <li>• loratadine/pseudoephedrine 24-hour OTC tabs               <ul style="list-style-type: none"> <li>○ QL – 1 tablet/day</li> <li>○ ST – previous trial and failure of a preferred single-agent 2nd generation antihistamine</li> </ul> </li> </ul> <p><b>Liquid Formulation</b></p> <ul style="list-style-type: none"> <li>• cetirizine 1 mg/ml OTC syrup               <ul style="list-style-type: none"> <li>○ AGE – under 18 years</li> <li>○ QL – 10 mL/day</li> </ul> </li> <li>• cetirizine 1 mg/mL Rx syrup               <ul style="list-style-type: none"> <li>○ AGE – under 18 years</li> <li>○ QL – 10 mL/day</li> </ul> </li> <li>• loratadine 1 mg/1ml OTC syrup               <ul style="list-style-type: none"> <li>○ AGE – under 18 years</li> <li>○ QL – 10 mL/day</li> </ul> </li> <li>• levocetirizine Rx oral solution               <ul style="list-style-type: none"> <li>○ QL – 10 mL/day</li> <li>○ ST – must have trial of loratadine solution/syrup or cetirizine solution/syrup</li> </ul> </li> </ul>	<p><b>Note: New patients must first try cetirizine and loratadine within 90 days prior to receiving a non-preferred agent. Patients with an existing PA are not subject to the step edit.</b></p> <ul style="list-style-type: none"> <li>• desloratadine Rx tabs</li> <li>• desloratadine Rx ODT tabs</li> </ul> <p><b>Combinations</b></p> <ul style="list-style-type: none"> <li>• Clarinex-D Rx tabs               <ul style="list-style-type: none"> <li>○ QL – 2 tablets/day</li> <li>○ ST – previous trial and failure of loratadine/pseudoephedrine 12-hour OTC tab</li> </ul> </li> </ul> <p><b>Liquid Formulation</b></p> <ul style="list-style-type: none"> <li>• Clarinex 0.5 mg/ml Rx syrup               <ul style="list-style-type: none"> <li>○ QL – 10 mL/day</li> <li>○ ST – must have trial on both cetirizine and loratadine within the past 90 days</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>RESPIRATORY - Continued</b>			
<b>Antiviral Monoclonal Antibody</b>	<ul style="list-style-type: none"> <li>N/A</li> </ul>	PA criteria must be met for the following: <ul style="list-style-type: none"> <li>Synagis</li> </ul>	<a href="#">Antiviral Monoclonal Antibodies PA</a>  <a href="#">Antiviral Monoclonal Antibodies PA Form</a>
<b>Beta Adrenergics and Corticosteroids</b>  <i>Note: All agents are limited to 1 diskus or inhaler per month unless otherwise specified</i>	<ul style="list-style-type: none"> <li>Advair HFA 45/21 mcg, 115/21 mcg, 230/21 mcg</li> <li>Airduo Respiclick</li> <li>Dulera 200-5 mcg</li> <li>fluticasone/salmeterol (generic Advair Diskus) 100/50 mcg, 250/50 mcg, 500/50 mcg</li> <li>Trelegy Ellipta               <ul style="list-style-type: none"> <li>Asthma ST – must have tried and failed Advair or Symbicort therapy for at least 90 days of the past 120 days</li> <li>COPD ST – must have tried and failed Anoro Ellipta therapy for at least 90 of the past 120 days</li> </ul> </li> </ul> <p>QL of 3 units per 30 days for ages 19 and younger/2 units per 30 days for ages 20 and over apply to the following:</p> <ul style="list-style-type: none"> <li>Dulera 50-5 mcg, 100-5 mcg</li> <li>Symbicort 80-4.5 mcg, 160-4.5 mcg</li> </ul>	<ul style="list-style-type: none"> <li>Airduo Digihaler</li> <li>Airsupra               <ul style="list-style-type: none"> <li>AGE – 18 years of age and older</li> <li>QL – 2 inhalers per 30 days</li> </ul> </li> <li>Breo Ellipta</li> <li>Breztri Aerosphere               <ul style="list-style-type: none"> <li>ST – must have tried and failed Trelegy Ellipta or have contraindication or intolerance to use</li> </ul> </li> <li>fluticasone/salmeterol HFA (ABA Advair HFA) 45-21 mcg, 115-21 mcg, 230-21 mcg</li> <li>fluticasone/salmeterol Respiclick (ABA Airduo Respiclick) 55-13 mcg, 113-14 mcg, 232-14 mcg               <ul style="list-style-type: none"> <li>ST – must have tried at least 90 days of therapy with Airduo Respiclick</li> </ul> </li> <li>fluticasone/vilanterol</li> <li>Wixela</li> </ul> <p>QL of 3 units per 30 days for ages 19 and younger/2 units per 30 days for ages 20 and over apply AND Generic Medically Necessary PA criteria apply to the following:</p> <ul style="list-style-type: none"> <li>budesonide/formoterol 80-4.5 mcg, 160-4.5 mcg</li> <li>Breyna</li> </ul>	
<b>Beta Agonists – Long Acting</b>	<ul style="list-style-type: none"> <li>Serevent</li> </ul>	<ul style="list-style-type: none"> <li>arformoterol</li> <li>formoterol</li> <li>Striverdi Respimat</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>RESPIRATORY - Continued</b>			
