

**DIVISION OF MEDICAID AND LONG-TERM CARE**  
Nebraska Department of Health and Human Services

**PHARMACEUTICAL AND THERAPEUTICS (P&T)  
COMMITTEE MEETING MINUTES**

Wednesday, May 14<sup>th</sup> at 9:00 AM CST  
Mahoney State Park, Peter Kiewit Lodge  
28500 West Park Hwy, Ashland, NE 68003

**Committee Members Present:**

Andrew Bendlin, Pharm.D.  
Cassie Cowles, APRN  
Allison Dering-Anderson, Pharm.D. **(Chair)**  
Stephen Dolter, M.D.  
Jennifer Hill, M.D.  
Laura Klug, Pharm.D.  
Jessica Pohl, Pharm.D.  
Steven Rose, D.O.  
Bradley Sundsboe, Pharm.D.

**Division of Medicaid and Long-Term Care Staff Present:**

Dianne Garside, Pharm.D.  
Spencer Moore, Pharm.D.  
Leah Spencer, R.N., M.Ed.  
Lee Stutzman, Pharm.D.

**Prime Therapeutics Staff Present:**

Nikia Bennette-Carter, Pharm.D., Clinical Account Executive  
ShaLeigh Hammons, CPhT, Account Operations Executive  
Sandy Pranger, Pharm.D., Sr. Director, Clinical Account Services

**Managed Care Staff Present:**

Jamie Benson, Pharm.D., Nebraska Total Care  
Shannon Nelson, Pharm. D., Molina Healthcare  
Bernadette Ueda, Pharm. D., United Healthcare of Nebraska

**Committee Members Excused:**

Eric Avery, M.D.  
Claire Baker, M.D.  
Wade Fornander, M.D.  
C. Jose Friesen, M.D.  
Stephen Salzbrenner, M.D.  
Joyce Juracek, Pharm.D.  
Sarah Stewart- Bouckaert, Pharm.D.

**Committee Members Unexcused:**

N/A

**1. Opening of Public Meeting and Call to Order Committee Business**

- a. The meeting was called to order by the committee chair at 9:00 AM CST. The agenda was posted on the Nebraska Medicaid Pharmacy website (<https://nebraska.fhsc.com/PDL/PTcommittee.asp>) on Monday, April 14<sup>th</sup>. A copy of the Open Meetings Act and meeting materials distributed to members were made available at the physical meeting site for public viewing.
- b. Introduction of new committee member. Dianne Garside welcomed Stephen Rose, D.O., as the newest committee member.
- c. Roll Call: See list above.
- d. Conflict of Interest: No new conflicts of interest were reported.

- e. Approval of November 13<sup>th</sup>, 2024, P&T Committee Meeting Minutes.

Approval of November 13 <sup>th</sup> , 2024 P&T Committee Meeting Minutes							
<b>(1<sup>st</sup>) Motion:</b> Friesen							
<b>(2<sup>nd</sup>) Motion:</b> Hill							
<b>Discussion:</b> Approve as written.							
<b>Voting – P&amp;T Committee Members</b> <small>Does not include excused or unexcused members</small>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.			X
Dering-Anderson, Allison, Pharm.D. (Chair) <i><b>Votes only in the event of a tie</b></i>				Rose, Steven, D.O.			X
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.			X
Hill, Jennifer, M.D.	X						

- f. Department information: Dianne Garside provided an update on the department's new Nebraska Director of Medicaid & Long-Term Care, Drew Gonshorowski.

## 2. Public Testimony

Speaker Order	DRUG CLASS	Drug Name	PDL Status	Speaker Name	Affiliation
1	Antimigraine Agents, Other	Ajovy	P	Dave Miley	Teva
2	Cystic Fibrosis, Oral	Alyftrek	NP	Chad Duncan	Vertex Pharmaceuticals
3	Glucagon Agents	Gvoke	NP	Sue McLaughlin	Children's Nebraska
4	HAE Treatments	Orladeyo	NP	Giuseppe Miranda	BioCryst
5	Lipotropics, Other	Repatha	P	Nicole Nesselhauf	Amgen

## 3. Committee Closed Session

<b>(1<sup>st</sup>) Motion:</b> Dering-Anderson	<b>(2<sup>nd</sup>) Motion:</b> Dolter
<b>Committee Closed Session unanimously approved by all in attendance</b>	

## 4. Resume Open Session

<b>A motion was made to Resume Open Session and was unanimously approved by all in attendance.</b>
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*During the public open session, committee members vote publicly on decisions with regards to the Nebraska Preferred Drug List recommendations. Per the State of Nebraska P&T Committee By-Laws, the minutes reflect how each member voted or if the member was absent or not voting. The chairperson votes only in the event of a tie. The details of each vote and the associated PDL recommendations are presented in the following tables.*

a. Consent Agenda

Consent Agenda							
<b>(1<sup>st</sup>) Motion:</b> Dolter							
<b>(2<sup>nd</sup>) Motion:</b> Hill							
<b>Discussion:</b> Committee removed one Consent Agenda class and added it to Therapeutic Class Reviews: Glucagon Agents. The Committee approved the amended Consent Agenda.							
<b>Voting – P&amp;T Committee Members</b> <small>Does not include excused or unexcused members</small>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <i><b>Votes only in the event of a tie</b></i>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

Consent Agenda: Therapeutic categories (TC) with unchanged recommendations unless otherwise indicated.	
ANALGESICS, OPIOIDS LONG-ACTING	HEPATITIS C COURSES
ANDROGENIC AGENTS	HIV/AIDS
ANTICOAGULANTS	HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS
ANTIMIGRAINE, OTHER	HYPOGLYCEMICS, MEGLITINIDES
ANTIPARASITICS, TOPICAL	HYPOGLYCEMICS, TZDS
ANTIVIRALS, TOPICAL	LINCOSAMIDES/ OXAZOLIDINONES/ STREPTOGRAMINS
BONE RESORPTION SUPPRESSION AND RELATED AGENTS	MACROLIDES AND KETOLIDES
BPH- BENIGN PROSTATIC HYPERPLASIA AGENTS	NITROFUAN DERIVATIVES
CEPHALOSPORINS AND RELATED ANTIBIOTICS	PANCREATIC ENZYMES
<b>GLUCAGON AGENTS-(Removed)</b>	PENICILLINS
GROWTH HORMONE	PLATELET AGGREGATION INHIBITORS
H. PYLORI TREATMENT	THYROID HORMONES
HEPATITIS B AGENTS	UTERINE DISORDER TREATMENTS
HEPATITIS C AGENTS	VASODILATORS, CORONARY

**b. Therapeutic Class Reviews**

Review Agenda – ACNE AGENTS, TOPICAL							
<b>(1<sup>st</sup>) Motion:</b> Pohl							
<b>(2<sup>nd</sup>) Motion:</b> Cowles							
<b>Discussion:</b> Approved as written.							
<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

Review Agenda – ANALGESICS, OPIOIDS SHORT-ACTING							
<b>(1<sup>st</sup>) Motion:</b> Pohl							
<b>(2<sup>nd</sup>) Motion:</b> Hill							
<b>Discussion:</b> Approved as written.							
<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANGIOTENSIN MODULATOR COMBINATIONS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANGIOTENSIN MODULATORS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIBIOTICS, GASTROINTESTINAL

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIBIOTICS, INHALED

**(1<sup>st</sup>) Motion:** Rose

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Rose made a motion to keep TOBI-PODHALER in the preferred position on the PDL.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIBIOTICS, TOPICAL

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIBIOTICS, VAGINAL

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIEMETICS / ANTIVERTIGO AGENTS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIFUNGALS, ORAL

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						



## Review Agenda – ANTIFUNGALS, TOPICAL

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIMIGRAINE AGENTS, TRIPTANS

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – ANTIVIRALS, ORAL

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <i>Votes only in the event of a tie</i>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – BETA-BLOCKERS

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Avery, Eric, M.D.	X			Fornander, Wade, M.D. ( <b>Vice Chair</b> )	X		
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN				Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <i>Votes only in the event of a tie</i>	X			Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – BLADDER RELAXANT PREPARATIONS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – CALCIUM CHANNEL BLOCKERS

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – CONTRACEPTIVES, ORAL

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – CYSTIC FIBROSIS, ORAL

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – DIURETICS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – FLUOROQUINOLONES

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – GI MOTILITY

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – GLUCAGON AGENTS

**(1<sup>st</sup>) Motion:** Dolter

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Dolter made a motion to move Gvoke Pen and Syringe and the Glucagon Emergency Kit (Fresenius) from NP to P due to access for pediatric patients and Lilly discontinuing their preferred Glucagon Emergency Kit.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – HAE TREATMENTS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – HYPOGLYCEMICS, INCRETIN MIMETICS/ ENHANCERS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – HYPOGLYCEMICS, INSULIN AND RELATED AGENTS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – HYPOGLYCEMICS, METFORMINS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						



## Review Agenda – HYPOGLYCEMICS, SGLT2

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Bendlin

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <i><b>Votes only in the event of a tie</b></i>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – HYPOGLYCEMICS, SULFONYLUREAS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <i><b>Votes only in the event of a tie</b></i>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – IMMUNOSUPPRESSIVES, ORAL

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – LIPOTROPICS, OTHER

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** The committee asked the state to review the Repatha criteria that currently state failure to reach target LDL-C levels and add target levels for very high risk ASCVD < 55 mg/dL. Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – LIPOTROPICS, STATINS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Rose

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – MULTIPLE SCLEROSIS AGENTS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – OPIOID DEPENDENCE TREATMENTS

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – OPIOID REVERSAL AGENTS

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – PULMONARY/ ARTERIAL HYPERTENSION AGENTS

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – PEDIATRIC VITAMIN PREPARATIONS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Rose

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – PHOSPHATE BINDERS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – PRENATAL VITAMINS

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Cowles

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – PROTON PUMP INHIBITORS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Pohl

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – SINUS NODE INHIBITORS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – SKELETAL MUSCLE RELAXANTS

**(1<sup>st</sup>) Motion:** Pohl

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

## Review Agenda – TETRACYCLINES

**(1<sup>st</sup>) Motion:** Hill

**(2<sup>nd</sup>) Motion:** Dolter

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	Yes	No	Abstain	<b>Voting – P&amp;T Committee Members</b>	Yes	No	Abstain
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						



## Review Agenda – ULCERATIVE COLITIS

**(1<sup>st</sup>) Motion:** Cowles

**(2<sup>nd</sup>) Motion:** Hill

**Discussion:** Approved as written.

<b>Voting – P&amp;T Committee Members</b> <b>Does not include excused or unexcused members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>	<b>Voting – P&amp;T Committee Members</b>	<b>Yes</b>	<b>No</b>	<b>Abstain</b>
Bendlin, Andrew, Pharm.D.	X			Klug, Laura, Pharm.D.	X		
Cowles, Cassie, APRN	X			Pohl, Jessica, Pharm.D.	X		
Dering-Anderson, Allison, Pharm.D. (Chair) <b><i>Votes only in the event of a tie</i></b>				Rose, Steven, D.O.	X		
Dolter, Stephen, M.D.	X			Sundsboe, Bradley, Pharm.D.	X		
Hill, Jennifer, M.D.	X						

- c. Complete Copy of Proposed PDL

# Nebraska Medicaid Preferred Drug List

## with Prior Authorization Criteria

May 2025 P&T Proposed Changes

*Highlights* indicate proposed changes



Jim Pillen, Governor

For the most up to date list of covered drugs consult the **Drug Lookup** on the Nebraska Medicaid website at <https://ne.primetherapeutics.com/drug-lookup>.

- **PDMP Check Requirements** – Nebraska Medicaid providers are required to check the prescription drug history in the statewide PDMP before prescribing CII controlled substances to certain Medicaid beneficiaries (exemption to this requirement are for beneficiaries receiving cancer treatment, hospice/palliative care, or in long-term care facilities). If not able to check the PDMP, then provider is required to document good faith effort, including reasons why unable to conduct the check and may be required to submit documentation to the State upon request.
  - PDMP check requirements are under Section 5042 of the SUPPORT for Patients and Communities Act, consistent with section 1944 of the Social Security Act [42 U.S.C. 1396w-3a].
- **Opioids** – The maximum opioid dose covered is 90 Morphine Milligram Equivalents (MME) per day.

### Non-Preferred Drug Coverage

Class and drug-specific therapeutic trial and failure requirements are found within this document. Examples of non-preferred exception criteria include:

- Adverse reaction to preferred drugs
- Allergy to preferred drugs
- Contraindication to preferred drugs
- Documentation of inability to swallow solid dosage forms

Specific Class Prior Authorization forms can be found within the PDL class listings and at: <https://nebraska.fhsc.com/priorauth/paforms.asp>

- [Immunomodulators Self-Injectable PA Form](#)
- [Opioid Dependence Treatment PA Form](#)
- [Opioid Dependence Treatment Informed Consent](#)
- [Growth Hormone PA Form](#)
- [HAE Treatments PA Form](#)
- [Hepatitis C PA Form](#)

For all other class medically-necessary coverage, quantity, and high dose requests use the following:

[Documentation of Medical Necessity PA Form](#)

## ACNE AGENTS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
adapalene (generic Differin) <b>GEL (OTC/Rx), GEL PUMP</b> adapalene/BPO (generic Epiduo) benzoyl peroxide (BPO) <b>WASH, LOTION</b> benzoyl peroxide <b>GEL OTC</b> clindamycin/BPO (generic BenzaClin) <b>GEL, PUMP</b> clindamycin/BPO (generic Duac) clindamycin phosphate <b>PLEDGET</b> clindamycin phosphate <b>SOLUTION</b> erythromycin <b>GEL</b> erythromycin <b>SOLN</b> erythromycin-BPO (generic for Benzamycin) RETIN-A (tretinoin) <sup>AL</sup> <b>CREAM, GEL</b>	adapalene (generic Differin) <b>CREAM</b> adapalene/BPO (generic Epiduo Forte) ALTRENO (tretinoin) <sup>AL</sup> AMZEEQ (minocycline) ARAZLO (tazarotene) <sup>AL</sup> ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) AZELEX (azelaic acid) BENZEFOAM (benzoyl peroxide) benzoyl peroxide <b>CLEANSER, CLEANSING BAR OTC</b> benzoyl peroxide <b>FOAM</b> (generic BenzePro) benzoyl peroxide <b>GEL Rx</b> benzoyl peroxide <b>TOWELETTE OTC</b> CABTREO (clindamycin phosphate/BPO/adapalene) <sup>AL</sup> <b>GEL</b> clindamycin <b>FOAM, LOTION</b> clindamycin <b>GEL</b> clindamycin phosphate (generic for Clindagel) <b>GEL</b> clindamycin/BPO (generic Acanya) <b>GEL</b> clindamycin/BPO <b>PUMP</b> (generic Onexton) <sup>AL</sup> clindamycin/tretinoin (generic Veltin, Ziana) dapsone (generic Aczone) <b>DIFFERIN (adapalene) CREAM, LOTION, GEL-OTC, GEL PUMP</b> erythromycin <b>PLEDGET</b> EVOCLIN (clindamycin)	■ Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class

