DIVISION OF MEDICAID AND LONG-TERM CARE

Nebraska Department of Health and Human Services

PHARMACEUTICAL AND THERAPEUTICS (P&T) COMMITTEE MEETING MINUTES

Wednesday, May 10th, 2023 at 9:00 AM CST Mahoney State Park, Peter Kiewit Lodge 28500 West Park Hwy, Ashland, NE 68003

Committee Members Present:

Eric Avery, M.D. (Chair) Claire Baker, M.D. Andrew Bendlin, Pharm.D. Cassie Cowles, APRN

Allison Dering-Anderson, Pharm.D. (Vice Chair)

Stephen Dolter, M.D.

Gary Elsasser, Pharm.D. (Early PM Departure)

Wade Fornander, M.D. C. Jose Friesen, M.D. Jennifer Hill, M.D. Joyce Juracek, Pharm.D.

Lauren Nelson, M.D.

Jessica Pohl, Pharm.D.

Linda Sobeski, Pharm.D. (Early PM Departure)

Sarah Stewart-Bouckaert, Pharm.D.

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.

Magellan Medicaid Administration Staff Present:

Nikia Bennette-Carter, Pharm.D., Clinical Account Executive Elanah Figueroa, B.A., Account Operations Executive

Managed Care Staff Present:

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

Committee Members Excused:

Rachelle Kaspar-Cope, M.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 vice-chair at 9:02 AM CST. The agenda was posted on the Nebraska Medicaid Pharmacy website (https://nebraska.fhsc.com/PDL/PTcommittee.asp) on April 10th, 2023. 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 members. Dianne Garside welcomed Sarah Stewart-Bouckaert, Pharm.D. as the newest P&T committee member.
- c. Roll Call: See list above.
- d. Conflict of Interest: No new conflicts of interest were reported.

e. Approval of November 16th, 2022 P&T Committee Meeting Minutes.

Approval of November 16th, 2022 P&T Committee Meeting Minutes (1st) Motion: Dolter (2nd) Motion: Fornander **Discussion:** Approve as written. Abstain Abstain Yes Yes **Voting - P&T Committee Members** ŝ ŝ **Voting - P&T Committee Members** Avery, Eric, M.D. (Chair) Friesen, C. Jose, M.D. Х Votes only in the event of a tie Baker, Claire, M.D. Hill, Jennifer, M.D. Х Х Bendlin, Andrew, Pharm.D. Juracek, Joyce, Pharm.D. Х Х Cowles, Cassie, APRN Nelson, Lauren, M.D. Х Х Dering-Anderson, Allison, Pharm.D. Pohl, Jessica, Pharm.D. Х Х Dolter, Stephen, M.D. Sobeski, Linda, Pharm.D. (Not present during х voting) Elsasser, Gary, Pharm.D. Stewart-Bouckaert, Sarah, Pharm.D. Х Fornander, Wade, M.D. Sundsboe, Bradley, Pharm.D. Х Х

f. Department information: Dianne Garside notified the committee and public attendees of recent staffing changes within the state's pharmacy department, including the resignation of Ken Saunders. There are vacant positions open within the P&T committee board, including physician roles for family or internal medicine, psychiatry or neurology, and a public member position. Any interested candidates may reach out to Dianne Garside at dhhs.medicaidpharmacyunit@nebraska.gov for more information. A Request for Proposal (RFP) round was completed for the Heritage Health program and its MCO vendors. The awarded MCOs are Molina Healthcare, Nebraska Total Care, and United Healthcare and will be effective 1/1/2024. All plans will follow the State of Nebraska's Preferred Drug List (PDL).

2. Public Testimony

Speaker Order	DRUG CLASS	Drug Name	PDL Status	Speaker Name	Affiliation
1	Antimigraine Agents, Other	Ajovy	Р	Dave Miley	Teva
2	Antimigraine Agents, Other	Qulipta	NP	Bradley Jones	AbbVie
3	HAE Treatments	Takhzyro	NP	Doug McCann	Takeda
4	HIV/AIDS	Sunlenca	NP	Porscha Showers	Gilead Sciences
5	Hypoglycemics, Incretin Mimetics/Enhancers	Ozempic	Р	Shawn Hansen	Novo Nordisk
6	Immunosuppressives, Oral	Tavneos	NP	Larry Palmisano	Amgen
7	Lipotropics, Other	Repatha	NP	Patricia Woster	Amgen
8	PAH	Tyvaso DPI	NP	Sandeep Anand	United Therapeutics Corporation

3. Committee Closed Session

(1st) Motion: Dering-Anderson (2nd) Motion: Baker
Committee Closed Session unanimously approved by all in attendance.

4. Resume Open Session

(1st) Motion: Baker (2nd) Motion: Pohl
Resume Open Session unanimously approved by all in attendance.

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 <u>only in the event of a tie.</u> The details of each vote and the associated PDL recommendations are presented in the following tables.

a. Consent Agenda

Consent Agenda

(1st) Motion: Dering-Anderson

(2nd) Motion: Sobeski

Discussion: Committee **removed** five Consent Agenda categories and **added** them to Therapeutic Class Reviews: (Antiparasitics, Topical; Antivirals, Oral; Cephalosporins and Related Antibiotics; Hepatitis C Agents; and Immunosuppressive**s**, Oral). Committee approved the amended Consent Agenda.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	Х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	Х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	Х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	Х		
Dering-Anderson, Allison, Pharm.D.	Х			Pohl, Jessica, Pharm.D.	Х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	Х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	Х		

Consent Agenda: Therapeutic categories (TC) with ur	nchanged recommendations unless otherwise indicated.
Angiotensin Modulators	Hypoglycemics, Meglitinides
Antibiotics, Gastrointestinal	Hypoglycemics, Metformins
Antibiotics, Topical	Hypoglycemics, SGLT2
Antifungals, Topical	Hypoglycemics, Sulfonylureas
Antimigraine Agents, Triptans	Hypoglycemics, TZDs
Antiparasitics, Topical (Removed)	Immunosuppressives, Oral (Removed)
Antivirals, Oral (Removed)	Lincosamides / Oxazolidinones / Streptogramins
Bone Resorption Suppression and Related Agents	Lipotropics, Statins
Cephalosporins and Related Antibiotics (Removed)	Macrolides and Ketolides
Cystic Fibrosis	Nitrofuran Derivatives
Diuretics	Pancreatic Enzymes
Fluoroquinolones, Oral	Penicillins
Growth Hormone	Platelet Aggregation Inhibitors
H. Pylori Treatment	Sinus Node Inhibitors
Hepatitis B Agents	Thyroid Hormones
Hepatitis C Agents (Removed)	Uterine Disorder Treatments
Hypoglycemics, Alpha-glucosidase Inhibitors	

b. Therapeutic Class Reviews

Review Agenda – Acne Agents, Topical

(1st) Motion: Pohl

(2nd) Motion: Fornander

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	§.	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Analgesics, Opioids Long-Acting

(1st) Motion: Elsasser

(2nd) Motion: Pohl

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	Х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	Х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Analgesics, Opioids Short-Acting

(1st) Motion: Juracek

(2nd) Motion: Dering-Anderson

Discussion: The committee recommended moving tramadol/APAP from preferred to non-preferred PDL status.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	_o N	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN		х		Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.		х		Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Androgenic Agents

(1st) Motion: Dering-Anderson

(2nd) Motion: Sobeski

Discussion: The committee recommended removing the drug-specific prior authorization criteria for

Androderm/Androgel.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	Х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.		х	
Dolter, Stephen, M.D.	Х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	Х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	Х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Angiotensin Modulator Combinations

(1st) Motion: Juracek

(2nd) Motion: Fornander

Discussion: The committee recommended updating the beginning of the class criteria to state "non-preferred agents will be approved for patients who have failed **TWO** preferred agents within this drug class within the last 12 months" to match the other drug class criteria.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.		х		Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Antibiotics, Inhaled

(1st) Motion: Hill

(2nd) Motion: Dolter

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Antibiotics, Vaginal

(1st) Motion: Sobeski

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Anticoagulants

(1st) Motion: Dolter

(2nd) Motion: Cowles

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Antiemetics / Antivertigo Agents

(1st) Motion: Dering-Anderson

(2nd) Motion: Fornander

Discussion: Approve as written with the recommendation of placing the generic name behind Metozolv ODT on the

PDL criteria.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Antifungals, Oral

(1st) Motion: Sobeski

(2nd) Motion: Pohl

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	Х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	Х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	Х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	Х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	Х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	Х			Sundsboe, Bradley, Pharm.D.	Х		

Review Agenda – Antimigraine Agents, Other

(1st) Motion: Dering-Anderson

(2nd) Motion: Hill

Discussion: Approve as written.

				<u> </u>			
Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _O	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Antiparasitics, Topical

(1st) Motion: Baker

(2nd) Motion: Hill

Discussion: The committee recommended updating the drug class criteria to "Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class **within the last 6 months.**"

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No O	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

5. Committee Moved to Closed Session (Working Lunch)

(1st) Motion: Dering-Anderson (2nd) Motion: Pohl
Committee Moved to Closed Session unanimously approved by all in attendance.

6. Committee Open Session – Consideration of Therapeutic Class Reviews – Resume Open Session:

(1st) Motion: Baker (2nd) Motion: Cowles

Resume Open Session unanimously approved by all in attendance.

a. Therapeutic Class Reviews (continued)

Review Agenda – Antivirals, Oral

(1st) Motion: Dering-Anderson

(2nd) Motion: Sobeski

Discussion: Approve as written. Spencer Moore, Pharm.D. presented 2022 quarterly anti-influenza utilization data.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	x			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D. (Not present during voting)				Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	x			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Antivirals, Topical

(1st) Motion: Dolter

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	_S	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D. (Not present during voting)				Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Beta-Blockers

(1st) Motion: Hill

(2nd) Motion: Fornander

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	S.	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	x			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Bladder Relaxant Preparations

(1st) Motion: Hill

(2nd) Motion: Fornander

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	o _N	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	x			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – BPH - Benign Prostatic Hyperplasia Agents

(1st) Motion: Cowles

(2nd) Motion: Friesen

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	o _N	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Calcium Channel Blockers

(1st) Motion: Fornander

(2nd) Motion: Hill

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	9 N	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Cephalosporins and Related Antibiotics

(1st) Motion: Elsasser

(2nd) Motion: Hill

Discussion: The committee recommended requiring an ICD -10 diagnosis of gonorrhea for cefixime or pyelonephritis for cefpodoxime for approval without prior authorization.

Abstain Abstain Yes **Voting – P&T Committee Members** Yes 8 ٤ **Voting - P&T Committee Members** Does not include excused or unexcused members Avery, Eric, M.D. (Chair)

Votes only in the event of a tie Friesen, C. Jose, M.D. Х Baker, Claire, M.D. Hill, Jennifer, M.D. Х Х Bendlin, Andrew, Pharm.D. Juracek, Joyce, Pharm.D. Х Х Cowles, Cassie, APRN Nelson, Lauren, M.D. Х Dering-Anderson, Allison, Pharm.D. Pohl, Jessica, Pharm.D. Х Х Dolter, Stephen, M.D. Sobeski, Linda, Pharm.D. Х Х Elsasser, Gary, Pharm.D. Stewart-Bouckaert, Sarah, Pharm.D. Х Х Fornander, Wade, M.D. Sundsboe, Bradley, Pharm.D. Х Х

Review Agenda – Contraceptives, Oral

(1st) Motion: Hill

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	o _N	Abstain	Voting – P&T Committee Members	Yes	oN N	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – GI Motility, Chronic

(1st) Motion: Cowles

(2nd) Motion: Fornander

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	x			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Glucagon Agents

(1st) Motion: Fornander

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – HAE Treatments

(1st) Motion: Hill

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	x			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Hepatitis C Agents

(1st) Motion: Fornander

(2nd) Motion: Cowles

Discussion: The committee discussed feedback about sobriety restrictions and acknowledged that criteria will be reviewed by the DUR Board, and requested an update of any criteria changes at the November P&T meeting.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	2	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – HIV/AIDS

(1st) Motion: Baker

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	N _o	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Sobeski, Linda, Pharm.D.	х		
Elsasser, Gary, Pharm.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda - Hypoglycemics, Incretin Mimetics / Enhancers

(1st) Motion: Baker

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Hill, Jennifer, M.D.	х		
Baker, Claire, M.D.	х			Juracek, Joyce, Pharm.D.		х	
Bendlin, Andrew, Pharm.D.	х			Nelson, Lauren, M.D.	х		
Cowles, Cassie, APRN	х			Pohl, Jessica, Pharm.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Sobeski, Linda, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		
Friesen, C. Jose, M.D.	х						