<b>Beta Agonists – Short Acting</b>	<ul style="list-style-type: none"> <li>• albuterol – all strengths/formulations excluding tablets</li> </ul> <p>QL of 3 canisters per 30 days for ages 18 and younger/2 canisters per 30 days for ages 19 and over apply to the following:</p> <ul style="list-style-type: none"> <li>• albuterol HFA</li> <li>• Proair HFA</li> <li>• Proair Respiclick</li> <li>• Proventil HFA</li> <li>• Ventolin HFA</li> <li>• Xopenex HFA               <ul style="list-style-type: none"> <li>○ ST – must have tried albuterol HFA in the past 90 days</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• albuterol tablets (brand/generic)</li> <li>• levalbuterol neb solution               <ul style="list-style-type: none"> <li>○ QL – 2 prescriptions per 180 days, 1 box of 24 per prescription</li> </ul> </li> <li>• terbutaline</li> </ul> <p>QL of 3 canisters per 30 days for ages 18 and younger/2 canisters per 30 days for ages 19 and over apply to the following:</p> <ul style="list-style-type: none"> <li>• levalbuterol HFA</li> <li>• Proair Digihaler</li> </ul>	
<b>Bronchodilator Agents-Beta Adrenergic and Anticholinergic Combinations</b>  <i>Note: Must not concurrently use &gt;1 inhaled anticholinergic agent (excluding short-acting nebulization solution)</i>	<p><b>Short-Acting</b></p> <ul style="list-style-type: none"> <li>• Atrovent HFA               <ul style="list-style-type: none"> <li>○ QL – 2 inhalers/30 days</li> </ul> </li> <li>• Combivent Respimat               <ul style="list-style-type: none"> <li>○ QL – 2 inhalers/30 days</li> </ul> </li> <li>• ipratropium solution               <ul style="list-style-type: none"> <li>○ QL – 2 boxes/30 days</li> </ul> </li> <li>• ipratropium/albuterol solution               <ul style="list-style-type: none"> <li>○ QL – 3 boxes/30 days</li> </ul> </li> </ul> <p><b>Long-Acting</b></p> <ul style="list-style-type: none"> <li>• Spiriva Handihaler               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Anoro Ellipta               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Incruse Ellipta               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Spiriva Respimat 1.25 mcg               <ul style="list-style-type: none"> <li>○ ST – must have diagnosis of asthma</li> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Spiriva Respimat 2.5 mcg               <ul style="list-style-type: none"> <li>○ ST – must have trial and failure of Spiriva Handihaler for a least 14 days</li> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> </ul>	<p><b>Short-Acting</b> N/A</p> <p><b>Long-Acting</b></p> <ul style="list-style-type: none"> <li>• Bevespi Aerosphere               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Duaklir Pressair               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Lonhala Magnair               <ul style="list-style-type: none"> <li>○ QL – 1 kit (60 vials)/30 days</li> </ul> </li> <li>• Stiolto Respimat               <ul style="list-style-type: none"> <li>○ QL – 1 box (60 inhalations)/30 days</li> </ul> </li> <li>• Tudorza Pressair               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> <li>• Yupelri               <ul style="list-style-type: none"> <li>○ QL – 1 box (90mL)/30 days</li> </ul> </li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• tiotropium inhalation capsules               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30 days</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>RESPIRATORY - Continued</b>			
<b>Leukotriene Receptor Antagonists</b>	<ul style="list-style-type: none"> <li>• montelukast</li> </ul>	<ul style="list-style-type: none"> <li>• montelukast granules               <ul style="list-style-type: none"> <li>○ ST – must have prescriber documentation indicating tablet formulations are unsuitable for use</li> </ul> </li> <li>• zafirlukast</li> <li>• zileuton SR 12 HR</li> <li>• Zyflo</li> </ul>	
<b>Nasal Antihistamines/Nasal Anti-Inflammatory Steroids</b>	<p><b><i>Antihistamines/Anticholinergics</i></b></p> <ul style="list-style-type: none"> <li>• azelastine 0.1% nasal spray</li> <li>• ipratropium NS</li> </ul> <p><b><i>Steroids/Steroid Combinations</i></b></p> <ul style="list-style-type: none"> <li>• Dymista</li> <li>• Fluticasone</li> <li>• Omnaris</li> </ul>	<p><b><i>Antihistamines/Anticholinergics</i></b></p> <ul style="list-style-type: none"> <li>• azelastine 0.15% nasal spray</li> <li>• olopatadine</li> <li>• Patanase</li> </ul> <p><b><i>Steroids/Steroid Combinations</i></b></p> <ul style="list-style-type: none"> <li>• Beconase AQ</li> <li>• budesonide nasal suspension</li> <li>• flunisolide</li> <li>• mometasone nasal susp</li> <li>• Qnasl</li> <li>• Ryaltris</li> <li>• Zetonna</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• azelastine/fluticasone nasal spray</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>RESPIRATORY - Continued</b>			