## ACNE AGENTS, TOPICAL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	<p>FABIOR (tazarotene) <b>FOAM</b></p> <p>NEUAC (clindamycin/BPO)</p> <p>ONEXTON (clindamycin/BPO)</p> <p>OVACE PLUS (sulfacetamide sodium)</p> <p>RETIN-A MICRO (tretinoin)</p> <p>sulfacetamide</p> <p>sulfacetamide/sulfur</p> <p>sulfacetamide sodium/ sulfur</p> <p><b>CLEANSER</b></p> <p>SUMADAN (sulfacetamide/sulfur)</p> <p>tazarotene (generic Tazorac) <b>CREAM</b></p> <p>tazarotene <b>FOAM</b> (generic Fabior)</p> <p>tazarotene <b>GEL</b> (generic Tazorac)</p> <p>TRETIN-X (tretinoin)</p> <p>tretinoin (generic Avita, Retin-A) <sup>AL</sup></p> <p><b>CREAM, GEL</b></p> <p>tretinoin microspheres (generic Retin-A Micro) <sup>AL</sup> <b>GEL, GEL PUMP</b></p> <p>WINLEVI (clascoterone)<sup>AL</sup></p>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class</li> </ul>

## ANALGESICS, OPIOID LONG-ACTING

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BUTRANS (buprenorphine) <sup>QL</sup> <b>PATCH</b> fentanyl 25, 50, 75, 100 mcg <b>PATCH</b> <sup>QL</sup> morphine ER <b>TABLET</b> (generic MS Contin, Oramorph SR) OXYCONTIN <sup>CL</sup> (oxycodone ER) tramadol ER (generic Ultram ER) <sup>CL</sup>	BELBUCA (buprenorphine) <sup>QL</sup> <b>BUCCAL</b> buprenorphine PATCH (generic Butrans) <sup>QL</sup> fentanyl 37.5/62.5/87.5 mcg <b>PATCH</b> <sup>QL</sup> hydrocodone ER (generic Hysingla ER) <sup>QL</sup> hydrocodone bitartrate ER (generic Zohydro ER) hydromorphone ER (generic Exalgo) <sup>CL</sup> HYSINGLA ER (hydrocodone ER) methadone <b>TABLET</b> <sup>CL</sup> methadone <b>ORAL SYR</b> <sup>CL</sup> <b>methadone SOL TABLET</b> morphine ER (generic Avinza, Kadian) <b>CAPS</b> oxycodone ER (generic Oxycontin) oxymorphone ER (generic Opana ER) tramadol ER (generic ConZip) <sup>CL</sup>	<p>The Center for Disease Control (CDC) does not recommend long-acting opioids when beginning opioid treatment.</p> <ul style="list-style-type: none"> <li>Preferred agents require previous use of a long-acting opioid or documentation of a trial on a short acting agent within 90 days</li> <li>Non-preferred agents will be approved with failure on, or intolerance to TWO preferred agents within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Methadone:</b> Will only be approved for use in pain control or end of life care. Trial of preferred agent not required for end-of-life care</li> <li><b>Oxycontin®:</b> Pain contract required for maximum quantity authorization</li> </ul>

## ANALGESICS, OPIOID SHORT-ACTING <sup>QL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>ORAL</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the last 12 months</li> <li>Note: for short acting opiate tablets and capsules there is a maximum quantity limit of #150 per 30 days.</li> <li>Opiate limits for opiate naïve patients will consist of: <ul style="list-style-type: none"> <li>-prescriptions limited to a 7 day supply, AND</li> <li>-initial opiate prescription fill limited to maximum of 50 Morphine Milligram Equivalents (MME) per day</li> </ul> </li> <li>These limits may only be exceeded with patient specific documentation of medical necessity, with examples such as, cancer diagnosis, end-of-life care, palliative care, Sickle Cell Anemia, or prescriber attestation that patient is not recently opiate naive</li> </ul>
acetaminophen/codeine <b>ELIXIR, TAB</b> codeine <b>TAB</b> hydrocodone/APAP <b>SOLN, TAB</b> hydrocodone/ibuprofen hydromorphone <b>TAB</b> morphine <b>CONC SOLN, DISP SYR SOLN, IR-TAB</b> oxycodone <b>TAB, SOLN</b> oxycodone/APAP tramadol 50 <b>TAB<sup>AL</sup></b> (generic Ultram)	butalbital/caffeine/APAP/codeine butalbital compound w/codeine (butalbital/ASA/caffeine/codeine) carisoprodol compound-codeine (carisoprodol/ASA/codeine) dihydrocodeine/APAP/caffeine hydromorphone <b>LIQUID, SUPPOSITORY</b> (generic Dilaudid) levorphanol meperidine (generic Demerol) morphine <b>SUPPOSITORIES</b> NALOCET (oxycodone/APAP) oxycodone <b>CAPS</b> oxycodone/APAP <b>SOLN</b> oxycodone <b>CONCENTRATE</b> oxymorphone IR (generic Opana) pentazocine/naloxone ROXICODONE (oxycodone) ROXYBOND (oxycodone) SEGLENTIS (celecoxib/tramadol) <sup>AL</sup> tramadol 25mg tramadol 75mg <sup>NR</sup> tramadol 100mg (generic Ultram) <sup>AL</sup> tramadol (generic Qdolo) <sup>AL,QL</sup> <b>SOLN</b> tramadol/APAP (generic Ultracet)	

## ANALGESICS, OPIOID SHORT-ACTING <sup>QL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NASAL		
	butorphanol <b>SPRAY</b> <sup>QL</sup>	
BUCCAL/TRANSMUCOSAL <sup>CL</sup>		
	fentanyl <b>TRANSMUCOSAL</b> (generic Actiq) <sup>CL</sup> FENTORA (fentanyl) <sup>CL</sup>	

## ANDROGENIC AGENTS (TOPICAL) <sup>CL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANDROGEL (testosterone) <b>PUMP</b> <sup>CL</sup> testosterone <b>PUMP</b> (generic Androgel) <sup>CL</sup> TESTIM (testosterone) <b>TRANSDERMAL</b>	NATESTO (testosterone) <sup>CL</sup> testosterone PACKET (generic Androgel) <sup>CL</sup> testosterone <b>GEL, PACKET, PUMP</b> (generic Vogelxo) testosterone (generic Axiron) testosterone (generic Fortesta) testosterone (generic Testim)	<ul style="list-style-type: none"> <li>▪ Preferred agents approved for diagnosis of Primary hypogonadism (congenital or acquired) or Hypogonadotropic hypogonadism. Off label use for the following will be considered with documentation of necessity: female to male transsexual – gender dysphoria, weight gain, male osteoporosis, delayed puberty in males, corticosteroid-induced hypogonadism and osteoporosis, inoperable carcinoma of the breast, postpartum breast pain and engorgement, and menopause</li> <li>▪ In addition, non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 6 months</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li>▪ <b>Androgel®:</b> Approved for Males only</li> <li>▪ <b>Natesto®:</b> Approved for Males only with diagnosis of: Primary hypogonadism (congenital or acquired) OR Hypogonadotropic hypogonadism (congenital or acquired)</li> </ul>

## ANGIOTENSIN MODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ACE INHIBITORS		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed TWO preferred agents within this drug class within the last 12 months</li><li>Non-preferred combination products may be covered as individual prescriptions without prior authorization</li></ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"><li><b>Epaned/enalapril oral solution/Qbreilis oral solution:</b> Clinical reason why oral tablet is not appropriate</li></ul>
benazepril (generic Lotensin)	captopril (generic Capoten)	
enalapril (generic Vasotec)	EPANED (enalapril) <sup>CL</sup> <b>ORAL SOLN</b>	
lisinopril (generic Prinivil, Zestril)	enalapril (generic for Epaned) <sup>CL</sup> <b>ORAL SOLN</b>	
ramipril (generic Altace)	fosinopril (generic Monopril)	
	moexepiril (generic Univasc)	
	perindopril (generic Aceon)	
	QBRELIS (lisinopril) <sup>CL</sup> <b>ORAL SOLN</b>	
	quinapril (generic Accupril)	
	trandolapril (generic Mavik)	
ACE INHIBITOR/DIURETIC COMBINATIONS		
enalapril/HCTZ (generic Vaseretic)	benazepril/HCTZ (generic Lotensin HCT)	
lisinopril/HCTZ (generic Prinzide, Zestoretic)	captopril/HCTZ (generic Capozide)	
	fosinopril/HCTZ (generic Monopril HCT)	
	moexipril/HCTZ (generic Uniretic)	
	quinapril/HCTZ (generic Accuretic)	
ANGIOTENSIN RECEPTOR BLOCKERS		
irbesartan (generic Avapro)	candesartan (generic Atacand)	
losartan (generic Cozaar)	EDARBI (azilsartan)	
olmesartan (generic Benicar)	eprosartan (generic Teveten)	
valsartan (generic Diovan)	telmisartan (generic Micardis)	



## ANGIOTENSIN MODULATORS (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>ANGIOTENSIN RECEPTOR BLOCKER/DIURETIC COMBINATIONS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed TWO preferred agents within this drug class within the last 12 months</li> <li>Non-preferred combination products may be covered as individual prescriptions without prior authorization</li> </ul>
irbesartan/HCTZ (generic Avalide)	candesartan/HCTZ (generic Atacand-HCT)	
losartan/HCTZ (generic Hyzaar)	EDARBYCLOR (azilsartan/chlorthalidone)	
olmesartan/HCTZ (generic Benicar-HCT)	telmisartan/HCTZ (generic Micardis-HCT)	
valsartan/HCTZ (generic Diovan-HCT)		
<b>ANGIOTENSIN MODULATOR/ CALCIUM CHANNEL BLOCKER COMBINATIONS</b>		<ul style="list-style-type: none"> <li><b>Direct Renin Inhibitors/Direct Renin Inhibitor Combinations:</b> May be approved with a history of TWO preferred ACE Inhibitors or Angiotensin Receptor Blockers within the last 12 months</li> </ul> <p>Drug Specific Criteria</p> <ul style="list-style-type: none"> <li><b>Entresto/ sacubitril-valsartan:</b> May be approved in patients ages &gt;1 years old and with a diagnosis of heart failure</li> </ul>
amlodipine/benazepril (generic Lotrel)	amlodipine/olmesartan/HCTZ (generic Tribenzor)	
amlodipine/olmesartan (generic Azor)	amlodipine/telmisartan (generic Twynsta)	
amlodipine/valsartan (generic Exforge)	amlodipine/valsartan/HCTZ (generic Exforge HCT)	
	PRESTALIA (perindopril/amlodipine)	
	trandolapril/verapamil (generic Tarka)	
<b>DIRECT RENIN INHIBITORS</b>		
	aliskiren (generic Tekturna) <sup>QL</sup>	
<b>DIRECT RENIN INHIBITOR COMBINATIONS</b>		
	TEKTURNA/HCTZ (aliskiren/HCTZ)	
<b>NEPRILYSIN INHIBITOR COMBINATION</b>		
ENTRESTO (sacubitril/valsartan) <sup>CL,QL</sup>	ENTRESTO (sacubitril/valsartan) <sup>CL,NR,QL</sup> SPRINKLE CAP	

## ANTIBIOTICS, GASTROINTESTINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
metronidazole <b>TABLET</b>	AEMCOLO (rifamycin) <b>TAB</b>	<ul style="list-style-type: none"> <li>Note: Although azithromycin, ciprofloxacin, and trimethoprim/ sulfmethoxazole are not included in this review, they are available without prior authorization</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Alinia /nitazoxanide tablet:</b> Trial and failure with metronidazole is required for a diagnosis of giardiasis</li> <li><b>Dificid®:</b> For diagnosis of C. difficile diarrhea (pseudomembranous colitis), trial and failure or intolerance to oral vancomycin is required. For diagnosis of relapsed or recurrent C. difficile, an appropriate ICD-10 diagnosis code must be submitted for coverage.</li> <li><b>Flagyl®/Metronidazole 375mg capsules and / Metronidazole 750mg ER tabs:</b> Clinical reason why the generic regular release cannot be used</li> <li><b>tinidazole:</b> Approvable diagnoses include: <ul style="list-style-type: none"> <li>Giardia</li> <li>Amebiasis intestinal or liver abscess</li> <li>Bacterial vaginosis or trichomoniasis</li> </ul> </li> <li><b>vancomycin capsules:</b> Requires patient specific documentation of why preferred vancomycin solution is not appropriate for patient</li> <li><b>Xifaxan®:</b> Approvable diagnoses include: <ul style="list-style-type: none"> <li>Travelers's diarrhea resistant to quinolones</li> <li>Hepatic encephalopathy with treatment failure of lactulose or neomycin</li> <li>Diarrhea-Predominant IBS (IBS-D) 550mg strength only with treatment failure of Lomotil® AND Imodium®</li> </ul> </li> </ul>
neomycin	DIFICID (fidaxomicin) <sup>CL</sup> <b>TAB, SUSP</b>	
tinidazole (generic Tindamax) <sup>CL</sup>	<b>FIRVANQ (vancomycin)<sup>QL</sup> SOLN</b>	
<b>vancomycin (generic Firvanq)<sup>QL</sup> SOLN</b>	LIKMEZ (metronidazole) <b>SUSP</b>	
	metronidazole <sup>CL</sup> <b>CAPS</b>	
	<b>metronidazole 125mg<sup>NR</sup> TAB</b>	
	nitazoxanide	
	(generic Alinia) <b>TAB<sup>AL, CL, QL</sup></b>	
	paromomycin	
	SOLOSEC (secnidazole)	
	vancomycin <b>CAPS</b> (generic Vancocin) <sup>CL</sup>	
	VOWST (fecal microbiota spores) <sup>AL, QL</sup>	
	XIFAXAN (rifaximin) <sup>CL</sup>	

## ANTIBIOTICS, INHALED<sup>CL</sup>

Preferred Agents <sup>CL</sup>	Non-Preferred Agents	Prior Authorization/Class Criteria
BETHKIS (tobramycin) KITABIS PAK (tobramycin) tobramycin (generic Tobi)	ARIKAYCE (amikacin liposomal inh) <b>SUSP</b> CAYSTON (aztreonam lysine) <sup>QL</sup> <b>TOBI-PODHALER (tobramycin)<sup>QL</sup></b> tobramycin (generic Bethkis)	<ul style="list-style-type: none"> <li>Diagnosis of Cystic Fibrosis is required for all agents ICD10 Group = E84, ICD9 = 277.00, 277.01, 277.02, 277.03, 277.09</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Arikayce:</b> Requires diagnosis of refractory MAC lung disease defined as patients who did not achieve negative sputum cultures after a minimum of 6 consecutive months of a multidrug background regimen therapy</li> <li><b>Cayston®:</b> Trial of tobramycin via nebulizer and demonstration of TOBI® compliance required</li> <li><b>Tobi Podhaler®:</b> Requires trial of tobramycin via nebulizer or documentation of why nebulized tobramycin cannot be used</li> </ul>

## ANTIBIOTICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
bacitracin <b>OINT</b> bacitracin <b>OINT OTC</b> bacitracin/polymyxin (generic Polysporin) mupirocin <b>OINT</b> (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/pramoxine	bacitracin <b>PACKET-OTC</b> gentamicin <b>OINT, CREAM</b> mupirocin <b>CREAM</b> (generic Bactroban) <sup>CL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class within the last 12 months</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Mupirocin® Cream:</b> Clinical reason the ointment cannot be used</li> </ul>