Review Agenda – Hypoglycemics, Insulin and Related Agents

(1st) Motion: Baker

(2nd) Motion: Fornander

Discussion: The committee recommended moving HUMALOG (insulin lispro) U-200 KWIKPEN from non-preferred to preferred PDL status. The committee also recommended removing the drug-specific class criteria for Humulin R U-500 Kwikpen.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Hill, Jennifer, M.D.	х		
Baker, Claire, M.D.	х			Juracek, Joyce, Pharm.D.	х		
Bendlin, Andrew, Pharm.D.	х			Nelson, Lauren, M.D.	х		
Cowles, Cassie, APRN	х			Pohl, Jessica, Pharm.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Sobeski, Linda, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		
Friesen, C. Jose, M.D.	х						

Review Agenda – Immunosuppressants, Oral

(1st) Motion: Baker

(2nd) Motion: Hill

Discussion: The committee recommended adding drug specific class criteria for TAVNEOS (avacopan) CAPS that no trial of a preferred agent required with appropriate documentation of FDA approved indications and concurrent use of standard therapy.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	8	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Hill, Jennifer, M.D.	х		
Baker, Claire, M.D.	х			Juracek, Joyce, Pharm.D.	х		
Bendlin, Andrew, Pharm.D.	х			Nelson, Lauren, M.D.	х		
Cowles, Cassie, APRN	х			Pohl, Jessica, Pharm.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Sobeski, Linda, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		
Friesen, C. Jose, M.D.	х						

Review Agenda - Lipotropics, Other

(1st) Motion: Friesen

(2nd) Motion: Fornander

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.			х	Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Multiple Sclerosis Agents

(1st) Motion: Baker

(2nd) Motion: Hill

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Opioid Dependence Treatments

(1st) Motion: Dering-Anderson

(2nd) Motion: Fornander

Discussion: Approve as written. The committee recommended reviewing this therapeutic class again at the November 2023 P&T meeting for updated market availability of a higher strength of NARCAN (naloxone) NASAL SPRAY.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	Х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – PAH - Pulmonary Arterial Hypertension Agents

(1st) Motion: Hill

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Pediatric Vitamin Preparations

(1st) Motion: Juracek

(2nd) Motion: Dolter

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.			х	Sundsboe, Bradley, Pharm.D.			х

Review Agenda – Phosphate Binders

(1st) Motion: Fornander

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Prenatal Vitamins

(1st) Motion: Pohl

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Proton Pump Inhibitors

(1st) Motion: Fornander

(2nd) Motion: Friesen

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Skeletal Muscle Relaxants

(1st) Motion: Dering-Anderson

(2nd) Motion: Hill

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D. (Not present during voting)			
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Tetracyclines

(1st) Motion: Dering-Anderson

(2nd) Motion: Hill

Discussion: The committee recommended updating the class criteria to "Non-preferred agents will be approved for

patients who have failed sequential 3-day trial of two preferred agents within this drug class."

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	No	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	Х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Ulcerative Colitis

(1st) Motion: Dering-Anderson

(2nd) Motion: Cowles

Discussion: Approve as written.

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	N _o	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	Х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

Review Agenda – Vasodilators, Coronary

(1st) Motion: Friesen

(2nd) Motion: Nelson

Voting – P&T Committee Members Does not include excused or unexcused members	Yes	Š	Abstain	Voting – P&T Committee Members	Yes	No	Abstain
Avery, Eric, M.D. (Chair) Votes only in the event of a tie				Friesen, C. Jose, M.D.	х		
Baker, Claire, M.D.	х			Hill, Jennifer, M.D.	х		
Bendlin, Andrew, Pharm.D.	х			Juracek, Joyce, Pharm.D.	х		
Cowles, Cassie, APRN	х			Nelson, Lauren, M.D.	х		
Dering-Anderson, Allison, Pharm.D.	х			Pohl, Jessica, Pharm.D.	х		
Dolter, Stephen, M.D.	х			Stewart-Bouckaert, Sarah, Pharm.D.	х		
Fornander, Wade, M.D.	х			Sundsboe, Bradley, Pharm.D.	х		

b. Complete Copy of Proposed PDL

Nebraska Medicaid - Preferred Drug List with Prior Authorization Criteria

May 2023 P&T Proposed Changes

Red Highlights indicated proposed changes.

For the most up to date list of covered drugs consult the Drug Lookup on the Nebraska Medicaid Website at https://druglookup.fhsc.com/druglookupweb/?client=nestate

- 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], beginning October 1, 2021.
- Opioids The maximum opioid dose covered will decrease from 120 Morphine Milligram Equivalents (MME) per day
 to 90 Morphine Milligram Equivalents (MME) per day. (Beginning December 1, 2020)

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
- Buprenorphine Products PA Form
- Buprenorphine Products 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

For a complete list of Claims Limitations visit:

https://nebraska.fhsc.com/Downloads/neclaimlimitations.pdf

Unless otherwise specified, the listing of a particular brand or generic name includes all dosage forms of that drug.

CL - Prior Authorization / Class Criteria apply

QL - Quantity/Duration Limit

AL- Age Limit

NR - Product was not reviewed - New Drug criteria will apply

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
adapalene (generic differin) CREAM, GEL (OTC/Rx), GEL PUMP benzoyl peroxide (BPO) WASH, LOTION clindamycin/BPO (generic Benzaclin) PUMP clindamycin phosphate PLEDGET clindamycin phosphate SOLUTION erythromycin GEL erythromycin-BPO (generic for Benzamycin) RETIN-A (tretinoin) ^{AL} CREAM, GEL	adapalene/BPO (generic Epiduo) adapalene/BPO (generic Epiduo Forte) ALTRENO (tretinoin) ^{AL} AMZEEQ (minocycline) ARAZLO (tazarotene) ^{AL} ATRALIN (tretinoin) AVAR (sulfacetamide sodium/sulfur) AVITA (tretinoin) AZELEX (azelaic acid) BENZEFOAM (benzoyl peroxide) benzoyl peroxide CLEANSER, CLEANSING BAR OTC benzoyl peroxide FOAM (generic Benzepro) benzoyl peroxide GEL OTC benzoyl peroxide GEL Rx benzoyl peroxide TOWELETTE OTC clindamycin FOAM, LOTION clindamycin GEL clindamycin/BPO (generic Acanya) GEL clindamycin/BPO (generic Duac) clindamycin/BPO (generic Duac) clindamycin/BPO (generic For Benzamycin) EVOCLIN (clindamycin) FABIOR (tazarotene foam) NEUAC (clindamycin/BPO) ONEXTON (clindamycin/BPO) ONEXTON (clindamycin/BPO) OVACE PLUS (sulfacetamide sodium) RETIN-A ^{AL} GEL, CREAM (tretinoin) sulfacetamide sulfacetamide/sulfur SUMADAN (sulfacetamide/sulfur) tazarotene CREAM,GEL ^{NR} (generic Tazorac) tazarotene FOAM (generic Fabior) TRETIN-X (tretinoin) tretinoin CREAM, GELAL (generic Avita, Retin-A) tretinoin microspheres (generic for Retin-A Micro) AL WINLEVI (clascoterone)	Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class Output Description: Output Descripti

ANALGESICS, OPIOID LONG-ACTING

AN	ALGESICS, OPIOID LONG-ACTING		
	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
fe m C tr	UTRANS (buprenorphine)QL PATCH entanyl 25, 50, 75, 100 mcg PATCHQL norphine ER TAB (generic MS Contin, Oramorph SR) DXYCONTINCL (oxycodone ER) amadol ER (generic Ultram ER)CL (TAMPZA (oxycodone) ER	BELBUCA (buprenorphine)QL BUCCAL buprenorphine BUCCAL (generic for Belbuca)AL,QL buprenorphine PATCH (generic Butrans)QL EMBEDA (morphine sulfate/ naltrexone) DURAGESIC MATRIX (fentanyl)QL fentanyl 37.5, 62.5, 87.5 mcg PATCHQL hydrocodone ER (generic for Hysingla ER)QL hydrocodone bitartrate ER (generic for Zohydro ER) hydromorphone ER (generic for Exalgo)CL HYSINGLA ER (hydrocodone ER) KADIAN (morphine ER) methadone TABCL methadone ORAL SYRCL MORPHABOND ER (morphine sulfate) morphine ER (generic for Avinza, Kadian) CAPS NUCYNTA ER (tapentadol)CL oxycodone ER (generic Oxycontin) oxymorphone ER (generic Opana ER) tramadol ER (generic Conzip)CL	The Center for Disease Control (CDC) does not recommend long acting opioids when beginning opioid treatment. Preferred agents require previous use of a long acting opioid or documentation of a trial on a short acting agent within 90 days Non-preferred agents will be approved with failure on, or intolerance to TWO preferred agents within this drug class Drug-specific criteria: Methadone: 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 Oxycontin®: Pain contract required for maximum quantity authorization

ANALGESICS, OPIOID SHORT-ACTINGQL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acetaminophen/codeine ELIXIR, TAB codeine TAB hydrocodone/APAP SOLN, TAB hydrocodone/ibuprofen hydromorphone TAB morphine CONC SOLN, SOLN, TAB oxycodone TAB, SOLN oxycodone/APAP Tramadol 50 TAB ^{AL} (generic Ultram) tramadol/APAP (generic Ultracet)	APADAZ (benzhydrocodone/APAP) ^{CL} benzhydrocodone/APAP (generic Apadaz· ^{CL} butalbital/caffeine/APAP/codeine butalbital compound w/codeine (butalbital/ASA/caffeine/codeine) carisoprodol compound-codeine (carisoprodol/ASA/codeine) dihydrocodeine/APAP/caffeine dihydrocodeine/aspirin/caffeine hydromorphone LIQUID, SUPPOSITORY (generic Dilaudid) levorphanol meperidine (generic Demerol) morphine SUPPOSITORIES NALOCET (oxycodone/APAP) NUCYNTA (tapentadol) ^{CL} oxycodone CAPS oxycodone/APAP SOLN oxycodone CONCENTRATE oxymorphone IR (generic Opana) pentazocine/naloxone PROLATE (oxycodone/APAP) ^{NR} SOLN,TAI ROXICODONE (oxycodone) SEGLENTIS (celecoxib/tramadol) ^{AL} tramadol 100mg (generic Ultram) ^{AL} tramadol (generic Qdolo) ^{AL,QL} SOLN	 Non-preferred agents will be approved for patients who have failed THREE preferred agents within this drug class within the last 12 months Note: for short acting opiate tablets and capsules there is a maximum quantity limit of #150 per 30 days. Beginning Oct. 11, 2018: Opiate limits for opiate naïve patients will consist of -prescriptions limited to a 7 day supply, AND -initial opiate prescription fill limited to maximum of 50 Morphine Milligram Equivalents (MME) per day 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 Drug-specific criteria: Apadaz: Approval for 14 days or less Nucynta®: Approved only for diagnosis of acute pain, for 30 days or less

ANALGESICS, OPIOID SHORT-ACTINGQL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NASA	AL	
	utorphanol SPRAY ^{QL} .AZANDA (fentanyl citrate)	
		Drug-specific criteria:
BUCCAL/TRANS	SMUCOSAL ^{CL}	Abstral®/Actiq®/Fentora®/ Onsolis (fentanyl): Approved only for
fe	ABSTRAL (fentanyl) ^{CL} entanyl TRANSMUCOSAL (generic Actiq) ^{CL} FENTORA (fentanyl) ^{CL}	diagnosis of cancer AND current use of long-acting opiate

ANDROGENIC AGENTS (Topical)CL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANDRODERM (testosterone) CL TRANSDERM ANDROGEL (testosterone) PUMP CL TESTIM (testosterone) TRANSDERM testosterone PUMP (generic Androgel) CL	NATESTO (testosterone) ^{CL} testosterone PACKET (generic Androgel) ^{CL} testosterone GEL, PACKET, PUMP (generic Vogelxo) testosterone (generic Axiron) testosterone (generic Fortesta) testosterone (generic Testim)	 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 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 Drug-specific criteria: Androderm®/Androgel®: Approved for Males only with diagnosis of: Primary hypogonadism (congenital or acquired) OR Hypogonadotropic hypogonadism (congenital or acquired)