<b>Oral Inhaled Glucocorticoids</b>	<ul style="list-style-type: none"> <li>• Arnuity Ellipta               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30days</li> </ul> </li> <li>• Asmanex, Asmanex HFA               <ul style="list-style-type: none"> <li>○ QL – 1 inhaler/30days</li> </ul> </li> <li>• budesonide inhalation suspension               <ul style="list-style-type: none"> <li>○ AGE – 3 years and younger</li> <li>○ QL – 120 mL/30 days (0.25 mg/2 mL vial, 0.5 mg/2 mL vial); 60 mL/30 days (1 mg/2 mL vial)</li> </ul> </li> <li>• fluticasone propionate HFA</li> <li>• fluticasone Diskus</li> <li>• Pulmicort Flexhaler</li> <li>• QVAR Redihaler</li> </ul>	<ul style="list-style-type: none"> <li>• Alvesco</li> <li>• Armonair Digihaler</li> <li>• budesonide inhalation suspension               <ul style="list-style-type: none"> <li>○ AGE – 4 years and older</li> <li>○ QL – 120 mL/30 days (0.25 mg/2 mL vial, 0.5 mg/2 mL vial); 60 mL/30 days (1 mg/2 mL vial)</li> </ul> </li> <li>• Flovent Diskus</li> <li>• Flovent HFA</li> </ul>	
<b>Pulmonary Antihypertensives</b>	<p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• sildenafil inj/susp/tab</li> <li>• tadalafil</li> <li>• Tracleer</li> <li>• Tracleer dispersible tablet</li> </ul>	<p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Adempas</li> <li>• ambrisentan</li> <li>• bosentan</li> <li>• Liqrev</li> <li>• Opsumit</li> <li>• Opsynvi</li> <li>• Orenitram. Orenitram Titration Pack</li> <li>• Tadliq</li> <li>• Tyvaso, Tyvaso DPI</li> <li>• Upravi</li> <li>• Ventavis</li> <li>• Winrevair</li> </ul>	<p><a href="#">Pulmonary Antihypertensives PA Criteria</a></p> <p><a href="#">Pulmonary Antihypertensives PA Form</a></p>
<b>Respiratory and Allergy Biologics</b>	<p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Dupixent</li> <li>• Fasenra</li> <li>• Nucala</li> <li>• Tezspire</li> <li>• Xolair</li> </ul>	<p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Cinqair</li> </ul>	<p><a href="#">Respiratory and Allergy Biologics PA Criteria</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>TARGETED IMMUNOMODULATORS</b>			
<b>Targeted Immunomodulators</b>	PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Actemra</li> <li>• Adalimumab               <ul style="list-style-type: none"> <li>○ adalimumab-fkjp</li> <li>○ Hadlima</li> <li>○ Humira</li> </ul> </li> <li>• Adbry</li> <li>• Enbrel</li> <li>• Infliximab               <ul style="list-style-type: none"> <li>○ Infliximab (unbranded Remicade)</li> </ul> </li> <li>• Kineret</li> <li>• Olumiant</li> <li>• Orencia vials &amp; syringes</li> <li>• Otezla</li> <li>• Simponi, Simponi Aria</li> <li>• Taltz</li> <li>• Xeljanz               <ul style="list-style-type: none"> <li>○ Xeljanz oral solution                   <ul style="list-style-type: none"> <li>○ AND the member is 2 years of age or older and weighing 10 kg or more AND less than 18 years of age and weighing less than 40 kg, OR provider has submitted documentation supporting inability to swallow tablet formulation</li> </ul> </li> </ul> </li> </ul>	PSQC/PA criteria must be met for the following: <ul style="list-style-type: none"> <li>• Adalimumab               <ul style="list-style-type: none"> <li>○ Abrilada</li> <li>○ adalimumab-aacf</li> <li>○ adalimumab-adaz</li> <li>○ adalimumab-adbm</li> <li>○ Amjevita</li> <li>○ Cyltezo</li> <li>○ Hulio</li> <li>○ Hyrimoz</li> <li>○ Idacio</li> <li>○ Yuflyma</li> <li>○ Yusimry</li> </ul> </li> <li>• Arcalyst</li> <li>• Bimzelx</li> <li>• Cibirqo</li> <li>• Cimzia</li> <li>• Cosentyx</li> <li>• Entyvio, Entyvio Pen</li> <li>• Ilaris</li> <li>• Ilumya</li> <li>• Infliximab               <ul style="list-style-type: none"> <li>○ Avsola</li> <li>○ Inflectra</li> <li>○ Remicade</li> <li>○ Renflexis</li> </ul> </li> <li>• Kevzara</li> <li>• Litfulo</li> <li>• Omvoh</li> <li>• Rinvoq</li> <li>• Siliq</li> <li>• Skyrizi</li> <li>• Sotyktu</li> <li>• Spevigo</li> <li>• Stelara</li> <li>• Tremfya</li> <li>• Velsipity</li> <li>• Xeljanz XR</li> </ul>	<a href="#">Targeted Immunomodulators PA Criteria</a>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.



DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>TOPICAL AGENTS</b>			
<p><b>Dry Eye Disease or Keratoconjunctivitis</b></p> <p><i>See Dry Eye Disease or Keratoconjunctivitis PA Criteria for product-specific quantity limits</i></p> <p><i>*Note: No more than a 30-day supply may be dispensed at one time.</i></p>	<p>PSQC/PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Restasis single dose</li> <li>• Xiidra</li> </ul>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Cequa</li> <li>• cyclosporine single dose emulsion</li> <li>• Eysuvis</li> <li>• Miebo</li> <li>• Restasis Multidose</li> <li>• Tyrvaya</li> <li>• Verkazia</li> <li>• Vevye</li> </ul>	<p><a href="#">Dry Eye Disease or Keratoconjunctivitis PA criteria</a></p>
<p><b>Miotics-Intraocular Pressure Reducers</b></p>	<ul style="list-style-type: none"> <li>• Alphagan-P 0.1%</li> <li>• Alphagan-P 0.15%</li> <li>• apraclonidine</li> <li>• Azopt</li> <li>• Betoptic-S</li> <li>• brimonidine 0.2% solution</li> <li>• carteolol</li> <li>• Combigan</li> <li>• dorzolamide</li> <li>• dorzolamide/timolol</li> <li>• lopicol 1%</li> <li>• latanoprost</li> <li>• levobunolol</li> <li>• Lumigan 0.01% drops</li> <li>• metipranolol</li> <li>• pilocarpine</li> <li>• Rhopressa</li> <li>• Rocklatan</li> <li>• timolol solution</li> <li>• Travatan Z</li> </ul>	<ul style="list-style-type: none"> <li>• Betaxolol</li> <li>• Betimol</li> <li>• bimatoprost 0.03%</li> <li>• Cosopt PF</li> <li>• Iyuzeh <ul style="list-style-type: none"> <li>○ ST – must have tried and failed latanoprost OR prescriber has provided medical justification for use of Iyuzeh over latanoprost</li> </ul> </li> <li>• Phospholine Iodide</li> <li>• Simbrinza <ul style="list-style-type: none"> <li>○ ST – must provide documentation that separate components are not suitable for use (Azopt/brimonidine)</li> </ul> </li> <li>• timolol gel</li> <li>• Timoptic-XE</li> <li>• Vyzulta</li> <li>• Xelpros</li> <li>• Zioptan</li> </ul> <p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Vuity</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• brimonidine 0.1% solution</li> <li>• brimonidine 0.15% solution</li> <li>• brimonidine/timolol soln</li> <li>• brinzolamide suspension</li> <li>• tafluprost</li> <li>• travaprost 0.004%</li> </ul>	<p><a href="#">Presbyopia Agents PA criteria</a></p>