## ANTIBIOTICS, VAGINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CLEOCIN (clindamycin) <b>CREAM</b> <b>OVULES</b> metronidazole, <b>VAGINAL</b> NUVESSA (metronidazole)	clindamycin (generic Cleocin) <b>CREAM</b> CLINDESSE (clindamycin) VANDAZOLE (metronidazole) XACIATO (clindamycin phosphate) <b>GEL</b> <sup>AL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a therapeutic trial (duration = 3 days) with ONE preferred agent within this drug class within the last 6 months</li> </ul>

## ANTICOAGULANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
dabigatran etexilate (generic Pradaxa) <b>CAPS</b> ELIQUIS (apixaban) enoxaparin (generic Lovenox) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg <sup>CL,QL</sup> XARELTO DOSE PACK (rivaroxaban)	fondaparinux (generic Arixtra) FRAGMIN (dalteparin) PRADAXA (dabigatran) <b>CAPS</b> , <b>PELLETS</b> SAVAYSA (edoxaban) <sup>CL,QL</sup> XARELTO (rivaroxaban) <sup>CL</sup> <b>SUSP</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 12 months</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Coumadin®</b>: Clinical reason generic warfarin cannot be used</li> <li><b>Savaysa®</b>: Approved diagnoses include:            Stroke and systemic embolism (SE) risk reduction in nonvalvular atrial fibrillation (NVAf) OR            Treatment of deep vein thrombosis (DVT) and pulmonary embolism (PE) following 5-10 days of parenteral anticoagulant therapy</li> <li><b>Xarelto 2.5mg</b>: Use limited to reduction of risk of major cardiovascular events (cardiovascular death, myocardial infarction, and stroke) in patients with chronic coronary artery disease or peripheral artery disease</li> <li><b>Xarelto Suspension</b>: Approved for patients ≤12 years of age or if there is a clinical reason why a preferred solid dosage form cannot be used.</li> </ul>

## ANTIEMETICS/ANTIVERTIGO AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>CANNABINOIDS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the same group</li> </ul>
dronabinol (generic Marinol) <sup>AL</sup>		
<b>5HT3 RECEPTOR BLOCKERS</b>		<p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Akynzeo®:</b> Approved for Moderately/Highly emetogenic chemotherapy with dexamethasone and a 5-HT3 antagonist</li> <li><u>Regimens include:</u> AC combination (Doxorubicin or Epirubicin with Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide, Azacitidine, Bendamustine, Busulfan, Carmustine, Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine, Dacarbazine, Dactinomycin, Daunorubicin, Epirubicin, Etoposide, Hexamethylmelamine, Idarubicin, Ifosfamide, Imatinib, Interferon <math>\alpha</math>, Irinotecan, Mechlorethamine, Melphalan, Methotrexate, Oxaliplatin, Procarbazine, Streptozotocin, Temozolomide</li> <li><b>Diclegis/doxylamine-pyridoxine/ Bonjesta:</b> Approved only for treatment of nausea and vomiting of pregnancy</li> <li><b>Sancuso®:</b> Documentation of oral dosage form intolerance</li> </ul>
ondansetron (generic Zofran/Zofran ODT) <sup>QL</sup>	ANZEMET (dolasetron) granisetron (generic Kytril) <b>ondansetron 16mg ODT (generic Zofran ODT)<sup>NR</sup></b> SANCUSO (granisetron) <sup>CL</sup>	
<b>NK-1 RECEPTOR ANTAGONIST</b>		
aprepitant (generic Emend) <b>CAPS</b> <sup>QL</sup>	AKYNZEO (netupitant/palonosetron) <sup>CL</sup> aprepitant (generic Emend) <b>PACK</b> EMEND (aprepitant) <b>CAPS, PACK, POWDER</b> <sup>QL</sup>	
<b>TRADITIONAL ANTIEMETICS</b>		
DICLEGIS (doxylamine/pyridoxine) <sup>CL,QL</sup>	BONJESTA (doxylamine/pyridoxine) <sup>CL,QL</sup>	
dimenhydrinate (generic Dramamine) <b>OTC</b>	COMPRO (prochlorperazine)	
meclizine (generic Antivert)	doxylamine/pyridoxine (generic Diclegis) <sup>CL,QL</sup>	
metoclopramide (generic Reglan)	prochlorperazine <b>SUPPOSITORY</b> (generic Compazine)	
phosphoric acid/dextrose/fructose (generic Emetrol) <b>SOLN</b>	promethazine <b>SUPPOSITORY</b> 50mg	
prochlorperazine (generic Compazine)	trimethobenzamide <b>TAB</b> (generic Tigan)	
promethazine (generic Phenergan) <b>SYRUP, TAB</b>		
promethazine 12.5mg, 25mg <b>SUPPOSITORY</b>		
scopolamine (generic Transderm-Scop) <b>TRANSDERMAL</b>		
<b>TRANSDERM-SCOP (scopolamine) TRANSDERMAL</b>		

## ANTIFUNGALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clotrimazole (mucous membrane, troche)	BREXAFEMME (ibrexafungerp) <sup>QL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class</li> </ul>
fluconazole <b>SUSP, TAB</b> (generic Diflucan)	CRESEMBA (isavuconazonium) <sup>CL</sup>	
griseofulvin <b>SUSP</b>	flucytosine (generic Ancobon) <sup>CL</sup>	Drug-specific criteria:
griseofulvin microsize (generic GRIS-PEG)	griseofulvin ultramicrosize (generic GRIS-PEG)	<ul style="list-style-type: none"> <li><b>Cresemba®:</b> Approved for diagnosis of invasive aspergillosis or invasive mucormycosis</li> </ul>
nystatin <b>SUSP</b>	itraconazole (generic Sporanox) <sup>CL</sup>	<ul style="list-style-type: none"> <li><b>Flucytosine:</b> Approved for diagnosis of: <u>Candida</u>: Septicemia, endocarditis, UTIs <u>Cryptococcus</u>: Meningitis, pulmonary infections</li> </ul>
terbinafine (generic Lamisil)	ketoconazole (generic Nizoral)	<ul style="list-style-type: none"> <li><b>Noxafil/ posaconazole DR tablets, oral suspension, PowderMix® for delayed oral suspension: For prophylaxis of invasive Aspergillus and Candida infections, no preferred agent</b> trial is required in severely immunocompromised patients (i.e., Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropenic hematologic malignancies, Graft vs. Host disease (GVHD), Immunosuppression secondary to hematopoietic stem cell transplant</li> </ul>
	ORAVIG (miconazole) <sup>QL</sup> <b>BUCCAL</b>	<ul style="list-style-type: none"> <li><b>Noxafil® Powdermix:</b> pediatric patients 2 years of age and older who weigh 40 kg or less</li> </ul>
	NOXAFIL (posaconazole) <sup>AL</sup> <b>TAB</b>	
	NOXAFIL (posaconazole) <sup>AL,CL</sup>	<ul style="list-style-type: none"> <li><b>Noxafil/ posaconazole Suspension:</b> Oropharyngeal/esophageal candidiasis refractory to itraconazole and/or fluconazole and;</li> </ul>
	<b>POWDERMIX</b>	Prophylaxis of invasive Aspergillus and Candida infections
	nystatin <b>TAB</b>	<ul style="list-style-type: none"> <li><b>Sporanox®/itraconazole:</b> Approved for diagnosis of Aspergillosis, Blastomycosis, Histoplasmosis, Onychomycosis due to terbinafine-resistant dermatophytes, Oropharyngeal/ esophageal candidiasis refractory to fluconazole</li> </ul>
	posaconazole (generic Noxafil) <sup>AL,CL</sup>	
	TOLSURA (itraconazole) <sup>CL</sup>	<ul style="list-style-type: none"> <li><b>Sporanox® Liquid:</b> Clinical reason solid oral cannot be used</li> </ul>
	VIVJOA (oteseconazole) <b>CAPS</b>	<ul style="list-style-type: none"> <li><b>Tolsura:</b> Approved for diagnosis of Aspergillosis, Blastomycosis, and Histoplasmosis and requires a trial and failure of generic itraconazole</li> </ul>
	voriconazole (generic VFEND) <sup>CL</sup>	<ul style="list-style-type: none"> <li><b>Vfend/voriconazole:</b> No trial for diagnosis of Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Graft vs. Host disease (GVHD), Candidemia (candida krusei), Esophageal Candidiasis, Blastomycosis, <i>S. apiospermum</i> and <i>Fusarium spp.</i>, Oropharyngeal/esophageal candidiasis refractory to fluconazole</li> </ul>

## ANTIFUNGALS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTIFUNGAL		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class within the last 6 months</li></ul> Drug-specific criteria: <ul style="list-style-type: none"><li><b>Extina/ Ketodan/ ketoconazole foam:</b> Requires trial and failure or contraindication to other ketoconazole forms</li><li><b>Jublia and tavaborole:</b> Approved diagnoses include Onychomycosis of the toenails due to <i>T.rubrum</i> OR <i>T. Mentagrophytes</i></li><li><b>ciclopirox nail lacquer:</b> No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine</li></ul>
clotrimazole <b>CREAM</b> (generic Lotrimin) RX, OTC	ALEVAZOL (clotrimazole) OTC	
clotrimazole <b>SOLN RX</b> (generic Lotrimin)	ciclopirox <b>CREAM, GEL, SUSP</b> (generic Ciclodan, Loprox)	
ketoconazole <b>CREAM, SHAMPOO</b> (generic Nizoral)	ciclopirox <b>SHAMPOO</b> (generic Loprox)	
miconazole <b>CREAM, POWDER</b> OTC	clotrimazole <b>SOLN OTC</b>	
nystatin	DESENEX <b>POWDER OTC</b> (miconazole)	
terbinafine OTC (generic Lamisil AT)	econazole (generic Spectazole)	
tolnaftate <b>AERO-POWDER OTC, CREAM-OTC, SOLN-OTC</b> (generic Tinactin)	ERTACZO (sertaconazole)	
	FUNGOID (miconazole) <b>OTC</b>	
	JUBLIA (efinaconazole) <sup>CL</sup>	
	ketoconazole <b>FOAM</b> <sup>CL</sup> (generic Extina, Ketodan)	
	LOPROX (ciclopirox) <b>CREAM, SUSP</b>	
	LOTRIMIN AF <b>CREAM</b> OTC (clotrimazole)	
	LOTRIMIN ULTRA (butenafine)	
	luliconazole (generic Luzu)	
	miconazole OTC <b>OINT, SPRAY, SOLN</b>	
	miconazole/zinc oxide/petrolatum (generic Vusion)	
	naftifine <b>CREAM, GEL</b> (generic Naftin)	
	oxiconazole (generic Oxistat)	
	tavaborole <b>SOLN</b> <sup>CL</sup> (generic Kerydin)	
	tolnaftate <b>POWDER OTC</b>	
	TRIPENICOL (undecylenic acid) <sup>NR</sup> <b>CREAM OTC</b>	
	VOTRIZA-AL (clotrimazole) <b>LOTION OTC</b>	
ANTIFUNGAL/STEROID COMBINATIONS		
clotrimazole/betamethasone <b>CREAM</b> (generic Lotrisone)	clotrimazole/betamethasone <b>LOTION</b> (generic Lotrisone)	
nystatin/triamcinolone (generic Mycolog)		
<b>CREAM, OINT</b>		

## ANTIMIGRAINE AGENTS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AIMOVIG (erenumab-aooe) <sup>CL, QL</sup> AJOVY (fremanezumab-vfrm) <sup>CL, QL</sup> <b>PEN, Autoinjector</b> AJOVY (fremanezumab-vfrm) <b>Autoinjector 3-pack</b> <sup>CL, QL</sup> EMGALITY 120 mg/mL (galcanezumab-gnlm) <sup>CL, QL</sup> <b>PEN, SYRINGE</b> NURTEC ODT (rimegepant) <sup>AL, CL, QL</sup> QULIPTA (atogepant) <sup>AL, CL, QL</sup> UBRELVY (ubrogepant) <sup>AL, CL, QL</sup> <b>TAB</b>	diclofenac (generic Cambia) <b>POWDER</b> dihydroergotamine mesylate <b>NASAL</b> ELYXYB (celecoxib) <sup>AL, QL</sup> <b>SOLN</b> EMGALITY 100 mg (galcanezumab-gnlm) <sup>CL, QL</sup> <b>SYR</b> MIGERGOT (ergotamine/cafeine) <b>RECTAL</b> MIGRANAL (dihydroergotamine) <b>NASAL</b> REYVOW (lasmiditan) <sup>AL, CL, QL</sup> <b>TAB</b> ZAVZPRET (zavegepant) <sup>AL, QL</sup> <b>NASAL</b>	<ul style="list-style-type: none"> <li>All non-preferred agents will require a failed trial or contraindication of a preferred agent of the same indication</li> <li><b>For Acute Treatment:</b> agents will be approved for patients who have a failed trial or a contraindication to two triptans.</li> <li><b>For Prophylactic Treatment:</b> Require <math>\geq 4</math> migraines per month for <math>\geq 3</math> months and has tried and failed a <math>\geq 1</math> month trial of two medications listed in the 2012 American Academy of Neurology/American Headache Society guidelines (examples include: antidepressants (amitriptyline, venlafaxine), Beta blockers (propranolol, metoprolol, atenolol), anti-epileptics (valproate, topiramate), ACE (lisinopril)</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Emgality 100mg</b> will only be approved for treatment of Episodic Cluster Headache</li> <li><b>Nurtec ODT:</b> for use in acute treatment, will be approved for patients who have a failed trial or a contraindication to two triptans. For use in preventative treatment, will be approved for patients who have a failed trial of ONE preferred injectable CGRP</li> <li><b>Qulipta:</b> May be approved for patients who have a failed trial of ONE preferred injectable CGRP</li> </ul>



## ANTIMIGRAINE AGENTS, TRIPTANS<sup>QL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ORAL		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class</li></ul> Drug-specific criteria: <ul style="list-style-type: none"><li><b>Zembrace:</b> approved for patients who have failed ALL preferred agents</li></ul>
rizatriptan (generic Maxalt)	almotriptan (generic Axert)	
rizatriptan ODT (generic Maxalt MLT)	eletriptan (generic Relpax)	
sumatriptan	frovatriptan (generic Frova)	
	IMITREX (sumatriptan)	
	naratriptan (generic Amerge)	
	RELPAK (eletriptan) <sup>QL</sup>	
	sumatriptan/naproxen (generic Treximet)	
	zolmitriptan (generic Zomig)	
NASAL		
sumatriptan (generic Imitrex Nasal)	TOSYMRA (sumatriptan)	
	zolmitriptan (generic Zomig)	
INJECTABLE		
sumatriptan <b>SYRINGE, VIAL</b>	sumatriptan <b>KIT</b>	
	ZEMBRACE SYMTOUCH (sumatriptan)	

## ANTIPARASITICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NATROBA (spinosad)	CROTAN (crotamiton) <b>LOTION</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class within the past 6 months</li> </ul>
permethrin 1% OTC (generic Nix)	EURAX (crotamiton) <b>CREAM, LOTION</b>	
permethrin 5% RX (generic Elimite)	ivermectin (generic Sklice) <b>LOTION</b>	
pyrethrin/piperonyl butoxide <b>SHAMPOO</b>	malathion (generic Ovide)	
(generic RID, A-200)	spinosad (generic Natroba)	
	VANALICE (piperonyl butoxide/pyrethrins)	

## ANTIVIRALS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTI-HERPETIC DRUGS		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed a 10-day trial of ONE preferred agent within the same group</li></ul>
acyclovir (generic Zovirax)	acyclovir (generic for Zovirax) <sup>CL</sup> <b>SUSP</b>	
famciclovir (generic Famvir)		
valacyclovir (generic Valtrex)		
ANTI-INFLUENZA DRUGS		Drug-specific criteria:
oseltamivir (generic Tamiflu) <sup>QL</sup> <b>CAPS, SUSP</b>	rimantadine (generic Flumadine)	<ul style="list-style-type: none"><li><b>Acyclovir Susp:</b> Prior authorization NOT required for children ≤ 12 years old</li><li><b>Xofluza:</b> Requires clinical, patient specific reason that a preferred agent cannot be used</li><li><b>Paxlovid:</b> Requires a diagnosis of COVID-19 and is limited to 1 dose pack per 30 days</li><li></li></ul>
	RELENZA (zanamivir) <sup>QL</sup>	
	TAMIFLU (oseltamivir) <sup>QL</sup> <b>CAPS, SUSP</b>	
	XOFLUZA (baloxavir marboxil) <sup>AL,CL,QL</sup>	
ANTI-COVID-19 DRUGS		
PAXLOVID (nirmatrelvir and ritonavir) <sup>NR,QL</sup>		

## ANTIVIRALS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acyclovir <b>OINT</b>	acyclovir CREAM, (generic Zovirax)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred ORAL Antiviral agent</li> </ul>
docosanol <b>OTC</b>	DENAVIR (penciclovir) <sup>AL</sup>	
	penciclovir (generic Denavir) <sup>AL</sup>	
	XERESE (acyclovir/hydrocortisone)	

## BETA BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>BETA BLOCKERS</b>		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed TWO diagnosis-appropriate preferred agents within this drug class</li></ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"><li><b>Coreg CR/carvedilol:</b> Requires clinical reason generic IR product cannot be used</li><li><b>Hemangeol®:</b> Covered for diagnosis of Proliferating Infantile Hemangioma</li><li><b>Sotylize®:</b> Covered for diagnosis of life –threatening ventricular arrhythmias OR maintenance of normal sinus rhythm in highly symptomatic atrial fibrillation/flutter (AFIB/AFL)</li></ul> <p>Requires clinical reason generic sotalol cannot be used</p>
atenolol (generic Tenormin)	acebutolol (generic Sectral)	
atenolol/chlorthalidone (generic Tenoretic)	betaxolol (generic Kerlone)	
bisoprolol (generic Zebeta)	<b>BYSTOLIC (nebivolol)</b>	
bisoprolol/HCTZ (generic Ziac)	INDERAL LA/XL (propranolol ER)	
HEMANGEOL (propranolol) <sup>AL</sup> <b>SOLN</b>	INNOPRAN XL (propranolol ER)	
metoprolol (generic Lopressor)	KAPSPARGO SPRINKLE (metoprolol ER)	
metoprolol ER (generic Toprol XL)	LOPRESSOR (metoprolol)	
nebivolol (generic Bystolic)	metoprolol/HCTZ (generic	
propranolol (generic Inderal) <b>TAB, SOLN</b>	Lopressor HCT)	
propranolol ER (generic Inderal LA)	nadolol (generic Corgard)	
	pindolol (generic Viskin)	
	propranolol/HCTZ (generic Inderide)	
	timolol (generic Blocadren)	
	TOPROL XL (metoprolol ER)	
<b>BETA- AND ALPHA-BLOCKERS</b>		
carvedilol (generic Coreg)	carvedilol ER <sup>CL</sup> (generic Coreg CR)	
labetalol (generic Trandate)		
<b>ANTIARRHYTHMIC</b>		
sotalol (generic Betapace)	SOTYLIZE (sotalol)	

## BLADDER RELAXANT PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
fesoterodine (generic Toviaz) MYRBETRIQ (mirabegron) <sup>AL</sup> <b>TAB</b> oxybutynin IR, ER (generic Ditropan/Ditropan XL)	darifenacin ER (generic Enablex) flavoxate HCL GEMTESA (vibegron) <sup>AL,QL</sup> <b>mirabegron ER TAB (generic</b> <b>Myrbetriq)<sup>NR</sup></b> MYRBETRIQ (mirabegron) <b>SUSP</b> <sup>AL,CL,QL</sup> oxybutynin 2.5mg OXYTROL (oxybutynin) <b>TRANSDERMAL</b> solifenacin (generic Vesicare) tolterodine IR, ER (generic Detrol/ Detrol LA) TOVIAZ (fesoterodine ER) trospium IR, ER (generic Sanctura/ Sanctura XR) VESICARE (solifenacin) VESICARE LS <b>SUSP</b> (solifenacin) <sup>AL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> </ul> Drug-specific criteria: <ul style="list-style-type: none"> <li><b>Myrbetriq suspension:</b> Covered for pediatric patients <math>\geq 3</math> years old with a diagnosis of Neurogenic Detrusor Overactivity (NDO)</li> </ul>

## BONE RESORPTION SUPPRESSION AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>BISPHOSPHONATES</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within the same group</li> <li>Drug-specific criteria: <ul style="list-style-type: none"> <li><b>Actonel® Combinations:</b> Covered as individual agents without prior authorization</li> <li><b>Atelvia DR®:</b> Requires clinical reason alendronate cannot be taken on an empty stomach</li> <li><b>Binosto®:</b> Requires clinical reason why alendronate tablets OR Fosamax® solution cannot be used</li> <li><b>Forteo/ teriparatide:</b> Covered for high risk of fracture High risk of fracture: <ul style="list-style-type: none"> <li>BMD -3 or worse</li> <li>Postmenopausal women with history of non-traumatic fractures</li> <li>Postmenopausal women with 2 or more clinical risk factors <ul style="list-style-type: none"> <li>Family history of non-traumatic fractures</li> <li>DXA BMD T-score <math>\leq</math> -2.5 at any site</li> <li>Glucocorticoid use <math>\geq</math> 6 months at 7.5 dose of prednisolone equivalent</li> <li>Rheumatoid Arthritis</li> </ul> </li> <li>Postmenopausal women with BMD T-score <math>\leq</math> -2.5 at any site with any clinical risk factors <ul style="list-style-type: none"> <li>More than 2 units of alcohol per day</li> <li>Current smoker</li> </ul> </li> <li>Men with primary or hypogonadal osteoporosis</li> <li>Osteoporosis associated with sustained systemic glucocorticoid therapy</li> <li>Trial of calcitonin-salmon not required</li> <li>Maximum of 24 months treatment per lifetime</li> </ul> </li> </ul> </li> </ul>
alendronate (generic Fosamax) <b>TAB</b> ibandronate (generic Boniva) <sup>QL</sup>	alendronate <b>SOLN</b> (generic Fosamax) <sup>QL</sup> ATELVIA DR (risedronate) BINOSTO (alendronate) FOSAMAX PLUS D <sup>QL</sup> risedronate (generic Actonel) <sup>QL</sup>	
<b>OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS</b>		
calcitonin-salmon <b>NASAL</b> FORTEO (teriparatide) <sup>CL,QL</sup> raloxifene (generic Evista)	EVISTA (raloxifene) teriparatide (generic Forteo) <sup>CL,QL</sup> TYMLOS (abaloparatide)	

## BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>ALPHA BLOCKERS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Alfuzosin/dutasteride/finasteride</b> <ul style="list-style-type: none"> <li>Covered for males only</li> </ul> </li> <li><b>Cardura XL®:</b> Requires clinical reason generic IR form cannot be used</li> <li><b>Flomax/ tamsulosin: Females covered for a 7-day supply with diagnosis of acute kidney stones.</b></li> <li><b>Jalyn/ dutasteride-tamsulosin:</b> Requires clinical reason why individual agents cannot be used</li> </ul>
alfuzosin (generic Uroxatral)	CARDURA XL (doxazosin)	
doxazosin (generic Cardura)	silodosin (generic Rapaflo)	
tamsulosin (generic Flomax)		
terazosin (generic Hytrin)		
<b>5-ALPHA-REDUCTASE (5AR) INHIBITORS</b>		
dutasteride (generic Avodart)	dutasteride/tamsulosin (generic Jalyn)	
finasteride (generic Proscar)		

## CALCIUM CHANNEL BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>SHORT-ACTING</b>		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li></ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"><li><b>Nifedipine:</b> May be approved without trial for diagnosis of Preterm Labor or Pregnancy Induced Hypertension (PIH)</li><li><b>Nimodipine:</b> Covered without trial for diagnosis of subarachnoid hemorrhage</li><li><b>Nimodipine solution:</b> Covered without trial for diagnosis of subarachnoid hemorrhage and; documented swallowing difficulty</li><li><b>Katerzia/ Norliqva:</b> May be approved with documented swallowing difficulty</li></ul>
<b>Dihydropyridines</b>		
	isradipine (generic Dynacirc)	
	nicardipine (generic Cardene)	
	nifedipine (generic Procardia)	
	nimodipine (generic Nimotop)	
	<b>nimodipine (generic Nymalize)<sup>NR</sup> SOLN</b>	
	<b>NYMALIZE (nimodipine) SOLN</b>	
<b>Non-dihydropyridines</b>		
diltiazem (generic Cardizem)		
verapamil (generic Calan/Isoptin)		
<b>LONG-ACTING</b>		
<b>Dihydropyridines</b>		
amlodipine (generic Norvasc)	felodipine ER (generic Plendil)	
nifedipine ER (generic Procardia XL/ Adalat CC)	<b>KATERZIA (amlodipine)<sup>QL</sup> SUSP</b>	
	levamlodipine (generic Conjupri)	
	nisoldipine (generic Sular)	
	<b>NORLIQVA (amolidipine)<sup>AL,CL,QL</sup> SOLN</b>	
<b>Non-dihydropyridines</b>		
diltiazem ER (generic Cardizem CD)	diltiazem ER (generic Cardizem LA)	
verapamil ER <b>TAB</b>	MATZIM LA (diltiazem ER)	
	TIAZAC (diltiazem)	
	verapamil ER <b>CAPS</b>	
	verapamil 360mg <b>CAPS</b>	
	verapamil ER (generic Verelan PM)	
	<b>verapamil SR (generic Verelan)<sup>NR</sup> CAPS</b>	

## CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>BETA LACTAM/BETA-LACTAMASE INHIBITOR COMBINATIONS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within the same group</li> </ul> <p>Drug Specific Criteria</p> <ul style="list-style-type: none"> <li><b>Cefixime</b>- May be approved for a diagnosis of gonorrhea, with an appropriate ICD-10 diagnosis code without a 3-day trial of a preferred agent</li> <li><b>Cefpodoxime</b>- May be approved for a diagnosis of pyelonephritis, with an appropriate ICD-10 diagnosis code without a 3-day trial of a preferred agent</li> </ul>
amoxicillin/clavulanate <b>TAB, SUSP</b>	amoxicillin/clavulanate <b>CHEWABLE</b>	
	amoxicillin/clavulanate ER (generic Augmentin XR)	
	AUGMENTIN (amoxicillin/clavulanate) <b>SUSP, TAB</b>	
<b>CEPHALOSPORINS – First Generation</b>		
cefadroxil <b>CAPS, SUSP</b> (generic Duricef)	cefadroxil <b>TAB</b> (generic Duricef)	
cephalexin <b>CAPS, SUSP</b>	cephalexin <b>TAB</b>	
<b>CEPHALOSPORINS – Second Generation</b>		
cefprozil (generic Cefzil)	cefaclor (generic Ceclor)	
cefuroxime <b>TAB</b> (generic Ceftin)		
<b>CEPHALOSPORINS – Third Generation</b>		
cefdinir (generic Omnicef)	cefixime (generic Suprax) <b>CAPS, SUSP</b>	
	cefpodoxime (generic Vantin)	



## CONTRACEPTIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<p>All reviewed agents are recommended preferred at this time</p> <p><i>Only those products for review are listed.</i></p> <p>Brand name products may be subject to Maximum Allowable Cost (MAC) pricing or require substitution with a generic equivalent</p> <p>Specific agents can be looked up using the Drug Look-up Tool at:</p> <p><a href="https://ne.primetherapeutics.com/drug-lookup">https://ne.primetherapeutics.com/drug-lookup</a></p> <p>EMZAHH (norethindrone)<sup>NR</sup></p> <p>FEIRZA (norethindrone acetate/ethinyl</p> <p>estradiol/ferrous fumarate)<sup>NR</sup></p> <p>FEMLYV ODT (norethindrone acetate and ethinyl estradiol)<sup>NR</sup></p> <p>MINZOYA (levonorgestrel and ethinyl estradiol tablets, and ferrous bisglycinate)<sup>NR</sup></p> <p>OPILL (norgestrel)<sup>NR</sup> <b>OTC</b></p> <p>VALTYA (ethynodiol diacetate and ethinyl estradiol)<sup>NR</sup></p>		

## CYSTIC FIBROSIS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	<p><b>ALYFTREK (vanzacaftor; tezacaftor; deutivacaftor)<sup>AL,CL,NR</sup> TAB</b></p> <p>BRONCHITOL (mannitol) <sup>AL,CL,QL</sup></p> <p>KALYDECO <b>PACKET, TAB</b> (ivacaftor)<sup>QL, AL</sup></p> <p>ORKAMBI (lumacaftor/ivacaftor) <b>PACKET, TAB</b> <sup>QL, AL</sup></p> <p>SYMDEKO (tezacaftor/ivacaftor)<sup>QL, AL</sup></p> <p>TRIKAFTA(elexacaftor, tezacaftor, ivacaftor)<sup>AL, CL</sup> <b>PACKET<sup>CL</sup>, TAB</b></p>	<p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li>▪ <b>Alyftrek:</b> Diagnosis of CF and documentation of at least one F508del mutation or another responsive mutation in the CFTR gene.</li> <li>▪ <b>Bronchitol:</b> Approved for diagnosis of CF and documentation that the patient has passed the BRONCHITOL Tolerance Test</li> <li>▪ <b>Kalydeco®:</b> Diagnosis of CF and documentation of the drug-specific, FDA-approved mutation of CFTR gene</li> <li>▪ <b>Orkambi®:</b> Diagnosis of CF and documentation of presence of two copies of the F580del mutation (homozygous) of CFTR gene</li> <li>▪ <b>Symdeko:</b> Diagnosis of CF and documentation of the drug specific, FDA approved mutation of CFTR gene.</li> <li>▪ <b>Trikafta:</b> Diagnosis of CF and documentation of at least one F508del mutation in the CFTR gene or a mutation that is responsive to Trikafta based on clinical and/or in vitro data</li> </ul>