ANGIOTENSIN MODULATORS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
benazepril (generic Lotensin) enalapril (generic Vasotec) lisinopril (generic Prinivil, Zestril) quinapril (generic Accupril) ramipril (generic Altace)	captopril (generic Capoten) EPANED (enalapril) ^{CL} ORAL SOLN enalapril (generic for Epaned) ^{CL} ORAL SOLN fosinopril (generic Monopril) moexepril (generic Univasc) perindopril (generic Aceon) QBRELIS (lisinopril) ^{CL} ORAL SOLN trandolapril (generic Mavik)	 Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the last 12 months Non-preferred combination products may be covered as individual prescriptions without prior authorization Drug-specific criteria: Epaned® and Qbrelis® Oral Solution: Clinical reason why oral tablet is not appropriate
ACE INHIBITOR/DIU	RETIC COMBINATIONS	
benazepril/HCTZ (generic Lotensin HCT) enalapril/HCTZ (generic Vaseretic) lisinopril/HCTZ (generic Prinzide,	captopril/HCTZ (generic Capozide) fosinopril/HCTZ (generic Monopril HCT) moexipril/HCTZ (generic Uniretic)	
ANGIOTENSIN RE	CEPTOR BLOCKERS	
irbesartan (generic Avapro) losartan (generic Cozaar) olmesartan (generic Benicar) valsartan (generic Diovan)	candesartan (generic Atacand) EDARBI (azilsartan) eprosartan (generic Teveten) telmisartan (generic Micardis)	

ANGIOT	FNSIN MODUL	_ATORS (Conti	nued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANGIOTENSIN RECEPTOR BLOCKER/DIURETIC COMBINATIONS		 Non-preferred agents will be approved for patients who have failed TWO
irbesartan/HCTZ (generic Avalide) losartan/HCTZ (generic Hyzaar) olmesartan/HCTZ (generic Benicar-HCT) valsartan/HCTZ (generic Diovan-HCT)	candesartan/HCTZ (generic Atacand-HCT) EDARBYCLOR (azilsartan/ chlorthalidone) telmisartan/HCTZ (generic Micardis-HCT)	preferred agents within this drug class within the last 12 months Non-preferred combination products may be covered as individual prescriptions without prior authorization
		Angiotensin Modulator/Calcium Channel Blocker Combinations:
	MODULATOR/ OCKER COMBINATIONS	Combination agents may be approved if there has been a trial and failure of preferred agent
amlodipine/benazepril (generic Lotrel) amlodipine/olmesartan (generic Azor) amlodipine/valsartan (generic Exforge) amlodipine/valsartan/HCTZ (generic Exforge HCT)	amlodipine/olmesartan/HCTZ (generic Tribenzor) amlodipine/telmisartan (generic Twynsta) PRESTALIA (perindopril/amlodipine) trandolapril/verapamil (generic Tarka)	
		 Direct Renin Inhibitors/Direct Renin Inhibitor Combinations: May be approved with a history of TWO preferred ACE Inhibitors or
DIRECT RENI	N INHIBITORS	Angiotensin Receptor Blockers within the last 12 months
	aliskiren (generic Tekturna) ^{QL}	_Drug Specific Criteria • Entresto: May be approved in
	DIRECT RENIN INHIBITOR COMBINATIONS	
	TEKTURNA/HCT (aliskiren/HCTZ)	 patients ages >1 years old and with a diagnosis of heart failure
NEPRILYSIN INHIBI	TOR COMBINATION	
ENTRESTO (sacubitril/valsartan) ^{CL,QL}		
ANGIOTENSIN RECEPTOR BLOCKE	ER/BETA-BLOCKER COMBINATIONS	
	BYVALSON (nevibolol/valsartan)	

ANTIBIOTICS, GASTROINTESTINAL	New Performation	Drive Authority of the IOlean Oritoria
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FIRVANQ (vancomycin) SOLN metronidazole TAB neomycin tinidazole (generic Tindamax) ^{CL}	DIFICID (fidaxomicin) CL TAB, SUSP metronidazole ^{CL} CAPS nitazoxanide (generic Alinia) TABAL, CL, QL paromomycin SOLOSEC (secnidazole) vancomycin CAPS (generic Vancocin) ^{CL} XIFAXAN (rifaximin) ^{CL}	 Note: Although azithromycin, ciprofloxacin, and trimethoprim/ sulfmethoxazole are not included in this review, they are available without prior authorization Drug-specific criteria: Alinia®: Trial and failure with metronidazole is required for a diagnosis of giardiasis Dificid®: 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. Flagyl ER®: Trial and failure with metronidazole is required Flagyl @Metronidazole 375mg capsules and Flagyl ER®/Metronidazole 750mg ER tabs: Clinical reason why the generic regular-release cannot be used tinidazole:

ANTIBIOTICS, INHALEDCL

Preferred Agents ^{CL}	Non-Preferred Agents	Prior Authorization/Class Criteria
Preferred Agents ^{CL} BETHKIS (tobramycin) KITABIS PAK (tobramycin) TOBI-PODHALER (tobramycin) tobramycin (generic Tobi)	ARIKAYCE (amikacin liposomal inh) SUSP CAYSTON (aztreonam lysine) ^{QL} tobramycin (generic Bethkis)	

ANTIBIOTICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
bacitracin OINT bacitracin/polymyxin (generic Polysporin) mupirocin OINT (generic Bactroban) neomycin/polymyxin/bacitracin (generic Neosporin, Triple AB) neomycin/polymyxin/pramoxine neomycin/polymyxin/bacitracin/ pramoxine	CENTANY (mupirocin) gentamicin OINT, CREAM mupirocin CREAM (generic Bactroban) ^{CL}	 Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class within the last 12 months Drug-specific criteria: Mupirocin® Cream: Clinical reason the ointment cannot be used

ANTIBIOTICS, VAGINAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CLEOCIN OVULES (clindamycin) clindamycin CREAM (generic Cleocin) CLINDESSE (clindamycin) metronidazole, vaginal NUVESSA (metronidazole)	CLEOCIN CREAM (clindamycin) VANDAZOLE (metronidazole) XACIATO (clindamycin phosphate) GEL ^{AL,NR}	 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

ANTICOAGULANTS

ELIQUIS (apixaban) enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg ^{CL_OL} XARELTO (rivaroxaban) XARELTO (rivaroxaban) ELIQUIS (abigatran) PRADAXA (dabigatran)	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	enoxaparin (generic Lovenox) PRADAXA (dabigatran) warfarin (generic Coumadin) XARELTO (rivaroxaban) 10 mg, 15 mg, 20 mg XARELTO (rivaroxaban) 2.5 mg ^{CL,QL}	fondaparinux (generic Arixtra) FRAGMIN (dalteparin) PRADAXA (dabigatran) PELLETS ^{NR} SAVAYSA (edoxaban) ^{CL,QL}	for patients who have failed ONE preferred agent within this drug class within the last 12 months Drug-specific criteria: Coumadin®: Clinical reason generic warfarin cannot be used Savaysa®: 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 Xarelto 2.5mg: 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 Xarelto Suspension: Approved for patients ≤12 years of age or if there is a clinical reason why a preferred solid

ANTIEMETICS/ANTIVERTIGO AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
dronabinol (generic Marinol) ^{AL}	BINOIDS CESAMET (nabilone)	 Non-preferred agents will be approved for patients who have failed ONE preferred agent within this drug class within the same group
5HT3 RECEPTO	OR BLOCKERS	_ _Drug-specific criteria:
ondansetron (generic Zofran/Zofran ODT) ^{QL}	ANZEMET (dolasetron) granisetron (generic Kytril) SANCUSO (granisetron) ^{CL} ZUPLENZ (ondansetron)	Akynzeo®/Varubi®: Approved for Moderately/Highly emetogenic chemotherapy with dexamethasone and a 5-HT3 antagonist Regimens include: AC combination (Doxorubicin or Epirubicin with
NK-1 RECEPTO	R ANTAGONIST	Cyclophosphamide), Aldesleukin, Amifostine, Arsenic trioxide,
aprepitant CAPS (generic Emend) ^{QL}	AKYNZEO (netupitant/palonosetron) ^{CL} aprepitant PACK (generic Emend) ^{QL} EMEND (aprepitant) CAPS, PACK, PWD VARUBI (rolapitant) TAB ^{CL}	Armostine, Alserile triolide, Azacitidine, Bendamustine, Busulfan, Carmustine, Carbplatin, Cisplatin, Clofarabine, Cyclophosphamide, Cytarabine, Dacarbazine, Dactinomycin, Daunorubicin, Epirubicin, Etoposide, Hexamethylmelamine, Idarubicin,
TRADITIONAL	ANTIEMETICS	Ifosfamide, Imatinib, Interferon α,
DICLEGIS (doxylamine/pyridoxine) ^{CL,QL} dimenhydrinate (generic Dramamine) OTC meclizine (generic Antivert) metoclopramide (generic Reglan) phosphoric acid/dextrose/fructose SOLN (generic Emetrol) prochlorperazine, oral (generic Compazine) promethazine SYR, TAB (generic Phenergan) promethazine SUPPOSITORY 12.5mg, 25mg TRANSDERM-SCOP (scopolamine)	BONJESTA (doxylamine/pyridoxine).CL,QL COMPRO (prochlorperazine) doxylamine/pyridoxine (generic Diclegis)CL,QL metoclopramide ODT (generic Metozolv ODT) prochlorperazine SUPPOSITORY (generic Compazine) promethazine SUPPOSITORY 50mg scopolamine TRANSDERMAL trimethobenzamide TAB (generic Tigan)	Irinotecan, Mechlorethamine, Melphalan, Methotrexate, Oxaliplatin, Procarbazine, Streptozotocin, Temozolomide Diclegis®/Bonjesta: Approved only for treatment of nausea and vomiting of pregnancy Metozolv ODT®: Documentation of inability to swallow or Clinical reason oral liquid cannot be used Sancuso®/Zuplenz®: Documentation of oral dosage form intolerance

NTIFUNGALS, ORAL Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
elotrimazole (mucous membrane, roche) luconazole SUSP, TAB (generic Diflucan) griseofulvin SUSP griseofulvin microsized TAB (systatin SUSP, TAB (s	BREXAFEMME (ibrexafungerp) ^{QL} CRESEMBA (isavuconazonium) ^{CL} flucytosine (generic Ancobon) ^{CL} griseofulvin ultramicrosize (generic GRIS-PEG) itraconazole (generic Sporanox) ^{CL} ketoconazole (generic Nizoral) NOXAFIL (posaconazole) AL.CLNR POWDERMIX nystatin POWD posaconazole (generic Noxafil) ^{AL.CL} TOLSURA (itraconazole) ^{CL} VIVJOA (oteseconazole) CAPS ^{NR} voriconazole (generic VFEND) ^{CL}	 Non-preferred agents will be approved for patients who have failed a trial of TWO diagnosis-appropriate preferred agents within this drug class Drug-specific criteria: Cresemba®: Approved for diagnosis of invasive aspergillosis or invasive mucormycosis Flucytosine: Approved for diagnosis of: Candida: Septicemia, endocarditis, UTIs Cryptococcus: Meningitis, pulmonary infections Noxafil®: No trial for diagnosis of Neutropenia Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Neutropenic hematologic malignancies, Graft vs. Host disease(GVHD), Immunosuppression secondary to hematopoietic stem cell transplant Noxafil® Powdermix: pediatric patients 2 years of age and older who weigh 40 kg or less Noxafil® Suspension: Oropharyngeal/esophageal candidiasis refractory to itraconazole and/or fluconazole Onmel®: Requires trial and failure or contraindication to terbinafine Sporanox®/Itraconazole: Approved for diagnosis of Aspergillosis, Blastomycosis, Histoplasmosis, Onychomycosis due to terbinafine-resistant dermatophytes, Oropharyngeal/ esophageal candidiasis refractory to fluconazole Sporanox®: Requires trial and failure of generic itraconazole Sporanox®: Requires trial and failure of generic itraconazole Vfend®: No trial for diagnosis of Aspergillosis, Blastomycosis, and Histoplasmosis and requires a trial and failure of generic itraconazole Vfend®: No trial for diagnosis of Myelodysplastic Syndrome (MDS), Neutropenic Acute Myeloid Leukemia (AML), Graft vs. Host disease (GVHD), Candidemia (candida krusei), Esophageal Candidiasis, Blastomycosis, S. apiospermum and Fusarium spp., Oropharyngeal/esophageal candidiasis refractory to fluconazole