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>TOPICAL AGENTS - Continued</b>			
<b>Ophthalmic Antihistamines</b>	<ul style="list-style-type: none"> <li>• Alaway</li> <li>• azelastine</li> <li>• Bepreve</li> <li>• Ketotifen</li> <li>• olopatadine (Rx)</li> </ul>	<ul style="list-style-type: none"> <li>• epinastine</li> <li>• Zerviate</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• bepotastine besilate</li> </ul>	
<b>Ophthalmic Anti-Inflammatory Agents</b>	<p><b>NSAIDs</b></p> <ul style="list-style-type: none"> <li>• diclofenac 0.1% ophth soln</li> <li>• flurbiprofen 0.3% ophth soln</li> <li>• ketorolac 0.4% ophth soln</li> <li>• ketorolac 0.5% ophth soln</li> </ul> <p><b>Steroids</b></p> <ul style="list-style-type: none"> <li>• Alrex 0.2% ophth susp</li> <li>• dexamethasone sod phos 0.1% ophth soln</li> <li>• Durezol 0.05% ophth emul</li> <li>• FML Liquifilm 0.1% ophth susp</li> <li>• Lotemax 0.5% ophth gel/ointment/susp</li> <li>• Pred Forte 1% ophth susp</li> <li>• Pred Mild 0.12% ophth susp</li> <li>• Prednisolone sod phos 1% ophth soln</li> </ul>	<p><b>NSAIDs</b></p> <ul style="list-style-type: none"> <li>• Acuvail 0.45% ophth soln</li> <li>• bromfenac 0.09% ophth soln</li> <li>• Bromsite 0.075% ophth soln</li> <li>• Ilevro 0.3% ophth soln</li> <li>• Nevanac 0.1% ophth soln</li> <li>• Prolensa 0.07% ophth soln</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• bromfenac 0.07% ophth soln</li> <li>• bromfenac 0.075% ophth soln</li> </ul> <p><b>Steroids</b></p> <ul style="list-style-type: none"> <li>• Flarex 0.1% ophth susp</li> <li>• FML Forte 0.25% ophth susp</li> <li>• Inveltys 1% ophth susp</li> <li>• Lotemax SM 0.38% ophth gel</li> <li>• Maxidex 0.1% ophth susp</li> </ul> <p>Generic Medically Necessary PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• difluprednate 0.05% ophth emul</li> <li>• fluorometholone 0.1% ophth susp</li> <li>• loteprednol 0.2% ophth susp</li> <li>• loteprednol ophth gel/susp 0.5%</li> <li>• prednisolone 1% ophth susp</li> </ul>	
<b>Ophthalmic Mast Cell Stabilizers</b>	<ul style="list-style-type: none"> <li>• cromolyn</li> </ul>	<ul style="list-style-type: none"> <li>• Alocril</li> <li>• Alomide</li> </ul>	
<b>Otic Preparations</b>	<ul style="list-style-type: none"> <li>• acetic acid solution</li> <li>• Dermotic Oil</li> </ul>	<ul style="list-style-type: none"> <li>• acetic acid HC</li> <li>• fluocinolone acetonide oil</li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