## DIURETICS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>SINGLE-AGENT PRODUCTS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of <b>TWO</b> preferred agents within this drug class</li> <li><b>Eplerenone</b>: Will be approved with a failed trial or intolerance to spironolactone, a trial with two preferred agents is not required.</li> <li><b>Kerendia</b>: Approved for diagnosis of chronic kidney disease associated with Type-II diabetes in adults, <del>trial of a preferred agent not required</del></li> <li><b>spironolactone suspension</b>: May be approved without trial of a preferred agent if there is a clinical reason why preferred spironolactone solid dosage form cannot be used.</li> </ul>
amiloride <b>TAB</b>	CAROSPIR (spironolactone) <b>SUSP</b>	
bumetanide <b>TAB</b>	eplerenone <b>TAB</b> (generic Inspra) <sup>CL</sup>	
chlorthalidone <b>TAB</b> (generic Diuril)	ethacrynic acid <b>CAPS</b> (generic Edecrin)	
furosemide <b>SOLN, TAB</b> (generic Lasix)	spironolactone (generic Carospir) <b>SUSP</b>	
hydrochlorothiazide <b>CAPS, TAB</b> (generic Microzide)	THALITONE (chlorthalidone) <b>TAB</b>	
indapamide <b>TAB</b>	triamterene (generic Dyrenium)	
KERENDIA (finerenone) <b>TAB</b> <sup>CL,QL</sup>		
metolazone <b>TAB</b>		
spironolactone <b>TAB</b> (generic Aldactone)		
torsemide <b>TAB</b>		
<b>COMBINATION PRODUCTS</b>		
amiloride/HCTZ <b>TAB</b>		
spironolactone/HCTZ <b>TAB</b> (generic Aldactazide)		
triamterene/HCTZ <b>CAPS, TAB</b>		

## FLUOROQUINOLONES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ciprofloxacin <b>TAB</b> (generic Cipro)	BAXDELA (delafloxacin)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class</li> </ul>
levofloxacin <b>TAB</b> (generic Levaquin)	ciprofloxacin ER	
moxifloxacin (generic Avelox)	ciprofloxacin <b>SUSP</b> (generic Cipro)	
	levofloxacin <b>SOLN</b>	
	ofloxacin	
		Drug-specific criteria:
		<ul style="list-style-type: none"> <li><b>Baxdela</b>: Coverable with documented intolerance or failure of preferred MRSA agents (clindamycin, doxycycline, linezolid, sulfamethoxazole/trimethoprim)</li> <li><b>Ciprofloxacin/Levofloxacin Suspension</b>: Coverable with documented swallowing disorders</li> <li><b>Ofloxacin</b>: Trial of preferred not required for diagnoses of Pelvic Inflammatory Disease OR Acute Epididymitis (non-gonorrhea)</li> </ul>

## GI MOTILITY, CHRONIC

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
LINZESS (linaclotide) <sup>AL, QL</sup> lubiprostone (generic Amitiza) <sup>AL, QL</sup> RELISTOR (methylnaltrexone) <b>SYR</b> TRULANCE (plecanatide) <sup>AL, QL</sup>	alosetron (generic Lotronex) AMITIZA (lubiprostone) <sup>AL, QL</sup> IBSRELA (tenapanor) <sup>AL, QL</sup> MOTEGRITY (prucalopride succinate) MOVANTIK (naloxegol oxalate) <sup>QL</sup> prucalopride (generic Motegrity) <sup>NR</sup> RELISTOR (methylnaltrexone) <sup>QL</sup> <b>TAB, VIAL</b> SYMPROIC (naldemedine) VIBERZI (eluxodoline)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a 30-day trial of ONE preferred agent within this drug class with the same indication</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Ibsrela:</b> May be approved for diagnosis of IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.)</li> <li><b>Lotronex/ alosetron:</b> Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate</li> <li><b>Relistor® TAB:</b> Covered for diagnosis of opioid-induced constipation in adults with chronic, non-cancer pain after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.) and failure of Movantik</li> <li><b>Symproic:</b> Covered for diagnosis of opioid-induced constipation in adult patients with chronic non-cancer pain after trial on at least TWO OTC laxatives and failure of Movantik</li> <li><b>Viberzi®:</b> Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate</li> </ul>

## GLUCAGON AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) <sup>AL, QL</sup> <b>NASAL</b> GLUCAGON EMERGENCY (glucagon) <sup>QL</sup> <b>INJ KIT</b> (Amphastar) PROGLYCEM (diazoxide) <b>SUSP</b> ZEGALOGUE (dasiglucagon) <sup>AL, QL</sup> <b>AUTO-INJ</b>	diazoxide <b>SUSP</b> (generic Proglycem) GLUCAGON EMERGENCY (glucagon) <sup>QL</sup> <b>INJ KIT</b> (Fresenius) GVOKE (glucagon) <sup>AL, QL</sup> <b>KIT, PEN, SYR, VIAL</b> ZEGALOGUE (dasiglucagon) <sup>AL, QL</sup> <b>SYR</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

## GROWTH HORMONES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GENOTROPIN (somatropin)  <b>CARTRIDGE, DISP SYRINGE</b>  NORDITROPIN (somatropin)	HUMATROPE (somatropin)  NGENLA (somatrogon-ghla) <sup>AL</sup>  NUTROPIN AQ (somatropin)  OMNITROPE (somatropin)  SEROSTIM (somatropin)  SKYTROFA (lonapegsomatropin-tcgd)  SOGROYA (somapacitan-beco)  ZOMACTON (somatropin)	<a href="#">Growth Hormone PA Form</a>  <a href="#">Growth Hormone Criteria</a>

## H. PYLORI TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PYLERA (bismuth, metronidazole, tetracycline) <sup>QL</sup>	bismuth,metronidazole,tetracycline (generic Pylera) <sup>QL</sup> lansoprazole/amoxicillin/clarithromycin (generic Prevpac) <sup>QL</sup> OMECLAMOX-PAK (omeprazole, clarithromycin, amoxicillin) <sup>QL</sup> TALICIA (omeprazole/amoxicillin/rifabutin) VOQUEZNA (vonoprazan) <sup>QL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

## HAE TREATMENTS<sup>CL</sup>

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BERINERT (C1 esterase inhibitor, human) <b>INTRAVENOUS</b> HAEGARDA (C1 esterase inhibitor, human) <sup>AL,CL</sup> <b>SUB-Q</b> icatibant acetate (generic for FIRAZYR) <sup>AL</sup> <b>SUB-Q</b> <b>TAKHZYRO (lanadelumab-flyo)<sup>AL,CL</sup></b> <b>SYRINGE</b>	CINRYZE (C1 esterase inhibitor, human) <sup>AL,CL</sup> <b>INTRAVENOUS</b> FIRAZYR (icatibant acetate) <sup>AL</sup> <b>SUB-Q</b> ORLADEYO (berotralstat) <b>CAP<sup>AL,QL</sup></b> RUCONEST (recombinant human C1 inhibitor) <sup>AL</sup> <b>INTRAVENOUS</b> TAKHZYRO (lanadelumab-flyo) <sup>AL,CL</sup> <b>VIAL</b>	<a href="#">HAE Treatments PA Form</a> <ul style="list-style-type: none"> <li>All agents require documentation of diagnosis of Type I or Type II HAE and deficient or dysfunctional C1 esterase inhibitor enzyme. Concomitant use with ACE inhibitors, NSAIDs, or estrogen-containing products is contraindicated</li> <li>Non-preferred agents will be approved for patients who have a failed trial or a contraindication to ONE preferred agent within this drug class with the same indication.</li> </ul> Drug-Specific Criteria <ul style="list-style-type: none"> <li><b>Cinryze, Haegarda, Orladeyo, and Takhzyro</b>, require a history of two or more HAE attacks monthly</li> </ul>

## HEPATITIS B TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
entecavir <b>TAB</b>	adefovir dipivoxil BARACLUDE (entecavir) <b>SOLN,</b> <b>TAB</b> lamivudine hbv <b>TAB</b> VEMLIDY (tenofovir alafenamide fumarate)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul> Drug Specific Criteria <ul style="list-style-type: none"> <li><b>tenofovir disoproxil fumarate (generic Viread) tablet:</b> Diagnosis for use required. May be indicated for chronic hepatitis B or HIV-1 infection.               <ul style="list-style-type: none"> <li>See HIV/AIDS class for drug listing and placement</li> </ul> </li> </ul>

## HEPATITIS C TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>DIRECT ACTING ANTI-VIRAL</b>		<a href="#">Hepatitis C Treatments PA Form</a>  <a href="#">Hepatitis C Criteria</a>
MAVYRET (glecaprevir/pibrentasvir)  <b>TAB<sup>CL</sup>, PELLET<sup>AL,CL</sup></b>  sofosbuvir/velpatasvir (generic Epclusa) <sup>CL</sup>  VOSEVI (sofosbuvir/velpatasvir/voxilaprevir) <sup>CL</sup>	HARVONI 200/45MG, <b>TAB</b> (ledipasvir/sofosbuvir) <sup>CL</sup>  HARVONI (ledipasvir/sofosbuvir) <sup>CL</sup> <b>PELLET</b>  ledipasvir/sofosbuvir (generic Harvoni) <sup>CL</sup>  SOVALDI (sofosbuvir) <sup>CL</sup> <b>PELLET</b>  SOVALDI <b>TAB</b> (sofosbuvir) <sup>CL</sup>  ZEPATIER (elbasvir/grazoprevir) <sup>CL</sup>	<ul style="list-style-type: none"> <li>Non-preferred products require trial of preferred agents within the same group and/or will only be considered with documentation of why the preferred product within this drug class is not appropriate for patient</li> <li>Patients newly eligible for Medicaid will be allowed to complete treatment with the original that treatment was initially authorized by another payor</li> </ul> <p>Drug-specific criteria:</p> <p>Trial with with a preferred agent not required in the following:</p> <ul style="list-style-type: none"> <li><b>Harvoni/ ledipasvir-sofosbuvir:</b> <ul style="list-style-type: none"> <li>Post liver transplant for genotype 1 or 4</li> </ul> </li> <li><b>Vosevi:</b> Requires documentation of non-response after previous treatment course of Direct Acting Anti-viral agent (DAA) for genotype 1-6 without cirrhosis or with compensated cirrhosis</li> </ul>
<b>RIBAVIRIN</b>		
ribavirin 200mg <b>CAP, TAB</b>		
<b>INTERFERON</b>		
PEGASYS (pegylated interferon alfa-2a) <sup>CL</sup>		



## HIV / AIDS<sup>CL</sup>

Preferred Agents		Non-Preferred Agents	Prior Authorization/Class Criteria
CAPSID INHIBITOR			<ul style="list-style-type: none"><li>▪ All agents require:<ul style="list-style-type: none"><li>○ Diagnosis of HIV/AIDS required, OR</li><li>○ Diagnosis of Pre and Post Exposure Prophylaxis</li></ul></li><li>▪ Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li><li>▪ Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li></ul>
	SUNLENCA (lenacapavir) <sup>QL</sup>		
CCR5 ANTAGONISTS			
SELZENTRY <b>SOLN, TAB</b> (maraviroc)	maraviroc (generic Selzentry)		
FUSION INHIBITORS			
FUZEON <b>SUB-Q</b> (enfuvirtide) <sup>QL</sup>			
HIV-1 ATTACHMENT INHIBITOR			
	RUKOBIA ER (fostemsavir) <sup>AL,QL</sup>		
INTEGRASE STRAND TRANSFER INHIBITORS (INSTIs)			
ISENTRESS (raltegravir) <sup>QL</sup>	TIVICAY PD (dolutegravir)		
ISENTRESS HD (raltegravir)			
TIVICAY (dolutegravir)			
NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS (NNRTIs)			
EDURANT (rilpivirine)	etravirine (generic Intelence) <sup>QL</sup>		
efavirenz <b>CAPS, TABLET</b> (generic Sustiva)	nevirapine IR, ER (generic Viramune/Viramune XR)		
INTELENCE (etravirine) <sup>QL</sup>	SUSTIVA CAPS, TABLET (efavirenz)		
PIFELTRO (doravirine) <sup>QL</sup>	VIRAMUNE (nevirapine) <b>SUSP</b>		
NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITORS (NRTIs)			
abacavir <b>SOLN, TABLET</b> (generic Ziagen)	didanosine DR (generic Videx EC)		
EMTRIVA <b>CAPS, SOLN</b> (emtricitabine)	emtricitabine <b>CAPS</b> (generic for Emtriva)		
lamivudine <b>SOLN, TABLET</b> (generic Epivir)	RETROVIR (zidovudine)		
	stavudine <b>CAPS</b> (generic Zerit)		
zidovudine <b>CAPS, SYRUP, TABLET</b> (generic Retrovir)	ZIAGEN (abacavir)		
NUCLEOTIDE REVERSE TRANSCRIPTASE INHIBITORS (NRTIs)			
tenofovir <b>TABLET</b> (generic Viread)	VIREAD (tenofovir) <b>POWDER</b>		
PHARMACOKINETIC ENHANCER			
	TYBOST (cobicistat) <sup>QL</sup>		

## HIV / AIDS <sup>CL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>PROTEASE INHIBITORS</b>		<ul style="list-style-type: none"> <li>All agents require: <ul style="list-style-type: none"> <li>Diagnosis of HIV/AIDS required, OR</li> <li>Diagnosis of Pre and Post Exposure Prophylaxis</li> </ul> </li> <li>Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>
atazanavir <b>CAPS</b> (generic Reyataz) NORVIR (ritonavir) <b>TAB</b> PREZISTA (darunavir) <b>TAB</b> ritonavir TAB (generic Norvir)	APTIVUS <b>CAPS, SOLN</b> (tipranavir) CRIXIVAN (indinavir) DARUNAVIR PROPYLENE GLYCOLATE <sup>AL</sup> <b>TAB</b> darunavir ethanolate (generic Prezista) <sup>AL</sup> <b>TAB</b> fosamprenavir <b>TAB</b> (generic Lexiva) LEXIVA SUSP (fosamprenavir) LEXIVA TAB (fosamprenavir) NORVIR <b>POWDER, SOLN</b> (ritonavir) PREZISTA (darunavir) <b>SUSP</b> REYATAZ <b>POWDER</b> (atazanavir) VIRACEPT (nelfinavir)	
<b>COMBINATION PROTEASE INHIBITORS (PIs) or PIs plus PHARMACOKINETIC ENHANCER</b>		<ul style="list-style-type: none"> <li>All agents require: <ul style="list-style-type: none"> <li>Diagnosis of HIV/AIDS required; OR</li> <li>Diagnosis of Pre and Post Exposure Prophylaxis</li> </ul> </li> <li>Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>
EVOTAZ (atazanavir/cobicistat) <sup>QL</sup> lopinavir/ritonavir <b>SOLN, TAB</b> (generic Kaletra)	KALETRA <b>SOLN</b> (lopinavir/ritonavir) KALETRA <b>TAB</b> (lopinavir/ritonavir) PREZCOBIX (darunavir/cobicistat) <sup>QL</sup>	
<b>COMBINATION NUCLEOS(T)IDE REVERSE TRANSCRIPTASE INHIBITORS</b>		
abacavir/lamivudine (generic Epzicom) CIMDUO (lamivudine/tenofovir) <sup>QL</sup> DESCOVY (emtricitabine/tenofovir) <sup>QL</sup> emtricitabine/tenofovir (generic Truvada) lamivudine/zidovudine (generic Combivir)	abacavir/lamivudine/zidovudine (generic Trizivir) COMBIVIR (lamivudine/zidovudine) EPZICOM (abacavir sulfate/lamivudine) TRIZIVIR (abacavir/lamivudine/zidovudine) TRUVADA (emtricitabine/tenofovir)	