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Clotrimazole CREAM (generic Lotrimin) RX, OTC clotrimazole SOLN OTC ketoconazole CREAM, SHAMPOO (generic Nizoral) LAMISIL (terbinafine) SPRAY OTC miconazole CREAM, POWD OTC nystatin terbinafine OTC tolnaftate POWD, CREAM, POWD OTC (generic Tinactin)	ALEVAZOL (clotrimazole) OTC ciclopirox CREAM, GEL, SUSP (generic Ciclodan, Loprox) ciclopirox NAIL LACQUERCL (generic Penlac) ciclopirox SHAMPOO (generic Loprox) clotrimazole SOLN RX (generic Lotrimin) DESENEX POWD OTC (miconazole) econazole (generic Spectazole) ERTACZO (sertaconazole) EXELDERM (sulconazole) FUNGOID OTC JUBLIA (efinaconazole) ^{CL} ketoconazole FOAMCL (generic Extina, Ketodan) LOPROX (ciclopirox) SUSP, SHAMPOO, CREAM LOTRIMIN AF CREAM OTC (clotrimazole) LOTRIMIN ULTRA (butenafine) luliconazole (generic Luzu) MENTAX (butenafine) miconazole OTC OINT, SPRAY SOLN miconazole/zinc oxide/petrolatum (generic Vusion) naftifine CREAM, GEL (generic Naftin) oxiconazole (generic Bensal HP) tavaborole SOLNCL (generic Kerydin) tolnaftate SPRAY, OTC	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred ager within this drug class within the last months Drug-specific criteria: Extina: Requires trial and failure of contraindication to other ketoconazole forms Jublia and tavaborole: Approved diagnoses include Onychomycosis the toenails due to <i>T.rubrum OR T. Mentagrophytes</i> ciclopirox nail lacquer: No trial required in diabetes, peripheral vascular disease (PVD), immunocompromised OR contraindication to oral terbinafine
	ROID COMBINATIONS	
	clotrimazole/betamethasone LOTION (generic Lotrisone)	

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
Autoinjector DVY (fremanezumab-vfrm) Autoinjector 3-pack ^{CL,QL} GALITY 120 mg (galcanezumab-gnlm) CL, QL PEN, SYR RTEC ODT (rimegepant) AL,CL,QL RELVY (ubrogepant) AL,CL,QL TAB	AIMOVIG (erenumab-aooe) CL,QL CAFERGOT (ergotamine/caffeine) diclofenac POWDER (generic Cambia)NR dihydroergotamine mesylate NASAL ELYXYB (celecoxib)AL,QL SOLN EMGALITY 100 mg (galcanezumab-gnlm) CL,QL SYR ERGOMAR SUBLINGUAL (ergotamine tartrate) MIGERGOT (ergotamine/caffeine) RECTAL MIGRANAL (dihydroergotamine) NASAL QULIPTA (atogepant)AL,QL REYVOW (lasmiditan)AL, CL,QL TAB TRUDHESA (dihydroergotamine mesylate)AL,QL NASAL	 In addition, all non-preferred agents will require a failed trial or contraindication of a preferred agent the same indication For Acute Treatment: agents will be approved for patients who have a failed trial or a contraindication to a triptan. For Prophylactic Treatment: Required 4 migraines per month for ≥ 3 months and has tried and failed a ≥ 1 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, metroprolol, timolol, atenolol), anti-epileptics (valproate, topiramate), ACE/ARB (lisinopril, candesartan) Drug-specific criteria: Emgality 120mg is recommended indicated for preventative treatment of Migraine Emgality 100mg is recommended indicated will only be approved for treatment of Episodic Cluster Headache Nurtec ODT: for use in acute treatment, will be approved for patient who have a failed trial or a contraindication to a triptan. For use in preventative treatment, will be approved for patients who have a failed trial of ONE preferred injectable CGRP.

ANTIMIGRAINE AGENTS, TRIPTANSQL

ANTIMIGRAINE AGENTS, TRIPTANS ^{QL} Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
rizatriptan (generic Maxalt) rizatriptan ODT (generic Maxalt MLT) sumatriptan	almotriptan (generic Axert) eletriptan (generic Relpax) frovatriptan (generic Frova) IMITREX (sumatriptan) naratriptan (generic Amerge) RELPAX (eletriptan) ^{QL} sumatriptan/naproxen (generic Treximet) zolmitriptan (generic Zomig)	 Non-preferred agents will be approved for patients who have failed ALL preferred agents within this drug class Drug-specific criteria: Sumavel® Dosepro: Requires clinical reason sumatriptan injection cannot be used Onzetra, Zembrace: approved for patients who have failed ALL preferred agents
	NASAL	_
IMITREX (sumatriptan)	ONZETRA XSAIL (sumatriptan) sumatriptan (generic Imitrex Nasal) TOSYMRA (sumatriptan) zolmitriptan (generic Zomig) ZOMIG (zolmitriptan)	
INJECTABLE		
sumatriptan KIT, SYR, VIAL	IMITREX (sumatriptan) INJECTION SUMAVEL DOSEPRO (sumatriptan) ZEMBRACE SYMTOUCH (sumatriptan)	

ANTIPARASITICS, TOPICAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
NATROBA (spinosad) permethrin 1% OTC (generic Nix) permethrin 5% RX (generic Elimite) pyrethrin/piperonyl butoxide (generic RID, A-200)	CROTAN (crotamiton) LOTION EURAX (crotamiton) CREAM, LOTION ivermectin (generic Sklice) LOTION lindane malathion (generic Ovide) spinosad (generic Natroba) VANALICE (piperonyl butoxide/pyrethrins)	Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ANTI-HERPETIC DRUGS		Non-preferred agents will be approved for patients who have failed a 10-day
acyclovir (generic Zovirax) amciclovir (generic Famvir) valacyclovir (generic Valtrex)	acyclovir (generic for Zovirax) ^{CL} SUSP SITAVIG (acyclovir buccal) ^{CL}	trial of ONE preferred agent within the same group
		Drug-specific criteria: Acyclovir Susp: Prior authorization
ANTI-INFLUENZA DRUGS		NOT required for children ≤ 12 years
oseltamivir (generic Tamiflu) ^{QL}	rimantadine (generic Flumadine) RELENZA (zanamivir) ^{QL} TAMIFLU (oseltamivir) ^{QL} XOFLUZA (baloxavir marboxil) ^{AL,CL,QL}	old Sitavig®: Approved for recurrent herpes labialis (cold sores) in immunocompetent adults Xofluza: Requires clinical, patient specific reason that a preferred agent cannot be used

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acyclovir OINT	acyclovir CREAM, (generic Zovirax) DENAVIR (penciclovir) penciclovir (generic Denavir) ^{NR} XERESE (acyclovir/hydrocortisone)	 Non-preferred agents will be approved for patients who have failed a trial with ONE preferred ORAL Antiviral agent

BETA BLOCKERS, ORAL Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atenolol (generic Tenormin) atenolol/chlorthalidone (generic Tenoretic) bisoprolol (generic Zebeta) bisoprolol/HCTZ (generic Ziac) BYSTOLIC (nebivolol) metoprolol (generic Lopressor) metoprolol ER (generic Toprol XL) propranolol (generic Inderal) propranolol ER (generic Inderal LA)	acebutolol (generic Sectral) betaxolol (generic Kerlone) HEMANGEOL (propranolol) ^{CL} SOLN INDERAL/INNOPRAN XL (propranolol ER) KAPSPARGO SPRINKLE (metoprolol ER) metoprolol/HCTZ (generic Lopressor HCT) nadolol (generic Corgard) nadolol/bendroflumethiazide nebivolol (generic Bystolic) pindolol (generic Viskin) propranolol/HCTZ (generic Inderide) timolol (generic Blocadren) TOPROL XL (metoprolol ER)	 Non-preferred agents will be approved for patients who have failed TWO diagnosis-appropriate preferred agents within this drug class Drug-specific criteria: Bystolic®: Only ONE trial is required with Diagnosis of Obstructive Lung Disease Coreg CR®: Requires clinical reason generic IR product cannot be used Hemangeol®: Covered for diagnosis of Proliferating Infantile Hemangioma Sotylize®: Covered for diagnosis of life —threatening ventricular arrhythmias OR maintenance of normal sinus rhythm in highly symptomatic atrial fibrillation/flutter (AFIB/AFL) Requires clinical reason generic sotalo cannot be used
	PHA-BLOCKERS	
carvedilol (generic Coreg) labetalol (generic Trandate)	carvedilol ER ^{CL} (generic Coreg CR)	
ANTIARE	RHYTHMIC	
sotalol (generic Betapace)	SOTYLIZE (sotalol) ^{CL}	

BLADDER RELAXANT PREPARATIONS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MYRBETRIQ (mirabegron) TAB ^{AL,QL} oxybutynin IR, ER (generic Ditropan/Ditropan XL) TOVIAZ (fesoterodine ER)	darifenacin ER (generic Enablex) fesoterodine ^{NR} (generic Toviaz) flavoxate GELNIQUE (oxybutynin) GEMTESA (vibegron) ^{AL,QL} MYRBETRIQ (mirabegron) SUSP ^{AL,CL,QL} OXYTROL (oxybutynin) solifenacin (generic Vesicare) tolterodine IR, ER (generic Detrol/ Detrol LA) trospium IR, ER (generic Sanctura/ Sanctura XR) VESICARE (solifenacin) VESICARE LS SUSP (solifenacin)	 Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class Drug-specific criteria: Myrbetriq®: Covered without trial in contraindication to anticholinergic agents Myrbetriq suspension: Covered for pediatric patients ≥ 3 years old with a diagnosis of Neurogenic Detrusor Overactivity (NDO)

BISPHOSPHONATES alendronate (generic Fosamax) TAB ibandronate (generic Boniva) ^{QL} BINOSTO (alendronate) etidronate (generic Actonel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) Calcitonin-salmon NASAL FORTEO (teriparatide) raloxifene (generic Evista) Postmenopausal women history of non-traumatic feacures BINOSTO (abaldoparatide) PTUg-specific criteria: Actonel® Combinations: Comparation of the properties of th	
alendronate (generic Fosamax) TAB ibandronate (generic Boniva) ^{QL} alendronate SOLN (generic Fosamax) CL ATELVIA DR (risedronate) BINOSTO (alendronate) etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL} OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS calcitonin-salmon NASAL FORTEO (teriparatide) ^{QL,QL} raloxifene (generic Evista) EVISTA (raloxifene) teriparatide (generic Forteo) CL,QL TYMLOS (abaloparatide) TYMLOS (abaloparatide) EVISTA (raloxifene) teriparatide (generic Forteo) CL,QL TYMLOS (abaloparatide) Binosto®: Requires clinical required for diagnosis of hete ossification Forteo®: Covered for high ris fracture High risk of fracture: BMD -3 or worse Postmenopausal women history of non-traumatic fractures Postmenopausal women more clinical risk factors Facture Forteo®: Covered for high risk fractors Postmenopausal women more clinical risk factors Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical alendronate cannot be used Etidronate disodium: Trial required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women history of non-traumatic fractures Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical required clinical required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women history of non-traumatic fractures Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical required clinical required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women more clinical risk factors Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Etidronate cannot be used Etidronate disodium: Trial required for diagnosis of hete ossification Forteo®: Covered for high risk f	ia
alendronate (generic Fosamax) TAB ibandronate (generic Boniva) ^{QL} alendronate SOLN (generic Fosamax) CL ATELVIA DR (risedronate) BINOSTO (alendronate) etidronate disodium (generic Didronel) FOSAMAX PLUS D ^{QL} risedronate (generic Actonel) ^{QL} OTHER BONE RESORPTION SUPPRESSION AND RELATED DRUGS calcitonin-salmon NASAL FORTEO (teriparatide) ^{QL,QL} raloxifene (generic Evista) EVISTA (raloxifene) teriparatide (generic Forteo) CL,QL TYMLOS (abaloparatide) TYMLOS (abaloparatide) EVISTA (raloxifene) teriparatide (generic Forteo) CL,QL TYMLOS (abaloparatide) Binosto®: Requires clinical required for diagnosis of hete ossification Forteo®: Covered for high ris fracture High risk of fracture: BMD -3 or worse Postmenopausal women history of non-traumatic fractures Postmenopausal women more clinical risk factors Facture Forteo®: Covered for high risk fractors Postmenopausal women more clinical risk factors Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical alendronate cannot be used Etidronate disodium: Trial required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women history of non-traumatic fractures Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical required clinical required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women history of non-traumatic fractures Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Atelvia DR®: Requires clinical required clinical required for diagnosis of hete ossification Forteo®: Covered for high risk fractors Postmenopausal women more clinical risk factors Drug-specific criteria: Actone® Combinations: Cc individual agents without prior authorization Etidronate cannot be used Etidronate disodium: Trial required for diagnosis of hete ossification Forteo®: Covered for high risk f	approved
 Glucocorticoid use ≥ at 7.5 dose of predniequivalent Rheumatoid Arthritis Postmenopausal women T-score ≤ -2.5 at any site clinical risk factors More than 2 units of per day Current smoker Men with primary or hypodosteoporosis Osteoporosis associated sustained systemic glucotherapy 	approved trial of the same vered as all reason consumax® of the rectures with 2 or consumated and the solone with BMD with any alcohol gonadal with corticoid
 Trial of calcitonin-salmon required Maximum of 24 months to per lifetime 	