DRUG CLASS	PREFERRED	NON-PREFERRED	PA CRITERIA (if applicable)
<b>TOPICAL AGENTS - Continued</b>			
<b>Topical Anti-Inflammatory Agents – NSAIDS</b>	<ul style="list-style-type: none"> <li>• diclofenac 1% gel</li> <li>• Pennsaid topical solution</li> </ul>	<p>All of the following require physician documentation indicating oral medications are unsuitable for use AND trial and failure of diclofenac 1% gel and Pennsaid topical solution, OR medical justification for use over preferred agents</p> <ul style="list-style-type: none"> <li>• diclofenac solution</li> <li>• diclofenac epolamine patch               <ul style="list-style-type: none"> <li>○ QL – 2 patches per day</li> </ul> </li> <li>• Flector patch               <ul style="list-style-type: none"> <li>○ QL – 2 patches per day</li> </ul> </li> <li>• Licart ER patch               <ul style="list-style-type: none"> <li>○ QL – 1 patch per day</li> </ul> </li> </ul>	
<b>Topical Antiparasitics</b>  <i>Unless otherwise specified, all products are limited to one bottle or one tube per claim</i>	<ul style="list-style-type: none"> <li>• Natroba</li> <li>• permethrin 5% cream</li> <li>• permethrin 1% lotion</li> </ul>	<ul style="list-style-type: none"> <li>• Crotan</li> <li>• ivermectin lotion</li> <li>• Lindane shampoo</li> <li>• Malathion</li> <li>• spinosad</li> <li>• VanaLice</li> </ul>	
<b>Topical Immunomodulators</b>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• pimecrolimus cream</li> <li>• tacrolimus ointment</li> </ul>	<p>PA criteria must be met for the following:</p> <ul style="list-style-type: none"> <li>• Eucrisa</li> <li>• Opzelura</li> <li>• Zoryve 0.3% cream</li> <li>• Zoryve 0.3% foam</li> </ul>	<a href="#">Topical Immunomodulators PA criteria</a>
<b>Topical Post-Herpetic Neuralgia Agents</b>	<ul style="list-style-type: none"> <li>• lidocaine patches               <ul style="list-style-type: none"> <li>○ QL – 3 boxes/30 days</li> </ul> </li> <li>• Lidoderm               <ul style="list-style-type: none"> <li>○ QL – 3 boxes/30 days</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• Synera</li> <li>• ZTlido               <ul style="list-style-type: none"> <li>○ QL – 3 boxes/30 days</li> </ul> </li> <li>• Qutenza               <ul style="list-style-type: none"> <li>○ ST – must have tried lidocaine patches and over-the-counter capsaicin cream</li> <li>○ QL – 4 patches/3 months</li> </ul> </li> </ul>	