## HIV / AIDS <sup>CL</sup> (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>COMBINATION PRODUCTS – MULTIPLE CLASSES</b>		<ul style="list-style-type: none"> <li>All agents require: <ul style="list-style-type: none"> <li>Diagnosis of HIV/AIDS required, OR</li> <li>Diagnosis of Pre and Post Exposure Prophylaxis</li> </ul> </li> <li>Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific documentation of why the preferred products within this drug class are not appropriate for patient, including, but not limited to, drug resistance or concomitant conditions not recommended with preferred agents</li> <li>Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy</li> </ul>
BIKTARVY (bictegravir/emtricitabine/tenofovir) <sup>QL</sup>	ATRIPLA (efavirenz/emtricitabine/tenofovir)	
COMPLERA (rilpivirine/emtricitabine/tenofovir)	efavirenz/lamivudine/tenofovir (generic for Symfi) <sup>QL</sup>	
DELSTRIGO (dolutegravir/lamivudine/tenofovir) <sup>QL</sup>	efavirenz/lamivudine/tenofovir (generic for Symfi Lo) <sup>QL</sup>	
DOVATO (dolutegravir/lamivudine) <sup>QL</sup>	TRIUMEQ PD (abacavir, dolutegravir, and lamivudine) <b>SUSP</b>	
efavirenz/emtricitabine/tenofovir (generic Atripla) <sup>CL</sup>		
GENVOYA (elvitegravir/cobicistat/emtricitabine/tenofovir) <sup>QL, AL</sup>		
JULUCA (dolutegravir/rilpivirine) <sup>QL</sup>		
ODEFSEY (emtricitabine/rilpivirine/tenofovir) <sup>QL</sup>		
STRIBILD (elvitegravir/cobicistat/emtricitabine/tenofovir) <sup>QL</sup>		
SYMFI (efavirenz/lamivudine/tenofovir) <sup>QL</sup>		
SYMFI LO (efavirenz/lamivudine/tenofovir) <sup>QL</sup>		
SYMTUZA (darunavir/cobicistat/emtricitabine/tenofovir) <sup>QL</sup>		
TRIUMEQ (dolutegravir/abacavir/lamivudine)		

## HYPOGLYCEMICS, ALPHA-GLUCOSIDASE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acarbose (generic for Precose)	miglitol (generic for Glyset)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>

## HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>GLUCAGON-LIKE PEPTIDE-1 RECEPTOR AGONIST (GLP-1 RA)<sup>AL,CL,QL</sup></b>		<b><u>GLP-1 RA Criteria</u></b>
OZEMPIC (semaglutide) <sup>AL,QL</sup>	BYDUREON BCISE <b>PEN</b> (exenatide) <sup>AL,QL</sup>	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin <b>OR</b>  A diagnosis of ASCVD associated with a diagnosis of Type II diabetes  (no metformin trial required)
TRULICITY (dulaglutide) <sup>AL,QL</sup>	BYETTA (exenatide) <sup>AL,QL</sup> subcutaneous	
VICTOZA (liraglutide) <sup>AL,QL</sup> subcutaneous	<b>liraglutide (generic Victoza)<sup>AL,NR,QL</sup></b>	
	MOUNJARO (tirzepatide) <sup>AL,QL</sup> <b>PEN</b>	
	RYBELSUS (semaglutide) <sup>AL,QL</sup> <b>TAB</b>	
<b>INSULIN/GLP-1 RA COMBINATIONS</b>		Non-preferred agents will be approved for patients who have:
	SOLIQUA (insulin glargine/lixisenatide)	
	XULTOPHY (insulin degludec/liraglutide)	<ul style="list-style-type: none"> <li>Failed a trial of TWO preferred agents within GLP-1 RA</li> </ul> AND
<b>AMYLIN ANALOG</b>		<b><u>Amylin Analog Criteria</u></b>
	SYMLIN (pramlintide) subcutaneous	ALL criteria must be met
<b>DIPEPTIDYL PEPTIDASE-4 (DPP-4) INHIBITOR<sup>AL,QL</sup></b>		<ul style="list-style-type: none"> <li>Concurrent use of short-acting mealtime insulin</li> <li>Current therapy compliance</li> <li>No diagnosis of gastroparesis</li> <li>HbA1C ≤ 9% within last 90 days</li> <li>Monitoring of glucose during initiation of therapy</li> </ul>
JANUMET (sitagliptin/metformin)	alogliptin (generic Nesina)	<b><u>DPP-4 Inhibitor Criteria</u></b>
JANUMET XR (sitagliptin/metformin)	alogliptin/metformin (generic Kazano)	
JANUVIA (sitagliptin)	alogliptin/pioglitazone (generic Oseni)	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin.
JENTADUETO (linagliptin/metformin)	GLYXAMBI (empagliflozin/linagliptin)	
TRADJENTA (linagliptin)	JENTADUETO XR (linagliptin/metformin)	Non-preferred DPP-4s will be approved for patients who have failed a trial of ONE preferred agent within the DPP-4 inhibitor class
	KOMBIGLYZE XR (saxagliptin/metformin)	
	ONGLYZA (saxagliptin)	
	QTERN (dapagliflozin/saxagliptin)	
	saxagliptin (generic Onglyza)	
	saxagliptin/metformin ER (generic Kombiglyze ER)	
	<b>sitagliptin (generic Zituvio)<sup>NR</sup></b>	
	<b>sitagliptin/ metformin (Zituvimet)<sup>NR</sup></b>	
	STEGLUJAN (ertugliflozin/sitagliptin)	
	TRIJARDY XR (empagliflozin/linagliptin/metformin)	
	ZITUVIMET (sitagliptin and metformin) TABLET <sup>NR, QL</sup>	

HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIPEPTIDYL PEPTIDASE-4 (DPP-4) INHIBITOR <sup>AL,QL</sup>		
	ZITUVIMET XR (sitagliptin and metformin) TABLET <sup>NR, QL</sup>	
	ZITUVIO (sitagliptin)	

## HYPOGLYCEMICS, INSULIN AND RELATED DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
HUMULIN (insulin) <b>VIAL</b>	ADMELOG (insulin lispro) <b>PEN, VIAL</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>
HUMULIN 70/30 <b>VIAL</b>	AFREZZA (regular insulin) <b>INHALATION</b>	
HUMULIN U-500 <b>VIAL</b>	APIDRA (insulin glulisine) <b>SOLOSTAR, VIAL</b>	<p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Afrezza®</b>: Approved for T1DM on long-acting insulin with no current history of smoking or chronic lung disease</li> <li><b>Humulin® R U-500 Kwikpen</b>: May be approved for patients who require &gt;200 units/day</li> <li><b>Humalog U-200 Pen</b>: May be approved for patients who require &gt; 100 units/day</li> </ul>
HUMULIN 500 U/M <b>PEN<sup>CL</sup></b>	BASAGLAR (insulin glargine, rec) <b>PEN, TEMPO PEN</b>	
HUMULIN OTC <b>PEN</b>	FIASP (insulin aspart) <b>CARTRIDGE, PEN, VIAL</b>	
HUMULIN 70/30 OTC <b>PEN</b>	HUMALOG U-100 <b>TEMPO PEN</b>	
insulin aspart (generic for Novolog) <b>CARTRIDGE, PEN, VIAL</b>	HUMALOG (insulin lispro) <sup>CL</sup> U-200 <b>KWIKPEN</b>	
insulin aspart/insulin aspart protamine <b>PEN, VIAL</b> (generic for Novolog Mix)	HUMALOG (insulin lispro) U-100 <b>CARTRIDGE, PEN, VIAL</b>	
insulin lispro (generic for Humalog) <b>PEN, VIAL, JR KWIKPEN</b>	HUMALOG JR. (insulin lispro) U-100 <b>KWIKPEN</b>	
insulin lispro/lispro protamine <b>KWIKPEN</b> (Humalog Mix Kwikpen)	HUMALOG MIX <b>VIAL</b> (insulin lispro/lispro protamine)	
LANTUS SOLOSTAR <b>PEN</b> (insulin glargine)	HUMALOG MIX <b>KWIKPEN</b> (insulin lispro/lispro protamine)	
LANTUS (insulin glargine) <b>VIAL</b>	insulin degludec (generic Tresiba) 100U/mL <b>PEN, VIAL</b>	
	insulin degludec (generic Tresiba) 200U/mL <b>PEN</b>	
	insulin glargine <b>PEN, VIAL</b>	
	insulin glargine (Toujeo)	
	insulin glargine max (Toujeo Max)	
	insulin glargine-YFGN <b>PEN, VIAL</b> (generic for Semglee-YFGN)	
	LEVEMIR (insulin detemir) <b>PEN, VIAL</b>	
	LYUMJEV <b>KWIKPEN, VIAL</b> (insulin lispro-aabc)	
	LYUMJEV (insulin lispro-aabc) <b>TEMPO PEN</b>	

## HYPOGLYCEMICS, INSULIN AND RELATED DRUGS, continued

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	NOVOLIN (insulin) <b>NOVOLIN (insulin) PEN-OTC</b> NOVOLIN 70/30 (insulin) <b>PEN-OTC VIAL-OTC</b> <b>NOVOLOG (insulin aspart) CARTRIDGE, FLEXPEN, VIAL</b> NOVOLOG MIX (insulin aspart/aspart protamine) <b>FLEXPEN, VIAL</b> REZVOGLAR (insulin glargine-aglr) <b>KWIKPEN</b> SEMGLEE (insulin glargine) <b>PEN, VIAL</b> SEMGLEE YFGN (insulin glargine) <b>PEN, VIAL</b> TOUJEO SOLOSTAR (insulin glargine) TRESIBA (insulin degludec)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Afrezza®</b>: Approved for T1DM on long-acting insulin with no current history of smoking or chronic lung disease</li> <li><b>Humulin® R U-500 Kwikpen</b>: May be approved for patients who require &gt;200 units/day</li> <li><b>Humalog U-200 Pen</b>: May be approved for patients who require &gt; 100 units/day</li> </ul>

## HYPOGLYCEMICS, MEGLITINIDES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
repaglinide (generic for Prandin)	nateglinide (generic for Starlix) <sup>CL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients with: Failure of a trial of ONE preferred agent in another Hypoglycemic class OR  T2DM and inadequate glycemic control</li> </ul>

## HYPOGLYCEMICS, METFORMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
metformin IR & ER (generic Glucophage/Glucophage XR)	<p>metformin<sup>NR</sup> 750 mg</p> <p>metformin ER (generic Fortamet/Glumetza)</p> <p>metformin <b>SOLN</b> (generic Riomet)</p> <p>RIOMET ER (metformin ER)<sup>AL</sup></p>	<ul style="list-style-type: none"> <li><b>Metformin ER (generic Fortamet®)/Glumetza®:</b> Requires clinical reason why generic Glucophage XR® cannot be used</li> <li><b>Metformin solution:</b> Prior authorization not required for age &lt;7 years</li> </ul>



## HYPOGLYCEMICS, SGLT2

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) <sup>CL,QL</sup>	BRENZAVVY (bexagliflozin) <sup>NR</sup>	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin, <b>OR</b>
JARDIANCE (empagliflozin) <sup>CL,QL</sup>	dapagliflozin <sup>CL,NR,QL</sup> (generic Farxiga)	
SYNJARDY (empagliflozin/metformin) <sup>AL,CL,QL</sup>	dapagliflozin/metformin <sup>CL,QL</sup> (generic Xigduo)	A diagnosis of ASCVD or Heart Failure, or chronic kidney disease associated with a diagnosis of Type II diabetes (no metformin trial required)
XIGDUO XR (dapagliflozin/metformin) <sup>CL,QL</sup>	INPEFA (sotagliflozin) <sup>QL</sup> <b>TAB</b>	
	INVOKAMET (canagliflozin/ metformin) <sup>CL,QL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class</li> </ul>
	INVOKAMET XR (canagliflozin/metformin) <sup>QL</sup>	
	INVOKANA (canagliflozin) <sup>CL</sup>	<p>Drug Specific Criteria:</p> <ul style="list-style-type: none"> <li>- <b>Farxiga/ dapagliflozin:</b> May be approved for a diagnosis of Heart Failure without a diagnosis of diabetes</li> <li>- May be approved for a diagnosis of chronic kidney disease at risk of progression without a diagnosis of diabetes</li> <li>• <b>Jardiance:</b> May be approved for a diagnosis of Heart Failure without a diagnosis of diabetes</li> </ul>
	SEGLUROMET (ertugliflozin/metformin) <sup>QL</sup>	
	STEGLATRO (ertugliflozin) <sup>QL</sup>	
	SYNJARDY XR (empagliflozin/ metformin) <sup>AL,QL</sup>	

## HYPOGLYCEMICS, SULFONYLUREAS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
glimepiride (generic Amaryl) glipizide IR & ER (generic Glucotrol/ Glucotrol XL) glyburide (generic Diabeta/Glynase)		<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li></ul>
<b>SULFONYLUREA COMBINATIONS</b>		
glipizide/metformin glyburide/metformin (generic Glucovance)		

## HYPOGLYCEMICS, TZD

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>THIAZOLIDINEDIONES (TZDs)</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of THE preferred agent within this drug class</li> </ul>
pioglitazone (generic for Actos)		
<b>TZD COMBINATIONS</b>		<ul style="list-style-type: none"> <li><b>Combination products:</b> Require clinical reason why individual ingredients cannot be used</li> </ul>
	pioglitazone/glimepiride (generic for Duetact)	
	pioglitazone/metformin (generic for Actoplus Met)	

## IMMUNOSUPPRESSIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathioprine (generic Imuran)	ASTAGRAF XL (tacrolimus)	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class
azathioprine (generic Azasan) <sup>NR</sup>	AZASAN (azathioprine)	
cyclosporine, modified (generic Neoral) <b>CAPS</b>	cyclosporine <b>CAP, SOFTGEL</b>	<ul style="list-style-type: none"> <li>Patients established on existing therapy will be allowed to continue</li> </ul>
everolimus (generic for Zortress) <sup>AL</sup>	cyclosporine, modified (generic Neoral) <b>SOLN</b>	
mycophenolate mofetil (generic Cellcept) <b>CAPS, TAB</b>	ENVARUSUS XR (tacrolimus)	Drug Specific Criteria
mycophenolic acid (generic Myfortic)	GENGRAF (cyclosporine, modified)	
RAPAMUNE (sirolimus) <b>TAB</b>	<b>CAP, SOLN</b>	<ul style="list-style-type: none"> <li><b>Tavneos</b> (avacopan) <ul style="list-style-type: none"> <li>No trial of a preferred agent required with appropriate FDA indications with concurrent use of standard therapy, including glucocorticoids</li> </ul> </li> </ul>
sirolimus (generic Rapamune)	mycophenolate mofetil (generic Cellcept) <b>SUSP</b>	
<b>SOLN, TAB</b>	MYFORTIC (mycophenolate sodium)	
tacrolimus	<b>MYHIBBIN (mycophenolate)<sup>AL,NR</sup> SUSP</b>	
	PROGRAF (tacrolimus) <b>CAPS, PACKET</b>	
	REZUROCK (belumosudil) <sup>AL,QL</sup> <b>TAB</b>	
	SANDIMMUNE (cyclosporine)	
	<b>CAPS, SOLN</b>	
	TAVNEOS (avacopan) <sup>QL</sup> <b>CAPS</b>	
	ZORTRESS (everolimus) <sup>AL</sup>	

## LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clindamycin <b>CAPS</b>	CLEOCIN (clindamycin) <b>CAPS</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>
clindamycin palmitate <b>SOLN</b>	CLEOCIN PALMITATE (clindamycin)	
linezolid <b>TAB</b>	linezolid <b>SUSP</b>	
	SIVEXTRO (tedizolid phosphate)	
	ZYVOX (linezolid) <b>SUSP, TAB</b>	

## LIPOTROPICS, OTHER

Preferred Agents		Non-Preferred Agents	Prior Authorization/Class Criteria
BILE ACID SEQUESTRANTS			<ul style="list-style-type: none"><li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li></ul> Drug-specific criteria: <ul style="list-style-type: none"><li><b>Colesevelam:</b> Trial not required for diabetes control and monotherapy with metformin, sulfonylurea, or insulin has been inadequate</li><li><b>Juxtapid/ Kynamro:</b><ul style="list-style-type: none"><li>Approved for diagnosis of homozygous familial hypercholesterolemia (HoFH) OR</li><li>Treatment failure/maximized dosing/contraindication to ALL the following: statins, ezetimibe, niacin, fibric acid derivatives, omega-3 agents, bile acid sequestrants</li><li>Require faxed copy of REMS PA form</li></ul></li></ul>
cholestyramine (generic Questran)	colesevelam (generic Welchol) <b>TAB, PACKET</b>		
colestipol <b>TAB</b> (generic Colestid)	colestipol <b>GRANULES</b> (generic Colestid)		
	QUESTRAN LIGHT (cholestyramine)		
TREATMENT OF HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA			
	JUXTAPID (lomitapide) <sup>CL</sup>		
	KYNAMRO (mipomersen) <sup>CL</sup>		
TREATMENT OF FAMILIAL CHYLOMICRONEMIA SYNDROME (FCS)			
	TRYNGOLZA (olezarsen) <sup>AL,CL,QL</sup> <b>INJ</b>		
FIBRIC ACID DERIVATIVES			
fenofibrate (generic Tricor)	fenofibric acid (generic Fibracor/Trilipix)		
fenofibrate (generic Lofibra)	fenofibrate (generic Antara/Fenoglidle/Lipofen/Triglidle)		
gemfibrozil (generic Lopid)			
NIACIN			
niacin ER (generic Niaspan)	NIACOR (niacin IR)		
OMEGA-3 FATTY ACIDS			
omega-3 fatty acids (generic Lovaza)	icosapent (generic Vascepa) <sup>CL</sup>		
	omega-3 OTC		
CHOLESTEROL ABSORPTION INHIBITORS			
ezetimibe (generic Zetia)	NEXLETOL (bempedoic acid)		
	NEXLIZET (bempedoic acid/ezetimibe) <sup>QL</sup>		

## LIPOTROPICS, OTHER (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>PROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 9 (PCSK9) INHIBITORS</b>		<b>Drug-Specific Criteria</b>
PRALUENT (alirocumab) <sup>CL</sup>  REPATHA (evolocumab) <sup>CL</sup>  <b>SURECLICK, SYR</b>		<b>Praluent and Repatha:</b> May be approved for diagnoses of: <ul style="list-style-type: none"> <li>Atherosclerotic cardiovascular disease (ASCVD) in adults</li> <li>Heterozygous familial hypercholesterolemia (HeFH)               <ul style="list-style-type: none"> <li>Praluent ≥ 8 years of age</li> <li>Repatha ≥ 10 years of age</li> </ul> </li> <li>Homozygous familial hypercholesterolemia (HoFH)               <ul style="list-style-type: none"> <li>Praluent ≥ 18 years of age</li> <li>Repatha ≥ 10 years of age</li> </ul> </li> </ul> <b>AND</b> <ul style="list-style-type: none"> <li>Trial and failure or intolerance to a statin for 8 continuous weeks</li> <li>Concurrent use of a maximally tolerated statin must continue, except for statin-induced rhabdomyolysis or a contraindication to a statin</li> <li>Failure to reach target LDL-C levels:               <ul style="list-style-type: none"> <li>ASCVD – &lt; 70 mg/dL</li> <li>HeFH – &lt; 100 mg/dL</li> </ul> </li> </ul>

## LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>STATINS</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agent within this drug class, within the last 12 months</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Altoprev®:</b> One of the TWO trials must be IR lovastatin</li> <li><b>Combination products:</b> Require clinical reason why individual ingredients cannot be used</li> <li><b>fluvastatin ER:</b> Requires trial of TWO preferred agents AND trial of IR fluvastatin OR clinical reason IR cannot be used</li> <li><b>simvastatin/ezetimibe:</b> Approved for 3-month continuous trial of ONE standard dose statin</li> </ul>
atorvastatin (generic Lipitor) <sup>QL</sup>	ALTOPREV (lovastatin ER) <sup>CL</sup>	
lovastatin (generic Mevacor)	ATORVALIQ (atorvastatin) <sup>QL</sup> <b>SUSP</b>	
pravastatin (generic Pravachol)	EZALLOR SPRINKLE (rosuvastatin) <sup>QL</sup>	
rosuvastatin (generic Crestor)	fluvastatin IR/ER (generic Lescol/Lescol XL)	
simvastatin (generic Zocor)	LIVALO (pitavastatin) <sup>AL,QL</sup> pitavastatin (generic Livalo) <sup>AL,NR,QL</sup> ZYPITAMAG (pitavastatin)	
<b>STATIN COMBINATIONS</b>		
	atorvastatin/amlodipine (generic Caduet)  simvastatin/ezetimibe (generic Vytorin)	

## MACROLIDES AND KETOLIDES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>MACROLIDES</b>		<ul style="list-style-type: none"> <li>Non-preferred agents require clinical reason why preferred products within this drug class cannot be used AND ≥ 3-day trial on a preferred product</li> </ul>
azithromycin (generic Zithromax)	clarithromycin ER (generic Biaxin XL)	
clarithromycin <b>TAB, SUSP</b> (generic Biaxin)	E.E.S. <b>TAB</b> (erythromycin ethylsuccinate)	
E.E.S. <b>SUSP</b> (erythromycin ethylsuccinate)	ERY-TAB (erythromycin)	
	erythromycin ethylsuccinate <b>SUSP</b>	
	ERYPED <b>SUSP</b> (erythromycin)	
	ERYTHROCIN (erythromycin)	
	erythromycin base <b>TAB, CAPS</b>	

## MULTIPLE SCLEROSIS DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AVONEX (interferon beta-1a) <sup>QL</sup> COPAXONE 20mg (glatiramer) <sup>QL</sup> dimethyl fumarate (generic Tecfidera) fingolimod (generic Gilenya) <sup>QL</sup> KESIMPTA (Ofatumumab) <sup>CL,QL</sup> teriflunomide (generic Aubagio) <sup>QL</sup>	AUBAGIO (teriflunomide) <sup>QL</sup> BAFIERTAM (monomethyl fumarate) <sup>QL</sup> <b>BETASERON (interferon beta-1b)<sup>QL</sup></b> dalfampridine (generic Ampyra) <sup>QL</sup> <b>dimethyl fumarate (generic Tecfidera) STARTER PACK</b> EXTAVIA (interferon beta-1b) <sup>QL</sup> GILENYA (fingolimod) <sup>QL</sup> glatiramer (generic Copaxone) <sup>QL</sup> MAVENCLAD (cladribine) MAYZENT (siponimod) <sup>QL</sup> PLEGRIDY (peginterferon beta-1a) <sup>QL</sup> PONVORY (ponesimod) REBIF (interferon beta-1a) <sup>QL</sup> TASCENSO ODT (fingolimod) <b>TAB</b> <sup>AL</sup> TECFIDERA (dimethyl fumarate) VUMERITY (diroximel) <sup>QL</sup> ZEPOSIA (ozanimod) <sup>AL,CL,QL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agent within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Ampyra/ dalfampridine:</b> Approved for diagnosis of gait disorder associated with MS AND EDSS score ≤ 7</li> <li><b>Plegridy:</b> Approved for diagnosis of relapsing MS</li> <li><b>Kesimpta:</b> Approved for patients who have failed a trial of a preferred injectable agent within this class</li> <li><b>Zeposia:</b> Approved for a diagnosis of relapsing forms of multiple sclerosis (MS) with trial of ONE preferred agent OR a diagnosis of moderately to severely active ulcerative colitis and treatment failure of Humira.</li> </ul>

## NITROFURAN DERIVATIVES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
nitrofurantoin macrocrystals <b>CAPSULE</b> (generic Macrochantin) nitrofurantoin monohydrate-macrocrystals <b>CAPS</b> (generic Macrobid)	nitrofurantoin <b>SUSPENSION</b> (generic Furadantin)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of <b>ONE</b> preferred agent within this drug class</li> </ul>

## OPIOID DEPENDENCE TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
buprenorphine <b>SL</b> buprenorphine/naloxone <b>TAB (SL)</b> SUBOXONE <b>FILM</b> (buprenorphine/naloxone)	buprenorphine/naloxone <b>FILM</b> lofexidine (generic Lucemyra) <sup>CL,NR,QL</sup> LUCEMYRA (lofexidine) <sup>CL,QL</sup> ZUBSOLV (buprenorphine/naloxone)	<a href="#">Opioid Dependence Treatment PA Form</a> <a href="#">Opioid Dependence Treatment Informed Consent</a> <ul style="list-style-type: none"> <li>Non-preferred agents require a treatment failure of a preferred drug or patient-specific documentation of why a preferred product is not appropriate for the patient.</li> </ul> Drug-specific criteria: <ul style="list-style-type: none"> <li><b>Lucemyra/ lofexidine:</b> Approved for FDA approved indication and dosing per label. Trial of preferred product not required.</li> </ul>

## OPIOID-REVERSAL TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naltrexone <b>TAB</b> naloxone <b>NASAL(Rx), VIAL</b> NARCAN (naloxone) <b>NASAL (OTC)</b>	KLOXXADO (naloxone) <b>NASAL</b> naloxone (generic Narcan) <b>OTC NASAL</b> naloxone (generic Narcan) <b>(Rx) SYR</b> NARCAN (naloxone) <b>NASAL (Rx)</b> OPVEE (nalmeffene) <sup>AL</sup> <b>NASAL</b> REXTOVY (naloxone) <sup>NR</sup> <b>NASAL</b> ZIMHI (naloxone) <b>SYR</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved with documentation of why preferred products within this drug class are not appropriate for the patient</li> </ul>



## PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ambrisentan (generic Letairis) REVATIO (sildenafil) <sup>QL</sup> <b>TAB</b> sildenafil (generic Revatio) <sup>CL</sup> <b>SUSP</b> tadalafil (generic for Adcirca) <sup>CL</sup> TRACLEER (bosentan) <b>TAB</b>	ADEMPAS (riociguat) <sup>CL</sup> ADCIRCA (tadalafil) <sup>CL</sup> bosentan (generic Tracleer) <b>TAB</b> LETAIRIS (ambrisentan) LIQREV (sildenafil) <b>SUSP</b> OPSUMIT (macitentan) OPSYNVI (macitentan and tadalafil) <sup>NR</sup> <b>TAB</b> ORENITRAM ER (treprostinil) REVATIO (sildenafil) <sup>CL</sup> <b>SUSP</b> sildenafil (generic Revatio) <sup>CL</sup> <b>TAB</b> TADLIQ (tadalafil) <b>SUSP</b> TRACLEER (bosentan) <b>TAB FOR SUSPENSION</b> TYVASO (treprostinil) <b>INHALATION</b> TYVASO DPI (treprostinil) <b>INHALATION POWDER</b> UPTRAVI (selexipag) VENTAVIS (iloprost) <b>INHALATION</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Adcirca/Liqrev/Revatio/sildenafil tablets and suspension/tadalafil:</b> Approved for diagnosis of Pulmonary Arterial Hypertension (PAH)</li> <li><b>Adempas®:</b> PAH: Requires clinical reason preferred agent cannot be used  CTEPH: Approved for persistent/recurrent diagnosis after surgical treatment or inoperable CTEPH  NOT for use in Pregnancy</li> <li><b>Liqrev/ Revatio suspension:</b> Requires clinical reason why preferred sildenafil suspension cannot be used</li> </ul>

## PANCREATIC ENZYMES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CREON ZENPEP (pancrelipase)	PERTZYE (pancrelipase) VIOKACE (pancrelipase)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul>

## PEDIATRIC VITAMIN PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD MVI (mvi, ped mvi no. 19/FA, ped mvi no. 17) <b>OTC CHEW</b>	DEKAs PLUS <sup>AL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul> <p>Drug specific criteria:</p> <ul style="list-style-type: none"> <li><b>DEKAs Plus:</b> Approved for diagnosis of Cystic Fibrosis and does not require a trial of a preferred agent</li> </ul>
CHILDREN'S MVI-IRON <b>OTC CHEW</b> (ped mvi no. 91/iron fum)	DAVIMET W/ FLUORIDE (ped mvi no.247/ fluoride) <sup>NR</sup> <b>CHEW OTC</b>	
CHILDREN'S CHEWABLES <b>OTC</b> (ped mvi no. 25/FA, ped mvi no. 31 /iron/FA, ped mvi no.17/iron sulf)	FLORAFOL(mvi and fluoride) <sup>NR</sup> <b>CHEW OTC, DROPS-OTC<sup>NR</sup></b>	
FLUORIDE/VITAMINS A,C,AND D <b>DROPS</b> (ped mvi A,C,D3 no.21/ fluoride)	FLORAFOL FE PEDIATRIC <sup>NR</sup> <b>DROPS OTC</b>	
MULTIVITAMINS W/ FLUORIDE (PEDI MVI NO.2 W-FLUORIDE) <b>DROPS</b>	MULTI-VIT-FLOR (ped mvi no.205/fluoride) <b>CHEW</b>	
MULTIVITS W/ IRON & FLUORIDE <b>DROPS</b> (ped mvi no. 45/fluoride/iron)	PEDI MULTIVIT A,C,AND D3 NO.21 <b>DROPS<sup>NR</sup> OTC</b>	
PED MVI NO.17 W/ FLUORIDE <b>CHEW</b>	PEDI MVI NO.242/FLUORIDE <b>CHEW OTC</b>	
POLY-VITAMIN (ped mvi no. 212) <b>DROPS OTC</b>	POLY-VI-FLOR (ped mvi no.217/fluoride, ped mvi no. 205/fluoride) <b>CHEW</b>	
POLY-VITAMIN W/ IRON (ped mvi no. 207 w/ferrous sulf) <b>DROPS OTC</b>	POLY-VI-FLOR (ped mvi no.213 w/fluoride) <b>DROPS</b>	
TRI-VITAMIN W/ FLUORIDE (ped mvi A,C, D3 no. 21/fluoride)	POLY-VI-FLOR W/ IRON (ped mvi no. 205/fluoride/iron) <b>CHEW</b>	
	POLY-VI-FLOR W/ IRON (ped mvi no. 214/fluoride/iron) <b>DROPS</b>	