BPH (BENIGN PROSTATIC HYPERPLASIA) TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ALPHA	ABLOCKERS	 Non-preferred agents will be approved
alfuzosin (generic Uroxatral) doxazosin (generic Cardura) amsulosin (generic Flomax) erazosin (generic Hytrin)	CARDURA XL (doxazosin) silodosin (generic Rapaflo)	for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Alfuzosin/dutasteride/finasteride
5-ALPHA-REDUC	TASE (5AR) INHIBITORS	• Covered for males only
dutasteride (generic Avodart) finasteride (generic Proscar)	dutasteride/tamsulosin (generic Jalyn) ^{CL} ENTADFI (finasteride/tadalafil) ^{NR}	 Cardura XL®: Requires clinical reas generic IR form cannot be used Flomax®: Females covered for a 7 supply with diagnosis of acute kidnestones Jalyn®: Requires clinical reason whindividual agents cannot be used

CALCIUM CHANNEL BLOCKERS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SHORT	-ACTING	 Non-preferred agents will be approved
Dihydro	pyridines	for patients who have failed a trial of ONE preferred agent within this drug
	isradipine (generic Dynacirc) nicardipine (generic Cardene) nifedipine (generic Procardia) nimodipine (generic Nimotop) NYMALIZE (nimodipine) SOLN	class Drug-specific criteria: Nifedipine: May be approved without trial for diagnosis of Preterm Labor or Pregnancy Induced Hypertension (PIH)
Non-dihyo	Iropyridines	Nimodipine: Covered without trial for diagnosis of subarachnoid hemorrhage
diltiazem (generic Cardizem) verapamil (generic Calan/Isoptin)		Katerzia/ Norliqva: May be approved with documented swallowing difficulty
LONG	ACTING	
Dihydropyridines		
amlodipine (generic Norvasc) nifedipine ER (generic Procardia XL/ Adalat CC)	felodipine ER (generic Plendil) KATERZIA (amlodipine) ^{QL} SUSP levamlodipine (generic Conjupri) ^{NR} nisoldipine (generic Sular) NORLIQVA (amolidipine) ^{AL,CL,NR,QL} SOLN	
Non-dihyo	Iropyridines	
diltiazem ER (generic Cardizem CD) verapamil ER TAB	CALAN SR (verapamil) diltiazem ER (generic Cardizem LA) MATZIM LA (diltiazem ER) TIAZAC (diltiazem) verapamil ER CAPS verapamil 360mg CAPS verapamil ER (generic Verelan PM)	

CEPHALOSPORINS AND RELATED ANTIBIOTICS, ORAL

Preferred Agents	Non-Preferred Agents		Prior Authorization/Class Criteria
BETA LACTAM/BETA-LACTAM	MASE INHIBITOR COMBINATIONS	٠	Non-preferred agents will be approved
amoxicillin/clavulanate TAB, SUSP	amoxicillin/clavulanate CHEWABLE amoxicillin/clavulanate ER (generic Augmentin XR) AUGMENTIN (amoxicillin/clavulanate) SUSP, TAB		for patients who have failed a 3-day trial of ONE preferred agent within the same group
CEPHALOSPORIN	IS – First Generation	_	
cefadroxil CAPS, SUSP (generic Duricef) cephalexin CAPS, SUSP (generic Keflex)	cefadroxil TAB (generic Duricef) cephalexin TAB		
CEPHALOSPORINS -	- Second Generation	7	
cefprozil (generic Cefzil) cefuroxime TAB (generic Ceftin)	cefaclor (generic Ceclor) CEFTIN (cefuroxime) TAB, SUSP		
CEPHALOSPORINS	- Third Generation		
cefdinir (generic Omnicef)	cefixime (generic Suprax) CAPS, SUSP cefpodoxime (generic Vantin) SUPRAX (cefixime) CAPS, CHEWABLE TAB, SUSP, TAB		

CONTRACEPTIVES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
All reviewed agents are recommended preferred at this time Only those products for review are listed. Brand name products may be subject to Maximum Allowable Cost (MAC) pricing or require substitution with a generic equivalent Specific agents can be looked up using the Drug Look-up Tool at: https://druglookup.fhsc.com/druglookupweb //?client=nestate	FINZALA (ethinyl estradiol/norethindrone acetate) CHEW NR Her Style (levonogestrel) OTC NR norethindrone/ethinyl estradiol FE estrophasic (generic EstropFE)NR	

CYSTIC FIBROSIS, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	BRONCHITOL (mannitol) ^{AL,CL,QL} KALYDECO (ivacaftor) ^{QL, AL} PACKET, TAB ORKAMBI (lumacaftor/ivacaftor) ^{QL, AL} PACKET, TAB SYMDEKO (tezacaftor/ivacaftor) ^{QL, AL} TRIKAFTA (elexacaftor, tezacaftor, ivacaftor) ^{AL, CL}	 Bronchitol: Approved for diagnosis of CF and documentation that the patient has passed the BRONCHITOL Tolerance Test Kalydeco®: Diagnosis of CF and documentation of the drug-specific, FDA-approved mutation of CFTR gene Orkambi®: Diagnosis of CF and documentation of presence of two copies of the F580del mutation (homozygous) of CFTR gene Symdeko: Diagnosis of CF and documentation of the drug specific, FDA approved mutation of CFTR gene. Trikafta: Diagnosis of CF and documentation of at least one F508del mutation in the CFTR gene

IURETICS		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
SINGLE-AGEN amiloride TAB cumetanide TAB chlorothiazide TAB chlorothalidone TAB (generic Diuril) urosemide SOLN, TAB (generic Lasix) nydrochlorothiazide CAPS, TAB (generic Microzide) ndapamide TAB metolazone TAB spironolactone TAB (generic Aldactone) orsemide TAB	CAROSPIR (spironolactone) SUSP eplerenone TAB (generic Inspra)CL ethacrynic acid CAPS (generic Edecrin) KERENDIA (finerenone) CL,QL TAB THALITONE (chlorthalidone) TAB triamterene (generic Dyrenium)	 Non-preferred agents will be approve for patients who have failed a trial of TWO preferred agents within this dructlass Eplerenone: Will be approved with a failed trial or intolerance to spironolactone, a trial with two preferred agents is not required. Kerendia: For diagnosis of chronic kidney disease associated with Typediabetes in adults, trial of a preferred agent not required.
COMBINATION	N PRODUCTS	
amiloride/HCTZ TAB spironolactone/HCTZ TAB (generic Aldactazide) triamterene/HCTZ CAPS , TAB (generic Dyazide, Maxzide)		

FLUOROQUINOLONES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ciprofloxacin TAB (generic Cipro) evofloxacin TAB (generic Levaquin)	BAXDELA (delafloxacin) ciprofloxacin ER ciprofloxacin SUSP (generic Cipro) levofloxacin SOLN moxifloxacin (generic Avelox) ofloxacin	 Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class Drug-specific criteria: Baxdela: Coverable with documented intolerance or failure of preferred MRSA agents (clindamycin, doxycycline, linezolid, sulfamethoxazole/trimethoprim) Ciprofloxacin/Levofloxacin Suspension: Coverable with documented swallowing disorders Ofloxacin: Trial of preferred not required for diagnoses of Pelvic Inflammatory Disease OR Acute Epididymitis (non-gonorrhea)

GI MOTILITY, CHRONIC		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AMITIZA (lubiprostone) ^{AL,QL} LINZESS (linaclotide) ^{QL} MOVANTIK (naloxegol oxalate) ^{QL} RELISTOR (methylnaltrexone) SYR	alosetron (generic Lotronex) IBSRELA (tenapanor) AL,NR,QL Iubiprostone (generic Amitiza) AL,QL MOTEGRITY (prucalopride succinate) RELISTOR (methylnaltrexone) CL,QL TAB SYMPROIC (naldemedine) TRULANCE (plecanatide) QL VIBERZI (eluxodoline)	 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 Drug-specific criteria: Ibsrela: Covered for diagnosis of IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.) Lotronex®/ Viberzi®: Covered for diagnosis of IBS Diarrhea Predominant type with trial and failure of loperamide AND diphenoxylate Relistor®Tab: 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. Symproic: 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 Trulance®: Covered for diagnosis of either chronic idiopathic constipation or IBS with constipation after trial of at least TWO OTC laxatives (senna, bisacodyl, etc.)

GLUCAGON AGENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BAQSIMI (glucagon) ^{AL,QL} NASAL GLUCAGON EMERGENCY (glucagon) ^{QL} INJ KIT (Lilly) glucagon ^{QL} INJ PROGLYCEM (diazoxide) SUSP ZEGALOGUE (dasiglucagon) ^{AL, QL} AUTO-INJ	diazoxide SUSP (generic Proglycem) GLUCAGON EMERGENCY (glucagon) ^{QL} INJ KIT (Fresenius) GVOKE (glucagon) ^{AL,QL} KIT, PEN, SYR, VIAL ZEGALOGUE (dasiglucagon) ^{AL,QL} SYR	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

GROWTH HORMONES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
GENOTROPIN (somatropin) NUTROPIN AQ (somatropin) NORDITROPIN (somatropin)	HUMATROPE (somatropin) OMNITROPE (somatropin) SAIZEN (somatropin) SEROSTIM (somatropin) SKYTROFA (lonapegsomatropin-tcgd) ZOMACTON (somatropin) ZORBTIVE (somatropin)	Growth Hormone PA Form Growth Hormone Criteria

H. PYLORI TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PYLERA (bismuth, metronidazole, tetracycline) ^{QL}	lansoprazole/amoxicillin/clarithromycin (generic Prevpac) ^{QL} OMECLAMOX-PAK (omeprazole, clarithromycin, amoxicillin) ^{QL} TALICIA (omeprazole/amoxicillin/rifabutin)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

HAE TREATMENTSCL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BERINERT (C1 esterase inhibitor, human) INTRAVENOUS HAEGARDA (C1 esterase inhibitor, human)AL,CL SUB-Q icatibant acetate (generic for FIRAZYR)AL SUB-Q	CINRYZE (C1 esterase inhibitor, human) ^{AL,CL} INTRAVENOUS FIRAZYR (icatibant acetate) ^{AL} SUB-Q ORLADEYO (berotralstat) CAP ^{AL,QL} RUCONEST (recombinant human C1 inhibitor) ^{AL} INTRAVENOUS TAKHZYRO (lanadelumab-flyo) ^{AL,CL} VIAL, SYR ^{NR}	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 estrogencontaining products is contraindicated 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. Drug-Specific Criteria Cinryze, Haegarda, Orladeyo, and Takhzyro, require a history of two or more HAE attacks monthly, and trial and failure or contraindication to oral danazol

HEPATITIS B TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
entecavir TABLET	adefovir dipivoxil BARACLUDE (entecavir) SOLN, TAB EPIVIR HBV (lamivudine) TAB, SOLN lamivudine hbv TAB VEMLIDY (tenofovir alafenamide fumarate)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug Specific Criteria tenofovir disoproxil fumarate (generic Viread) tablet: Diagnosis for use required. May be indicated for chronic hepatitis B or HIV-1 infection. See HIV/AIDS class for drug listing and placement

HEPATITIS C TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DIRECT ACTING ANTI-VIRAL		Hepatitis C Treatments PA Form Hepatitis C Criteria
MAVYRET (glecaprevir/pibrentasvir) ^{CL} TAB, PELLET^{AL} OSEVI (sofosbuvir/velpatasvir/ voxilaprevir) ^{CL}	HARVONI 200/45MG, TAB (sofosbuvir/ledipasvir) ^{CL} HARVONI (ledipasvir/sofosbuvir) ^{CL} PELLET sofosbuvir/ledipasvir (generic Harvoni) ^{CL} SOVALDI (sofosbuvir) ^{CL} PELLET SOVALDI TAB (sofosbuvir) ^{CL} VIEKIRA PAK (ombitasvir/ paritaprevir/ritonavir/dasabuvir) ^{CL} ZEPATIER (elbasvir/grazoprevir) ^{CL}	 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 Patients newly eligible for Medicaid will be allowed to complete treatment with the original that treatment was initially authorized by another payor Drug-specific criteria: Trial with with a preferred agent not required in the following: Harvoni: Post liver transplant for genotype 1 or 4 Vosevi: Requires documentation of non-response after previous treatment course of Direct Acting Anti-viral agent
RIBAVIRIN		 (DAA) for genotype 1-6 without cirrhosis or with compensated cirrhosis
pavirin 200mg CAPSULE, TABLET		
INTERF	ERON	-
EGASYS (pegylated interferon alfa-2a) CL		
GASYS (pegylated interferon alfa-2a) CL		