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

MISCELLANEOUS INFORMATION

[Preferred Brand Drug List](#)  
[OTC Drug Formulary](#)  
[Pharmacy Supplements Formulary](#)  
[OTC Contraceptive Agents Formulary](#)  
[Brand Medically Necessary/Generic Medically Necessary Prior Authorization Form](#)  
[IHCP Early Refill Prior Authorization Request Form](#)  
[Non-Drug-Specific PA Criteria](#)  
[PBM Call Center LTC ProDUR and Home Health PA Request Form](#)  
[PBM Call Center Prior Authorization Form](#)  
[Vaccine Utilization Edits](#)  
[Vaccine Utilization Edits for VFC-Enrolled Pharmacies](#)

[Mental Health Medications Medical Necessity Prior Authorization Form](#)  
[Antipsychotic Therapy PA with QL](#)  
[Sedative Hypnotics Benzodiazepine PA Criteria](#)  
[Benzodiazepine and Opioid Concurrent Therapy PA Form](#)  
[SSRI/SNRI/NRI Duplicate Therapy PA Criteria with QL](#)  
[Stimulants PA Criteria](#)  
[Hetlioz PA Criteria](#)  
[Hetlioz PA Form](#)  
[Igalmi PA Criteria](#)  
[Narcolepsy Agents PA Criteria](#)  
[Narcolepsy Agents PA Form](#)  
[Nuplazid PA Criteria](#)  
[Utilization Edits for Mental Health Medications](#)

Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.

## MISCELLANEOUS INFORMATION - Continued

[Allergy Specific Immunotherapy PA Criteria](#)

[Amyloid Beta-Directed Antibodies](#)

[Aromatase Inhibitors PA Criteria](#)

[Complement Inhibitor Agents PA](#)

[Corticotropin](#)

[Cushing Syndrome Agents](#)

[Cushing Syndrome Agents PA Form](#)

[Cystic Fibrosis Inhaled Agents PA Criteria](#)

[Cystic Fibrosis Agents PA Criteria](#)

[Cystic Fibrosis Agents PA Form](#)

[Daliresp PA Criteria](#)

[Daliresp PA Form](#)

[Daybue PA Criteria](#)

[Disposable Insulin Delivery Devices PA](#)

[Egrifta PA Criteria](#)

[Elevidys PA Criteria](#)

[Elmiron PA Criteria](#)

[Epidermolysis Bullosa Agents PA](#)

[Gralise, Horizant, and Lyrica CR PA Criteria](#)

[Gralise, Horizant, and Lyrica CR PA Form](#)

[HCG PA Criteria](#)

[Hemophilia B Gene Therapy PA](#)

[Hepatitis B Agents PA Criteria](#)

[High Dollar Compounded PA Criteria](#)

[High Dollar Compounded PA Request Form](#)

[Immunoglobulin A Nephropathy \(IgAN\) Agents PA](#)

[Lucemyra PA Criteria](#)

[Lucemyra PA Form](#)

[MASH/MASLD Agents](#)

[Mepron PA Criteria](#)

[Muscular Dystrophy Agents PA Criteria](#)

[Muscular Dystrophy Agents PA Form](#)

[Non-SUPDL Agents PA and ST](#)

[Nuedexta PA Criteria](#)

[Nuedexta PA Form](#)

[Oxervate PA Criteria](#)

[Pompe Disease Agents PA Criteria](#)

[Prenatal Vitamins High Dollar Limit PA](#)

[Roctavian PA Criteria](#)

[Sickle Cell Agents PA Criteria](#)

[Sickle Cell Agents PA Form](#)

[Skyclarys PA criteria](#)

[Solaraze PA Criteria](#)

[Somatostatin Analog PA Criteria](#)

[Spinal Muscular Atrophy Agents PA Criteria](#)

[Spinal Muscular Atrophy Agents PA Form](#)

[Topical Doxepin PA](#)

[Topical Lidocaine QL](#)

[Topical Steroid PA](#)

[Topical Agents PA Form](#)

[Tzield PA](#)

[Tzield PA Form](#)

[Vyndaqel and Vyndamax PA Criteria](#)

[Wegovy PA](#)

[Zurzuva PA criteria](#)

**Effective for FFS claims submitted on or after September 1, 2024. Effective for Managed Care claims submitted on or after September 15, 2024. V1.2**

Inclusion or reference to any given drug does not indicate market availability of the drug. Drugs that will be, or have been, withdrawn from the market will be removed from the SUPDL as part of routine periodic updating of the SUPDL.