## PEDIATRIC VITAMIN PREPARATIONS, continued

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	<p>QUFLORA (ped mvi no.84/fluoride, ped mvi no. 63/fluoride, ped mvi no. 83/fluoride)</p> <p>QUFLORA FE (ped mvi 142/iron/fluoride, ped mvi no. 151/iron/fluoride) <b>CHEW</b></p> <p>QUFLORA (ped mvi no.157/ fluoride) <b>OTC</b></p> <p><b>SOLUVITA A,C,D WITH FLUORIDE DROPS<sup>NR</sup> OTC</b></p>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class</li> </ul> <p>Drug specific criteria:</p> <ul style="list-style-type: none"> <li><b>DEKAs Plus:</b> Approved for diagnosis of Cystic Fibrosis and does not require a trial of a preferred agent</li> </ul>

## PENICILLINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
amoxicillin <b>CAPS, CHEWABLE TAB, SUSP, TAB</b> ampicillin <b>CAPS</b> dicloxacillin penicillin VK		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class</li> </ul>

## PHOSPHATE BINDERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcium acetate <b>TAB</b> sevelamer carbonate (generic Renvela) <b>PWD PACK, TAB</b>	AURYXIA (ferric citrate) calcium acetate <b>CAPS</b> <b>CALPHRON OTC (calcium acetate)</b> lanthanum (generic FOSRENOL) PHOSLYRA (calcium acetate) RENAGEL (sevelamer HCl) <b>TAB</b> RENVELA (sevelamer carbonate) <b>PWD PACK, TAB</b> sevelamer HCl (generic Renagel) VELPHORO (sucroferric oxyhydroxide) XPHOZAH (tenapanor) <b>TAB</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class within the last 6 months</li> </ul>

## PLATELET AGGREGATION INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
aspirin BRILINTA (ticagrelor) clopidogrel (generic Plavix) dipyridamole (generic Persantine) prasugrel (generic Effient)	aspirin/dipyridamole (generic Aggrenox) ticlopidine (generic Ticlid) YOSPRALA (aspirin/omeprazole)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class OR documented clopidogrel resistance</li> </ul>

Additional covered agents can be looked up using the Drug Look-up Tool at:

<https://ne.primetherapeutics.com/drug-lookup>

## PRENATAL VITAMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FE C/FA	CITRANATAL B-CALM	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of or are intolerant to TWO preferred agents within this drug class</li> </ul>
PNV 2/IRON B-G SUC-P/FA/OMEGA-3	COMPLETENATE CHEW TABLET	
PNV NO.118/IRON FUMARATE/FA <b>CHEW TAB</b>	DERMACINRX PRENATRIX <b>OTC</b>	
PNV NO.15/IRON FUM & PS CMP/FA	DERMACINRX PRETRATE <b>TAB</b>	
PNV WITH CA, NO.72/IRON/FA	ENBRACE HR	
PNV WITH CA, NO.74/IRON/FA <b>OTC</b>	MARNATAL-F	
PNV#16/IRON FUM & PS/FA/OM-3	MULTI-MAC <b>OTC</b>	
PNV119/IRON FUMARATE/FA/DSS	NATAL PNV (pnv no.164/iron/folate no.6)	
PRENATAL MULTI <b>OTC</b>	NEO-VITAL RX TAB <b>OTC<sup>NR</sup></b>	
PRENATAL VIT #76/IRON, CARB/FA	NESTABS	
PRENATAL VIT/FE FUMARATE/FA <b>OTC</b>	NESTABS ABC	
SELECT-OB + DHA	NESTABS DHA	
STUART ONE <b>OTC</b>	NESTABS ONE	
TRICARE	OB COMPLETE ONE	
TRINATAL RX 1	OB COMPLETE PETITE	
VITAFOL <b>CHEW TAB</b>	OB COMPLETE PREMIER	
VITAFOL FE+	OB COMPLETE <b>TAB</b>	
VITAFOL ULTRA	OB COMPLETE WITH DHA <b>OTC</b>	
VITAFOL-OB	PNV 11-IRON FUM-FOLIC ACID-OM3	
VITAFOL-OB+DHA	PNV COMBO#47/IRON/FA #1/DHA	
VITAFOL-ONE	PNV W-CA NO.40/IRON FUM/FA CMB NO.1	
	PNV WITH CA NO.68/IRON/FA NO.1/DHA	
	PRENATAL + DHA <b>OTC</b>	
	PRENATE AM	
	PRENATE <b>CHEW TAB</b>	
	PRENATE DHA	
	PRENATE ELITE	
	PRENATE ENHANCE	
	PRENATE ESSENTIAL	
	PRENATE MINI	
	PRENATE PIXIE	
	PRENATE RESTORE	
	PRENATE STAR	
	PRIMACARE	
	SELECT-OB <b>CHEW TAB</b>	
	TRISTART DHA	
	VITAFOL NANO	
	WESTGEL DHA	

## PROTON PUMP INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
esomeprazole magnesium (generic Nexium) <b>RX</b> <sup>QL</sup>	DEXILANT (dexlansoprazole)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed an 8-week trial of THREE preferred agents.</li> </ul> <p><b>Pediatric Patients:</b></p> <p>Patients <math>\leq 4</math> years of age – No PA required for Prevacid 30mg/ lansoprazole 30mg capsules (used to compound suspensions).</p> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Prilosec®OTC/Omeprazole OTC:</b> EXCLUDED from coverage Acceptable as trial instead of Omeprazole 20mg</li> <li><b>Prevacid (lansoprazole) Solutab:</b> may be approved after trial of compounded suspension. Patients <math>\geq 5</math> years of age- Only approve non-preferred for GI diagnosis if: <ul style="list-style-type: none"> <li>Child can not swallow whole generic omeprazole capsules OR,</li> <li>Documentation that contents of capsule may not be sprinkled in applesauce</li> </ul> </li> </ul>
omeprazole (generic Prilosec) <b>RX</b>	dexlansoprazole (generic Dexilant)	
pantoprazole (generic Protonix) <sup>QL</sup>	esomeprazole magnesium (generic Nexium) <b>OTC</b> <sup>QL</sup>	
PROTONIX <b>SUSP</b> (pantoprazole)	esomeprazole strontium	
	KONVOMEF (omeprazole/sodium bicarb) <b>SUSP</b>	
	lansoprazole (generic Prevacid) <sup>QL</sup>	
	NEXIUM <b>SUSP</b> (esomeprazole)	
	omeprazole/sodium bicarbonate (generic Zegerid RX)	
	pantoprazole <b>GRANULES</b> <sup>QL</sup>	
	rabeprazole (generic Aciphex)	

## SINUS NODE INHIBITORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORLANOR <b>SOLN, TAB</b> (ivabradine) ivabradine (generic Corlanor) <sup>NR</sup> <b>TAB</b>	<ul style="list-style-type: none"> <li>▪ Diagnosis of Chronic Heart Failure with left ventricular ejection fraction less than or equal to 35%, AND</li> <li>▪ Sinus rhythm with resting heart rate greater than or equal to 70 beats per minute, AND</li> <li>▪ On maximally tolerated doses of beta-blockers OR have a contraindication to beta-blocker use</li> </ul>

## SKELETAL MUSCLE RELAXANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
baclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) <sup>QL</sup> methocarbamol (generic Robaxin) tizanidine <b>TAB</b> (generic Zanaflex)	baclofen (generic Fleqsuvy) <sup>QL</sup> <b>SUSP</b> baclofen (generic Ozobax) <sup>QL</sup> <b>SOLN</b> baclofen (generic Ozobax DS)  <b>SUSP</b> carisoprodol (generic Soma) <sup>CL,QL</sup> carisoprodol compound cyclobenzaprine ER (generic Amrix) <sup>CL</sup> dantrolene (generic Dantrium) FEXMID (cyclobenzaprine ER) FLEQSUVY (baclofen) <sup>QL</sup> <b>SUSP</b> LORZONE (chlorzoxazone) <sup>CL</sup> LYVISPAH (baclofen) <sup>QL</sup> <b>GRANULES</b> metaxalone (generic Skelaxin) NORGESIC FORTE (orphenadrine/ASA/caffeine) orphenadrine ER PARAFON FORTE (chlorzoxazone) <b>TANLOR (methocarbamol)<sup>NR</sup> TAB</b> tizanidine <b>CAPS</b> ZANAFLEX (tizanidine) <b>CAPS, TAB</b>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a 1-week trial of TWO preferred agents within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>cyclobenzaprine ER:</b> <ul style="list-style-type: none"> <li>Requires clinical reason why IR cannot be used</li> <li>Approved only for acute muscle spasms</li> <li>NOT approved for chronic use</li> </ul> </li> <li><b>carisoprodol:</b> <ul style="list-style-type: none"> <li>Approved for Acute, musculoskeletal pain - NOT for chronic pain</li> <li>Use is limited to no more than 30 days</li> <li>Additional authorizations will not be granted for at least 6 months following the last day of previous course of therapy</li> </ul> </li> <li><b>Dantrolene:</b> Trial NOT required for treatment of spasticity from spinal cord injury</li> <li><b>Lorzone®:</b> Requires clinical reason why chlorzoxazone cannot be used</li> <li><b>Soma® 250 mg:</b> Requires clinical reason why 350 mg generic strength cannot be used</li> <li><b>Zanaflex® Capsules:</b> Requires clinical reason generic cannot be used</li> </ul>



## TETRACYCLINES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR <b>CAPS</b> (generic Vibramycin)	demeclocycline (generic Declomycin) <sup>CL</sup>	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a sequential 3-day trial of TWO preferred agents within this drug class</li> </ul>
doxycycline monohydrate <b>SUSP, TAB</b> (generic Vibramycin)	DORYX MPC DR (doxycycline pelletized)	
minocycline HCl <b>TAB</b> (generic Dynacin/Myrac)	doxycycline hyclate IR <b>TAB</b> (generic Vibramycin)	<p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Demeclocycline:</b> Approved for diagnosis of SIADH</li> <li><b>doxycycline suspension:</b> May be approved with documented swallowing difficulty</li> </ul>
tetracycline	doxycycline hyclate DR (generic Doryx)	
	doxycycline monohydrate <b>50MG, 100MG CAPS</b>	
	doxycycline monohydrate 40MG, 75MG and 150MG <b>CAP</b> (generic Adoxa/Monodox/ Oracea)	
	minocycline HCl <b>CAPS</b> (generic Dynacin/ Minocin/Myrac)	
	minocycline HCl ER (generic Solodyn)	
	NUZYRA (omadacycline)	
	VIBRAMYCIN SUSP (doxycycline)	
	XIMINO (minocycline ER)QL	

## THYROID HORMONES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
levothyroxine <b>TAB</b> (generic Synthroid)	ADTHYZA (thyroid, pork)	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>
liothyronine <b>TAB</b> (generic Cytomel)	ERMEZA (levothyroxine) <b>SOLN</b>	
thyroid, pork <b>TAB</b>	EUTHYROX (levothyroxine)	
UNITHROID (levothyroxine)	LEVO-T (levothyroxine)	
	levothyroxine <b>CAPS</b> (generic Tirosint)	
	THYQUIDITY (levothyroxine) <b>SOLN</b>	

## ULCERATIVE COLITIS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
<b>ORAL</b>		<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul> <p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Asacol HD®/Delzicol DR®:</b> Requires clinical reason why preferred mesalamine products cannot be used</li> </ul>
APRISO (mesalamine) <b>mesalamine (generic Lialda)</b> PENTASA (mesalamine) Sulfasalazine IR, DR (generic Azulfidine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) <b>LIALDA (mesalamine)</b> mesalamine ER (generic Apriso) mesalamine ER (generic Pentasa) mesalamine (generic Asacol HD/Delzicol)	
<b>RECTAL</b>		
mesalamine <b>SUPPOSITORY</b> (generic Canasa) Sulfite-Free ROWASA (mesalamine)	CANASA (mesalamine) mesalamine <b>ENEMA</b> (generic Rowasa) ROWASA (mesalamine) UCERIS (budesonide)	

## UTERINE DISORDER TREATMENT

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MYFEMBREE (relugolix/ estradiol/ norethindrone acetate) <sup>AL, CL, QL</sup> ORIAHNN (elagolix/ estradiol/ norethindrone) <sup>AL, CL</sup> ORILISSA (elagolix sodium) <sup>QL, CL</sup>		<p>Drug-specific criteria:</p> <ul style="list-style-type: none"> <li><b>Myfembree, Orilissa, and OriaHnn:</b> Requires an FDA approved indication, must follow FDA dosing guidelines, and have had a trial and failure of an NSAID and oral contraceptive               <ul style="list-style-type: none"> <li>Total duration of treatment is max of 24 months</li> </ul> </li> </ul>

## VASODILATORS, CORONARY

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
isosorbide dinitrate <b>TAB</b>	BIDIL (isosorbide dinitrate/	<ul style="list-style-type: none"> <li>Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class</li> </ul>
isosorbide dinitrate/hydralazine (Bidil) <sup>CL</sup>	hydralazine) <sup>CL</sup>	
isosorbide mono IR/SR <b>TAB</b>	GONITRO (nitroglycerin)	Drug-specific criteria:
nitroglycerin <b>SUBLINGUAL, TRANSDERMAL</b>	isosorbide dinitrate <b>TAB (Oceanside Pharm MFR only)</b>	<ul style="list-style-type: none"> <li><b>BiDiI/ isosorbide dinitrate-hydralazine:</b> Approved for the treatment of heart failure as an adjunct therapy to standard therapy in self-identified black patients</li> <li><b>Verquvo:</b> Approved for use in patients following a recent hospitalization for HF within the past 6 months OR need for outpatient IV diuretics, in adults with symptomatic chronic HF and EF less than 45%</li> </ul>
nitroglycerin ER <b>TAB</b>	NITRO-BID <b>OINT</b> (nitroglycerin)	
	NITRO-DUR (nitroglycerin)	
	nitroglycerin <b>TRANSLINGUAL</b> (generic Nitrolingual)	
	VERQUVO (vericiguat) <sup>AL,CL,QL</sup>	

### 5. Adjournment / Old Business

- Dr. Dering-Anderson noted that the state of Michigan had initiated a lawsuit against a PBM.
- A vote to conclude the meeting was made at 11:19 AM CST.

(1 <sup>st</sup> ) Motion: Dering-Anderson	(2 <sup>nd</sup> ) Motion:
Vote to conclude meeting unanimously approved by all in attendance.	

**The next Nebraska Medicaid Pharmaceutical and Therapeutics (P&T) Committee meeting is scheduled for:**

**Date:**

Wednesday, October 29<sup>th</sup>, 2025

**Time:**

9:00 AM – 5:00 PM CST

**Location:**

Mahoney State Park, Peter Kiewit Lodge  
28500 West Park Hwy  
Ashland, NE 68003

Recorded by: ShaLeigh Hammons, CPhT – Account Manager Senior, Prime Therapeutics