HIV / AIDS ^{CL}		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CAPSII	D INHIBITOR	All agents require:
	SUNLENCA (lenacapavir) ^{NR, QL}	 Diagnosis of HIV/AIDS required; OR
CCR5 A	NTAGONISTS	 Diagnosis of Pre and Post Exposure Prophylaxis
SELZENTRY SOLN, TAB (maraviroc)	maraviroc (generic Selzentry)	 Non-preferred agents will be approved for patients who have a diagnosis of HIV/AIDS and patient specific
HIV-1 ATTAC	CHMENT INHIBITOR	documentation of why the preferred products within this drug class are not appropriate for patient, including, but
	RUKOBIA ER (fostemsavir) ^{AL,QL}	not limited to, drug resistance or concomitant conditions not
INTEGRASE STRAND TR	RANSFER INHIBITORS (INSTIS)	recommended with preferred agents
ISENTRESS (raltegravir) ^{QL} ISENTRESS HD (raltegravir) TIVICAY (dolutegravir)	TIVICAY PD (dolutegravir)	 Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy
NON-NUCLEOSIDE REVERSE TR	RANSCRIPTASE INHIBITORS (NNRTIS)	
EDURANT (rilpivirine) efavirenz CAPS, TABLET (generic Sustivalintelence (etravirine) ^{QL} PIFELTRO (doravirine) ^{QL}	etravirine (generic Intelence) ^{QL} nevirapine IR, ER (generic Viramune/Viramune XR) RESCRIPTOR (delavirdine) SUSTIVA CAPS, TABLET (efavirenz) VIRAMUNE (nevirapine) SUSP	
NUCLEOSIDE REVERSE TRA	NSCRIPTASE INHIBITORS (NRTIs)	
abacavir SOLN, TABLET (generic Ziagen EMTRIVA CAPS, SOLN (emtricitabine) lamivudine SOLN, TABLET (generic Epivir) zidovudine CAPS, SYRUP, TABLET (generic Retrovir)	didanosine DR (generic Videx EC) emtricitabine CAPS (generic for Emtriva) EPIVIR (lamivudine) RETROVIR (zidovudine) stavudine CAPS (generic Zerit) VIDEX (didanosine) SOLN ZIAGEN (abacavir)	
NUCLEOTIDE REVERSE TRA	ANSCRIPTASE INHIBITORS (NRTIs)	
tenofovir TABLET (generic Viread)	VIREAD (tenofovir) POWDER	
PHARMACOK	INETIC ENHANCER	
	TYBOST (cobicistat) ^{QL}	

HIV / AIDSCL (Continued)

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atazanavir CAPS (generic Reyataz) NORVIR (ritonavir) TAB	APTIVUS CAPS, SOLN (tipranavir) CRIXIVAN (indinavir) fosamprenavir TAB (generic Lexiva) INVIRASE (saquinavir) LEXIVA SUSP (fosamprenavir) NORVIR POWD, SOLN (ritonavir) PREZISTA (darunavir) SUSP, TAB REYATAZ POWD (atazanavir) ritonavir TAB (generic Norvir) VIRACEPT (nelfinavir)	 All agents require: Diagnosis of HIV/AIDS required; OR Diagnosis of Pre and Post Exposure Prophylaxis 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 Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy

HIV / AIDSCL (Continued) Preferred Agents Non-Preferred Agents Prior Authorization/Class Criteria All agents require: COMBINATION PROTEASE INHIBITORS (PIs) or PIs plus PHARMACOKINETIC Diagnosis of HIV/AIDS **ENHANCER** required: OR KALETRA **SOLN** (lopinavir/ritonavir) EVOTAZ (atazanavir/cobicistat)QL Diagnosis of Pre and Post KALETRA **TAB** (lopinavir/ritonavir) PREZCOBIX (darunavir/cobicistat)^{QL} lopinavir/ritonavir SOLN, TAB (generic Exposure Prophylaxis Kaletra) 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 Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy COMBINATION NUCLEOS(T)IDE REVERSE TRANSCRIPTASE INHIBITORS abacavir/lamivudine/zidovudine (generic abacavir/lamivudine (generic Epzicom) CIMDUO (lamivudine/tenofovir)QL Trizivir) DESCOVY (emtricitabine/tenofovir)QL COMBIVIR (lamivudine/zidovudine) emtricitabine/tenofovir (generic Truvada) EPZICOM (abacavir sulfate/lamivudine) lamivudine/zidovudine (generic TEMIXYS (lamivudine/tenofovir)QL TRIZIVIR (abacavir/lamivudine/zidovudine) Combivir) TRUVADA (emtricitabine/tenofovir)

Preferred Agents Non-Preferred Agents	Prior Authorization/Class Criteria
BIKTARVY (bictegravir/emtricitabine/tenofovir) tenofovir) DELSTRIGO (doravirine/lamivudine/tenofovir) DOVATO (dolutegravir/lamivudine) CENVOYA (elvitegravier/cobicistat/entricitabine/tenofovir) denofovir) CILUCA (dolutegravir/lamividine) CILUCA (dolutegravir/lamividine) CILUCA (dolutegravir/lamividine) CILUCA (dolutegravir/rilpivirine) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (dolutegravir/cobicistat/entricitabine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (davirenz/lamivudine) CILUCA (davirenz/lamivudine) CILUCA (davirenz/lamivudine) CILUCA (davirenz/lamivudine) CILUCA (davirenz/lamivudine) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) CILUCA (davirenz/lamivudine/tenofovir) (generic for Symfi Lo) CILUCA (davirenz/lamivudine/tenofovir) (generic for Symfi) Lo) CILUCA (davirenz/lamivudine/tenofovir) (generic f	All agents require: Diagnosis of HIV/AIDS required; OR Diagnosis of Pre and Post Exposure Prophylaxis 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 Patients undergoing treatment at the time of any preferred status change will be allowed to continue therapy

HYPOGLYCEMIC	A 2	DHY-CI	LICOSII	ASE	NNHIR	TOPS
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Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
acarbose (generic for Precose)	miglitol (generic for Glyset) GLYSET (miglitol)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

HYPOGI YCEMICS	INCRETIN MIMETICS/	FNHANCERS
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HYPOGLYCEMICS, INCRETIN MIMETICS/ENHANCERS				
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria		
GLUCAGON-LIKE PEPTIDE-1 RE	CEPTOR AGONIST (GLP-1 RA) ^{CL}	GLP-1 RA Criteria		
OZEMPIC (semaglutide) TRULICITY (dulaglutide) VICTOZA (liraglutide) subcutaneous	ADLYXIN (lixisenatide) SUB-Q BYDUREON BCISE (exenatide)QL PEN BYETTA (exenatide) SUB-Q MOUNJARO (tirazepatide)NR PEN RYBELSUS (semaglutide) TAB	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR A diagnosis of ASCVD associated with a diagnosis of Type II diabetes (no metformin trial required)		
		Non-preferred agents will be approved for patients who have: ■ Failed a trial of TWO preferred agents within GLP-1 RA AND ■ Diagnosis of diabetes with HbA1C ≥ 7		
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INSULIN/GLP-1 RA	A COMBINATIONS	 Trial of metformin, or contraindication or intolerance to metformin 		
	SOLIQUA (insulin glargine/lixisenatide) XULTOPHY (insulin degludec/liraglutide)			
AMYLIN	ANALOG	Amylin Analog Criteria		
	SYMLIN (pramlintide) subcutaneous	 ALL criteria must be met Concurrent use of short-acting mealtime insulin Current therapy compliance No diagnosis of gastroparesis HbA1C ≤ 9% within last 90 days Monitoring of glucose during initiation of therapy 		
DIPEPTIDYL PEPTIDASI	E-4 (DPP-4) INHIBITOR ^{QL}	DPP-4 Inhibitor Criteria		
JANUMET (sitagliptin/metformin) JANUMET XR (sitagliptin/metformin) JANUVIA (sitagliptin) JENTADUETO (linagliptin/metformin) TRADJENTA (linagliptin)	alogliptin (generic for Nesina) alogliptin/metformin (generic for Kazano) GLYXAMBI (empagliflozin/linagliptin) JENTADUETO XR (linagliptin/metformin) KOMBIGLYZE XR (saxagliptin/metformin) ONGLYZA (saxagliptin) alogliptin/pioglitazone (generic for Oseni) QTERN (dapagliflozin/saxagliptin) STEGLUJAN (ertugliflozin/sitagliptin) TRIJARDY XR (empagliflozin/linagliptin/metformin) AL	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to 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		

HYPOGLYCEMICS. INSULIN AND RELATED DRUGS

H	HYPOGLYCEMICS, INSULIN AND RELATED DRUGS					
	Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria			
	APIDRA (insulin glulisine) SOLOSTAR, VIAL HUMALOG (insulin lispro) U-100 CARTRIDGE, PEN, VIAL HUMALOG JR. (insulin lispro) U-100 KWIKPEN HUMALOG MIX VIAL (insulin lispro/lispro protamine) HUMALOG MIX KWIKPEN (insulin lispro/lispro protamine) HUMULIN (insulin) VIAL HUMULIN 70/30 VIAL HUMULIN TO/30 VIAL HUMULIN OTC PEN HUMULIN OTC PEN HUMULIN 70/30 OTC PEN insulin aspart (generic for Novolog) insulin aspart/insulin aspart protamine PEN, VIAL (generic for Novolog Mix) insulin glargine PEN, VIAL insulin lispro (generic for Humalog) PEN, VIAL, JR KWIKPEN LANTUS SOLOSTAR PEN (insulin glargine) LANTUS (insulin glargine) VIAL LEVEMIR (insulin detemir) PEN, VIAL NOVOLIN (insulin) PEN NOVOLOG (insulin aspart) CARTRIDGE, FLEXPEN, VIAL NOVOLOG MIX FLEXPEN, VIAL (insulin aspart/aspart protamine)	ADMELOG (insulin lispro) PEN, VIAL AFREZZA (regular insulin) INHALATION BASAGLAR (insulin glargine, rec) KWIKPEN FIASP (insulin aspart) CARTRIDGE, PEN, VIAL HUMALOG (insulin lispro) U-200 KWIKPEN insulin degludec (generic Tresiba) NR 100U/mL PEN, VIAL insulin degludec (generic Tresiba) NR 200U/mL PEN insulin Glargine-YFGN PEN, VIAL (generic for Semglee-YFGN) insulin lispro/lispro protamine KWIKPEN (Humalog Mix Kwikpen) LYUMJEV KWIKPEN, VIAL(insulin lisproaabc) LYUMJEV(insulin lispro-aabc) TEMPO PENNR NOVOLIN (insulin) NOVOLIN 70/30 VIAL(insulin) NOVOLOG MIX (insulin aspart/aspart protamine) VIAL SEMGLEE (insulin glargine) PEN, VIAL TOUJEO SOLOSTAR (insulin glargine) TRESIBA (insulin degludec)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Afrezza®: Approved for T1DM on long-acting insulin with no current history of smoking or chronic lung disease Humulin® R U-500 Kwikpen:			

HYPOGLYCEMICS, MEGLITINIDES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
repaglinide (generic for Prandin)	nateglinide (generic for Starlix) ^{CL} repaglinide/metformin (generic for Prandimet) ^{CL}	 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

HYPOGLYCEMICS, METFORMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
metformin IR & ER (generic Glucophage/Glucophage XR)	metformin ER (generic Fortamet/Glumetza) metformin SOLN (generic Riomet) RIOMET ER (metformin ER) ^{AL}	 Metformin ER (generic Fortamet®)/Glumetza®: Requires clinical reason why generic Glucophage XR® cannot be used Metformin solution: Prior authorization not required for age <7 years

HYPOGLYCEMICS, SGLT2

HYPOGLYCEMICS, SGLT2 Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
FARXIGA (dapagliflozin) ^{QL,CL} INVOKAMET (canagliflozin/metformin) ^{QL, CL} INVOKANA (canagliflozin) ^{CL} JARDIANCE (empagliflozin) ^{QL, CL} SYNJARDY (empagliflozin/metformin) ^{AL,CL,QL} XIGDUO XR (dapagliflozin/metformin) ^{QL,CL}	INVOKAMET XR (canagliflozin/metformin) ^{QL} SEGLUROMET (ertugliflozin/metformin) ^{QL} STEGLATRO (ertugliflozin) ^{QL} SYNJARDY XR (empagliflozin/metformin) ^{AL,QL}	Preferred agents require a diagnosis of Type II diabetes AND a trial and failure or intolerance to metformin OR A diagnosis of ASCVD or Heart Failure, or Chronic Kidney Disease associated with a diagnosis of Type II diabetes (no metformin trial required)
		 Non-preferred agents will be approved for patients who have failed a trial with ONE preferred agent within this drug class
		Drug Specific Criteria: Farxiga: May be approved for a diagnosis of heart failure with reduced ejection fraction (NYHA class II-IV) without a diagnosis of diabetes May be approved for a diagnosis of chronic kidney disease at risk of progression without a diagnosis of diabetes
		Jardiance: May be approved for a diagnosis of Heart Failure without a diagnosis of diabetes

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)	chlorpropamide tolazamide tolbutamide	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class
SULFONYLURE	A COMBINATIONS	
glipizide/metformin glyburide/metformin (generic Glucovance)		

HYPOGLYCEMICS. TZD

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
THIZAOLIDINEDIONES (TZDs)		 Non-preferred agents will be approved for patients who have failed a trial of THE preferred agent within this drug class
pioglitazone (generic for Actos)		
TZD COMBINATIONS		
	pioglitazone/glimepiride (generic for Duetact) pioglitazone/metformin (generic for Actoplus Met)	Combination products: Require clinical reason why individual ingredients cannot be used

IMMUNOSUPPRESSIVES, ORAL

IMMUNOSUPPRESSIVES, ORAL		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
azathioprine (generic Imuran) azathioprine (generic Azasan) ^{NR} cyclosporine, modified CAPS (generic Neoral) everolimus (generic for Zortress) ^{AL} mycophenolate CAPS , TA (generic Cellcept) RAPAMUNE (sirolimus) SOLN RAPAMUNE (sirolimus) TAB tacrolimus	ASTAGRAF XL (tacrolimus) AZASAN (azathioprine) cyclosporine CAP, SOFTGEL cyclosporine, modified SOLN (generic Neoral) ENVARSUS XR (tacrolimus) GENGRAF (cyclosporine, modified) CAP, SOLN mycophenolate SUSP (generic Cellcept) mycophenolic acid MYFORTIC (mycophenolate sodium) PROGRAF (tacrolimus) CAPS, PACKET REZUROCK (belumosudil)AL,QL TAB SANDIMMUNE (cyclosporine) CAPS, SOLN sirolimus SOLN, TAB (generic Rapamune) TAVNEOS (avacopan)QL CAPS ZORTRESS (everolimus) AL	Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Patients established on existing therapy will be allowed to continue

LINCOSAMIDES / OXAZOLIDINONES / STREPTOGRAMINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
clindamycin CAPS clindamycin palmitate SOLN linezolid TAB	CLEOCIN (clindamycin) CAPS CLEOCIN PALMITATE (clindamycin) linezolid SUSP SIVEXTRO (tedizolid phosphate) ZYVOX (linezolid) SUSP, TAB	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

LIPOTROPICS, OTHER

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BILE ACID SE	QUESTRANTS	Non-preferred agents will be approved
cholestyramine (generic Questran) colestipol TAB (generic Colestid)	colesevelam (generic Welchol) TAB, PACKET colestipol GRANULES (generic Colestid) QUESTRAN LIGHT (cholestyramine)	for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Colesevelam: Trial not required for diabetes control and monotherapy with metformin, sulfonylurea, or insulin has
TREATMENT OF HOMOZYGOUS FA	MILIAL HYPERCHOLESTEROLEMIA	been inadequate
	JUXTAPID (lomitapide) ^{CL} KYNAMRO (mipomersen) ^{CL}	 Juxtapid®/ Kynamro®: Approved for diagnosis of homozygous familial hypercholesterolemia (HoFH) OR
FIBRIC ACID	DERIVATIVES	Treatment failure/maximized
fenofibrate (generic Tricor) fenofibrate (generic Lofibra) gemfibrozil (generic Lopid)	fenofibric acid (generic Fibricor/Trilipix) fenofibrate (generic Antara/Fenoglide/ Lipofen/Triglide)	dosing/contraindication to ALL the following: statins, ezetimibe, niacin, fibric acid derivatives, omega-3 agents, bile acid sequestrants
NIACIN		 Require faxed copy of REMS PA
niacin ER (generic Niaspan)	NIACOR (niacin IR)	- form - Vascepa®: Approved for TG ≥ 500
OMEGA-3 F	ATTY ACIDS	
omega-3 fatty acids (generic Lovaza) VASCEPA (icosapent)	icosapent (generic Vascepa) ^{CL} omega-3 OTC	
CHOLESTEROL ABS	ORPTION INHIBITORS	
ezetimibe (generic Zetia)	NEXLETOL (bempedoic acid) NEXLIZET (bempedoic acid/ ezetimibe) QL	

LIPOTROPICS, OTHER (continued)		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
PROPROTEIN CONVERTASE SUBTILI	SIN/KEXIN TYPE 9 (PCSK9) INHIBITORS	 Praluent®: Approved for diagnoses of: atherosclerotic cardiovascular
PRALUENT (alorocumab) ^{CL}	REPATHA (evolocumab) ^{CL}	disease (ASCVD) heterozygous familial hypercholesterolemia (HeFH) Homozygous familial hypercholesterolemia (HoFH) as an adjunct to other LDL-C lowering therapies
		AND
		 Maximized high-intensity statin WITH ezetimibe for at 3 continuous months Failure to reach target LDL-C levels: ASCVD - < 70 mg/dL, HeFH - < 100 mg/dL
		Repatha®: Approved for:
		adult diagnoses of atherosclerotic cardiovascular disease (ASCVD)
		 heterozygous familial hypercholesterolemia (HeFH) in adults and pediatric patients aged 10 years and older
		 homozygous familial hypercholesterolemia (HoFH) in adults and pediatric patients aged 10 years and older
		statin-induce rhabdomyolysis AND
		 Maximized high-intensity statin WITH ezetimibe for 3+ continuous months
		 Failure to reach target LDL-C levels: ASCVD - < 70 mg/dL, HeFH - < 100 mg/dL
		Concurrent use of maximally- tolerated statin must continue, except for statin-induced rhabdomyolysis or a contraindication to a statin

LIPOTROPICS, STATINS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
atorvastatin (generic Lipitor) ^{QL} lovastatin (generic Mevacor) pravastatin (generic Pravachol) rosuvastatin (generic Crestor) simvastatin (generic Zocor)	ALTOPREV (lovastatin ER) ^{CL} EZALLOR SPRINKLE (rosuvastatin) ^{QL} fluvastatin IR/ER (generic Lescol/ Lescol XL) LIVALO (pitavastatin) ZYPITAMAG (pitavastatin)	 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 Drug-specific criteria: Altoprev®: One of the TWO trials must be IR lovastatin Combination products: Require clinical reason why individual ingredients cannot be used
STATIN COI	atorvastatin/amlodipine (generic Caduet) simvastatin/ezetimibe (generic Vytorin)	 fluvastatin ER: Requires trial of TWO preferred agents AND trial of IR fluvastatin OR clinical reason IR cannot be used simvastatin/ezetimibe: Approved for 3-month continuous trial of ONE standard dose statin

MACROLIDES AND KETOLIDES, ORAL

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MACROLIDES		Non-preferred agents require clinical
azithromycin (generic Zithromax) clarithromycin TAB , SUSP (generic Biaxin) E.E.S. SUSP (erythromycin ethylsuccinate)	clarithromycin ER (generic Biaxin XL) E.E.S. TAB (erythromycin ethylsuccinate) ERY-TAB (erythromycin) erythromycin ethylsuccinate SUSP ERYPED SUSP (erythromycin) ERYTHROCIN (erythromycin) erythromycin base TAB, CAPS	reason why preferred products within this drug class cannot be used AND a 3-day trial on a preferred product

MULTIPLE SCLEROSIS DRUGS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
AVONEX (interferon beta-1a) ^{QL} BETASERON (interferon beta-1b) ^{QL} COPAXONE 20mg (glatiramer) ^{QL} dimethyl fumarate (generic for Tecfidera) fingolimod (generic Gilenya) ^{NR,QL} KESIMPTA (Ofatumumab) ^{QL,QL} teriflunomide (generic Aubagio) ^{NR,QL}	AUBAGIO (teriflunomide) ^{QL} BAFIERTAM (monomethyl fumarate) ^{QL} dalfampridine (generic Ampyra) ^{QL} EXTAVIA (interferon beta-1b) ^{QL} GILENYA (fingolimod) ^{QL} glatiramer (generic Copaxone) ^{QL} MAVENCLAD (cladribine) MAYZENT (siponimod) ^{QL} PLEGRIDY (peginterferon beta-1a) ^{QL} PONVORY (ponesimod) REBIF (interferon beta-1a) ^{QL} TASCENSO ODT (fingolimod) TAB ^{AL,NR} TECFIDERA (dimethyl fumarate) VUMERITY (diroximel) ^{QL} ZEPOSIA (ozanimod) ^{AL,CL,QL}	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agent within this drug class Drug-specific criteria: Ampyra®: Approved for diagnosis of gait disorder associated with MS AND EDSS score ≤ 7 Plegridy: Approved for diagnosis of relapsing MS Kesimpta: Approved for patients who have failed a trial of a preferred injectable agent within this class Zeposia: 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.

NITROFURAN DERIVATIVES

NITROFURAN DERIVATIVES		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
nitrofurantoin macrocrystals CAPS (generic Macrodantin) nitrofurantoin monohydrate-macrocrystals CAPS (generic Macrobid)	nitrofurantoin SUSP (generic Furadantin)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class

OPIOID DEPENDENCE TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
buprenorphine SL buprenorphine/naloxone TAB (SL) SUBOXONE (buprenorphine/ naloxone) FILM	buprenorphine/naloxone FILM LUCEMYRA (lofexidine) ^{CL,QL} ZUBSOLV (buprenorphine/naloxone)	Buprenorphine PA Form Buprenorphine Informed Consent 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. Drug-specific criteria: Lucemyra: Approved for FDA approved indication and dosing per label. Trial of preferred product not required.

OPIOID-REVERSAL TREATMENTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
naloxone NASAL SPRAY,SYR, VIAL naltrexone TAB	KLOXXADO (naloxone) NASAL NARCAN (naloxone) NASAL SPRAY ZIMHI (naloxone) ^{AL} SYR	 Non-preferred agents will be approved with documentation of why preferred products within this drug class are not appropriate for the patient

PAH (PULMONARY ARTERIAL HYPERTENSION AGENTS), ORAL AND INHALED

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
ambrisentan (generic Letairis) REVATIO (sildenafil) ^{CL} SUSP, TAB tadalafil (generic for Adcirca) ^{CL} TRACLEER (bosentan) TAB TYVASO (treprostinil) INHALATION VENTAVIS (iloprost) INHALATION	ADEMPAS (riociguat) ^{CL} ADCIRCA (tadalafil) ^{CL} bosentan (generic Tracleer) TAB LETAIRIS (ambrisentan) OPSUMIT (macitentan) ORENITRAM ER (treprostinil) sildenafil (generic Revatio) ^{CL} SUSP, TAB TADLIQ (tadalafil) ^{NR} SUSP TRACLEER (bosentan) TAB FOR SUSPENSION TYVASO DPI (treprostini) ^{NR} INHALATION POWDER UPTRAVI (selexipag)	 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 Drug-specific criteria: Adcirca®/Revatio®: Approved for diagnosis of Pulmonary Arterial Hypertension (PAH) Adempas®:

PANCREATIC ENZYMES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CREON ZENPEP (pancrelipase)	PERTZYE (pancrelipase) VIOKACE (pancrelipase)	 Non-preferred agents will be approved for patients who have failed a trial of TWO preferred agents within this drug class

PEDIATRIC VITAMIN PREPARATIONS

PEDIATRIC VITAMIN PREPARATIONS		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
CHILD CHEW + IRON (MULTIVITAMIN WITH IRON) CHEW	DEKAs PLUS ^{AL}	 Non-preferred agents will be approved for patients who have failed a trial of
CHILDREN'S CHEWABLES (PEDI MULTIVIT NO.31/IRON/FOLIC,	FLORIVA (PEDI MULTIVIT NO.85/FLUORIDE) CHEW	TWO preferred agents within this drug class
PEDI MULTIVIT NO.25/FOLIC ACID, PEDI MULTIVIT NO.23/FOLIC ACID)	FLORIVA PLUS (PEDI MULTIVIT NO.161/FLUORIDE DROP	 Drug specific criteria: DEKAs Plus: Approved for diagnosis of Cystic Fibrosis and does not require
MULTIVIT-FLUOR (PEDI MULTIVIT NO.17 W-FLUORIDE,	MULTI-VIT-FLOR (PEDI MULTIVIT NO.205/FLUORIDE) CHEW	a trial of a preferred agent
PEDI MULTIVIT NO.16 W-FLUORIDE) CHEW	POLY-VI-FLOR (PEDI MULTIVIT NO.33/FLUORIDE) CHEW	
MULTIVIT-FLUOR (PEDI MULTIVIT NO.2 W-FLUORIDE) DROP	POLY-VI-FLOR (PEDI MULTIVIT NO.37 W- FLUORIDE) DROPS	
MULTIVIT-IRON-FLUOR (PEDI MULTIVIT 45/FLUORIDE/IRON)	POLY-VI-FLOR /0.25mg IRON (PEDI MULTIVIT 37/FLUORIDE/IRON)	
PED MVIT A,C,D3 NO.21/FLUORIDE		
POLY-VI-SOL (PEDIATRIC MULTIVITAMIN NO.192) DROP	POLY-VI-FLOR /0.5mg IRON (PEDI MULTIVIT 33/FLUORIDE/IRON)	
POLY-VI-SOL WITH IRON (PEDI MV NO.189/FERROUS SULFATE) DROPS	POLY-VI-FLOR 0.5 MG (PED MVI NO. 217/Fluoride) GUMMY	
TRI-VI-SOL (VIT A PALMITATE/VIT C/VIT D3) DROP S	QUFLORA (PEDI MULTIVIT NO.157/FLUORIDE) GUMMIES	
TRI-VITE-FLUORIDE (PED MVIT A,C,D3 NO.21/FLUORIDE)	QUFLORA FE (PED MULTIVIT 142/IRON/FLUORIDE) CHEW	
	QUFLORA FE (PED MULTIVIT 151/IRON/FLUORIDE) DROP	
	QUFLORA PED (PEDI MULTIVIT NO.63 W- FLUORIDE) CHEW	
	QUFLORA PED (PEDI MULTIVIT 84 WITH FLUORIDE, PEDI MULTIVIT NO.83 W- FLUORIDE) DROP	
	TRI-VI-FLOR (PED MVIT A,C,D3 NO.38/FLUORIDE) DROPS	

PENICII I INS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
amoxicillin CAPS, CHEWABLE TAB, SUSP, TAB ampicillin CAPS dicloxacillin penicillin VK		 Non-preferred agents will be approved for patients who have failed a 3-day trial of ONE preferred agent within this drug class

PHOSPHATE BINDERS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
calcium acetate TAB CALPHRON OTC (calcium acetate) RENVELA (sevelamer carbonate) TAB, PWD PACK	AURYXIA (ferric citrate) calcium acetate CAPS ELIPHOS (calcium acetate) lanthanum (generic FOSRENOL) PHOSLO (calcium acetate) PHOSLYRA (calcium acetate) RENAGEL (sevelamer HCI) sevelamer HCI (generic Renagel) sevelamer carbonate (generic Renvela) VELPHORO (sucroferric oxyhydroxide)	 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

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)	 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

PRENATAL VITAMINS		
Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
COMPLETENATE CHEW TAB EXPECTA PRENATAL OTC FE C/FA FE C/VIT C/VIT B12/FA OTC MARNATAL-F O-CAL FA PNV2/IRON B-G SUC-P/FA/OMEGA-3 PNV 11-IRON FUM-FOLIC ACID-OM3 PNV NO.15/IRON FUMARATE/FA CHEW TAB PNV NO.15/IRON FUM & PS CMP/FA PNV W-CA NO.40/IRON FUM/FA CMB NO.1 PNV WITH CA,NO.72/IRON/FA PNV#16/IRON FG SUC-P/FA/OMEGA-3 PNV2/IRON B-G SUC-P/FA/OMEGA-3 PRENATAL VIT #76/IRON,CARB/FA PRENATAL VIT NO.78/IRON/FA PRENATAL VIT/FE FUMARATE/FA OTC PUREFE OB PLUS STUART ONE OTC TRINATAL RX 1 VITAFOL CHEW TAB VITAFOL ULTRA VP-PNV-DHA	CITRANATAL B-CALM COMPLETENATE CHEW TAB DERMACINRX PRENATRIX OTC DERMACINRX PRETRATE TAB OTC ENBRACE HR MULTI-MAC OTC NESTABS NESTABS ABC NESTABS ONE OB COMPLETE ONE OB COMPLETE PETITE OB COMPLETE PETITE OB COMPLETE WITH DHA PNV COMBO#47/IRON/FA #1/DHA PNV WITH CA NO.68/IRON/FA NO.1/DHA PNV WITH CA,NO.72/IRON/FA PNV WITH CA,NO.74/IRON/FA OTC PNV119/IRON FUMARATE/FA/DSS PRENATAL + DHA OTC PRENATE AM PRENATE CHEW TAB PRENATE ELITE PRENATE ESSENTIAL PRENATE ESSENTIAL PRENATE ESSENTIAL PRENATE STAR PRIMACARE SELECT-OB + DHA SELECT-OB CHEW TAB TENDERA-OB OTC TRICARE TRINATAL RX 1 TRISTART DHA VITAFOL-OB VITAFOL-ONE WESTGEL DHA WESTGEL DHA	 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

PROTON PUMP INHIBITORS

PROTON PUMP INHIBITORS Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
DEXILANT (dexlansoprazole) omeprazole (generic Prilosec) RX pantoprazole (generic Protonix) ^{QL} PROTONIX SUSP (pantoprazole)	dexlansoprazole (generic Dexilant) esomeprazole magnesium (generic Nexium) RX ^{QL} esomeprazole magnesium (generic Nexium) OTC ^{QL} esomeprazole strontium lansoprazole (generic Prevacid) ^{QL} NEXIUM SUSP (esomeprazole) omeprazole/sodium bicarbonate (generic Zegerid RX) pantoprazole GRANULES ^{QL} rabeprazole (generic Aciphex)	 Non-preferred agents will be approved for patients who have failed an 8-week trial of both preferred omeprazole Rx AND pantoprazole OR Protonix SUSP. Pediatric Patients: Patients ≤ 4 years of age – No PA required for Prevacid 30mg or omeprazole 20mg capsules (used to compound suspensions). Drug-specific criteria: Prilosec®OTC/Omeprazole OTC:

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
	CORLANOR SOLN , TAB (ivabradine)	 Diagnosis of Chronic Heart Failure with left ventricular ejection fraction less than or equal to 35%, AND Sinus rhythm with resting heart rate greater than or equal to 70 beats per minute, AND On maximally tolerated doses of betablockers OR have a contraindication to beta-blocker use

SKELETAL MUSCLE RELAXANTS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
paclofen (generic Lioresal) chlorzoxazone (generic Parafon Forte) cyclobenzaprine (generic Flexeril) ^{QL} methocarbamol (generic Robaxin) izanidine TAB (generic Zanaflex)	baclofen (generic for Ozobax) NR,QL SOLN carisoprodol (generic Soma) CL,QL carisoprodol compound cyclobenzaprine ER (generic Amrix) CL dantrolene (generic Dantrium) FEXMID (cyclobenzaprine ER) FLEQSUVY (baclofen) SUSP LORZONE (chlorzoxazone) CL LYVISPAH (baclofen) NR,QL GRANULES metaxalone (generic Skelaxin) NORGESIC FORTE (orphenadrine/ASA/caffeine) orphenadrine ER PARAFON FORTE (chlorzoxazone) tizanidine CAPS ZANAFLEX (tizanidine) CAPS, TAB	 Non-preferred agents will be approved for patients who have failed a 1-week trial of TWO preferred agents within this drug class Drug-specific criteria: cyclobenzaprine ER: Requires clinical reason why IR cannot be used Approved only for acute muscle spasms NOT approved for chronic use carisoprodol: Approved for Acute, musculoskeletal pain - NOT for chronic pain Use is limited to no more than 30 days Additional authorizations will not be granted for at least 6 months following the last dayy of previous course of therapy Dantrolene: Trial NOT required for treatment of spasticity from spinal cordinjury Lorzone®: Requires clinical reason why chlorzoxazone cannot be used Soma® 250mg: Requires clinical reason why 350mg generic strength cannot be used Zanaflex® Capsules: Requires clinical reason generic cannot be used

TETRACYCLINES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
doxycycline hyclate IR (generic Vibramycin) doxycycline monohydrate 50MG, 100MG CAPS doxycycline monohydrate SUSP, TAB (generic Vibramycin) minocycline HCl CAPS (generic Dynacin/ Minocin/Myrac)	demeclocycline (generic Declomycin) ^{CL} DORYX MPC DR (doxycycline pelletized) doxycycline hyclate DR (generic Doryx) doxycycline monohydrate 40MG, 75MG and 150MG CAP (generic Adoxa/Monodox/ Oracea) minocycline HCI TAB (generic Dynacin/Myrac) minocycline HCI ER (generic Solodyn) NUZYRA (omadacycline) tetracycline VIBRAMYCIN SUSP (doxycycline) XIMINO (minocycline ER) ^{QL}	 Non-preferred agents will be approved for patients who have failed a 3-day trial of TWO preferred agents within this drug class Drug-specific criteria: Demeclocycline: Approved for diagnosis of SIADH doxycycline suspension: May be approved with documented swallowing difficulty

THYROID HORMONES

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
levothyroxine TAB (generic Synthroid) liothyronine TAB (generic Cytomel) thyroid, pork TAB UNITHROID (levothyroxine)	ERMEZA (levothyroxine) EUTHYROX (levothyroxine) LEVO-T (levothyroxine) levothyroxine CAPS (generic Tirosint) THYQUIDITY (levothyroxine) SOLN TIROSINT CAPS (levothyroxine) TIROSINT-SOL LIQUID (levothyroxine)	 Non-preferred agents will be approved for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: Tirosint-Sol: May be approved with documented swallowing difficulty

ULCERATIVE COLITIS

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
APRISO (mesalamine) Sulfasalazine IR, DR (generic Azulfidine) LIALDA (mesalamine)	balsalazide (generic Colazal) budesonide DR (generic Uceris) DIPENTUM (olsalazine) mesalamine ER (generic Apriso)	 Non-preferred agents will be approve for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria:
	mesalamine ER (generic Pentasa) ^{NR} mesalamine (generic Asacol HD/ Delzicol/Lialda) PENTASA (mesalamine)	Asacol HD®/Delzicol DR®/ Pentasa® Requires clinical reason why preferred mesalamine products cannot be used
RE	CTAL	
Sulfite-Free ROWASA (mesalamine) mesalamine SUPPOSITORY (generic Canasa)	CANASA (mesalamine) mesalamine ENEMA (generic Rowasa) ROWASA (mesalamine) UCERIS (budesonide)	

UTERINE DISORDER TREATMENT

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
MYFEMBREE (relugolix/ estradiol/ norethindrone acetate) ^{AL, CL,QL} ORIAHNN (elagolix/ estradiol/ norethindrone) ^{AL,CL} ORILISSA (elagolix sodium) ^{QL,CL}		Drug-specific criteria: • Myfembree, Orilissa, and Oriahnn: Requires an FDA approved indication, must follow FDA dosing guidelines, and have had a trial and failure of an NSAID and oral contraceptive • Total duration of treatment is max of 24 months

Preferred Agents	Non-Preferred Agents	Prior Authorization/Class Criteria
BIDIL (isosorbide dinitrate/ hydralazine) ^{CL} cosorbide dinitrate TAB cosorbide dinitrate ER, SA TAB (generic Dilatrate-SR/Isordil) cosorbide mono IR/SR TAB itroglycerin SUBLINGUAL, TRANSDERMAL itroglycerin ER TAB	GONITRO (nitroglycerin) isosorbide dinitrate TAB (Oceanside Pharm MFR only) isosorbide dinitrate/hydralazine (Bidil) ^{CL,NR} NITRO-BID OINT (nitroglycerin) NITRO-DUR (nitroglycerin) nitroglycerin TRANSLINGUAL (generic Nitrolingual) VERQUVO (vericiguat) ^{AL,CL,QL}	 Non-preferred agents will be approve for patients who have failed a trial of ONE preferred agent within this drug class Drug-specific criteria: BiDil: Approved for the treatment of heart failure as an adjunct therapy to standard therapy in self-identified bla patients Verquvo: Approved for use in patient following a recent hospitalization for hwithin the past 6 months OR need for outpatient IV diuretics, in adults with symptomatic chronic HF and EF less than 45%

7. Adjournment / Old Business

- a. No old business topics were discussed by the committee.
- b. A vote to conclude the meeting was made at 2:40 PM CST.

(1st) Motion: Baker (2nd) Motion: Cowles

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, November 15th, 2023

Time:

9:00 AM - 5:00 PM CST

Location:

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

Recorded by: Elanah Figueroa, B.A. – Account Operations Executive Magellan Rx Management, Magellan Medicaid Administration, LLC.