State and Public School Life and Health Insurance Board Drug Utilization and Evaluation Committee Minutes July 19, 2021

The State and Public Life and Health Insurance Board, Drug Utilization and Evaluation Committee (DUEC) met on Monday, July 19th, 2021 at 1:00 p.m.

Voting Members present:

Dr. Scott Pace, Vice-Chairman

Dr. Hank Simmons, Chairman

Dr. Keith McCain -- teleconference

Dr. John Kirtley

Laura Mayfield - teleconference

Dr. Shane David - teleconference

Non-Voting Members present:

Shalada Toles, EBD Deputy Director

Dr. Dwight Davis

Dr. Micah Bard

Dr. Jill Johnson

Voting Members absent:

Non-Voting Members absent:

Dr. Sidney Keisner

OTHERS PRESENT

Rhoda Classen, Mary Massirer, Shay Burleson, Lauren Ballard, Laura Thompson, EBD; Frances Bauman, Nova Nordisk; Aaron Shaw, Jessica Akins, Takisha Sanders, HA; Sherry Bryant, EBRx/EBD; Octavia DeYoung, EBRx; Elizabeth Montgomery, ACHI; Brent Flaherty, Judith Paslaski, Suzanne Woodall, MedImpact; James Chapman, Abbvie; Nima Nabavi, Amgen; Trisha Grantham, AstraZeneca; Ronda Walthall, ARDOT; Mitch Rouse, Brooke Hollowoa, TSS, Autumn Sanson, AR Treasury, Debbie Rogers, DFA, Brent Parker, MERCK; Lance Johnson, Bi-Partisan Strategies; Lori Bowen, BLR; Stephen Carroll, AllCare Pharmacy; Kristie Banks, Mainstream; Robyn Keene, AAEA; Matt Kersting, Segal; Melissa Rifle, AGFC; Doug Wood, Viiv Healthcare

CALL TO ORDER

Meeting was called to order by Dr. Hank Simmons, Chair, and he announced that we do have a quorum today.

APPROVAL OF MINUTES

The request was made by Dr. Simmons to approve the March 8th, 2021 minutes. Dr. Kirtley made the motion to approve. Dr. Pace seconded; all were in favor.

Motion Approved.

I. New Business

A. New Drugs: by Dr. Jill Johnson and Dr. Sidney Keisner, UAMS

<u>Brand</u>	<u>Generic</u>	<u>Recommendation</u>		
Non-Specialty Drugs				
(1) VERQUVO	VERICIGUAT	Exclude, Code 8		
(2) ELEPSIA XR	LEVETIRACETAM	Exclude, Code 13		
(3) ROSZET	EZETIMIBE/ROSUVASTATIN CALCIUM	Exclude, Code 13		
(4) QELBREE	VILOXAZINE HCL	Exclude, Code 13		

(5) ZEGALOGUE AUTOINJECTOR	DASIGLUCAGON HCL	Exclude, Code 13
(6) NEXTSTELLIS	DROSPIRENONE/ESTETROL	Exclude, Code 13
(7) ACCRUFER	FERRIC MALTOL	Exclude, Code 13
	Specialty Drugs	
(1) CABENUVA	CABOTEGRAVIR/RILPIVIRINE	Exclude, Code 4, 13
(2) VOCABRIA	CABOTEGRAVIR	Exclude, Code 12, 13
(3) LUPKYNIS	VOCLOSPORIN	Exclude, Code 13
(4) TEPMETKO	TEPOTINIB HCL	Exclude, Code 1,13
(5) UKONIQ	UMBRALISIB TOSYLATE	Exclude, Code 1,13
(6) EVKEEZA	EVINACUMAB-DGNB	Exclude, Code 1,13
(7) COSELA	TRILACICLIB DIHYDROCHLORIDE	Cover w/PA
(8) AMONDYS-45	CASIMERSEN	Exclude
(9) PEPAXTO	MELPHALAN FLUFENAMIDE HCL	Exclude, Code 1,13
(10) MARGENZA	MARGETUXIMAB-CMKB	Exclude, code 8, 13
(11) FOTIVDA	TIVOZANIB HCL	Exclude, code 13
(12) PONVORY	PONESIMOD	Exclude, code 13
(13) JEMPERLI	DOSTARLIMAB-GXLY	Exclude, code 1
(14) ZYNLONTA	LONCASTUXIMAB TESIRINE-LPYL	Exclude, code 13

Dr. Pace made a motion to approve the non-specialty drug recommendations as presented. Dr. Kirtley seconded. All were in favor.

Motion Approved.

Dr. Simmons made a motion to approve the specialty drug recommendations as presented. Dr. Kirtley seconded. All were in favor.

Motion Approved.

II. Old Business

A. Second Review of Drugs: Dr. Jill Johnson, UAMS

<u>Brand</u>	<u>Generic</u>	<u>Recommendation</u>
(1) VUMERITY	DIROXIMEL FUMARATE	Exclude, Code 13
(2) DARZALEX FASPRO	DARATUMUMAB AND HYALURONIDASE-FIHJ	Cover w/PA
(3) ICLUSIG	PONATINIB	Cover w/PA
(4) REVCOVI®	ELAPEGADEMASE-LVLR	Cover w/PA, specialty
(5) BENLYSTA	BELIMUMAB	Cover w/PA for only lupus nephritis
(6) SGLT2 INHIBITORS		Remove PA from covered SGLT2; Rebid the class of SGLT2's

(7) PDE5 ENZYME INHIBITORS	SILDENAFIL (REVATIO)	Cover PO susp w/brand penalty (no PA or QL) - Age edit (<7y only)
	TADALAFIL	QL (6/30) on all strengths except 20 mg.
(8) XPOVIO	SELINEXOR	Cover w/PA (Selinexor/dex indication only)
(9) BLENREP	BELANTAMAB MAFODOTIN-BLMF	Covered w/PA (medical drug); revisit 1 year
(10) PADCEV	ENFORTUMAB VEDOTIN-EJFV	N/A Medical; Revisit 1 year
(11) KRYSTEXXA	PEGLOTICASE	Exclude from pharmacy and medical
(12) SIKLOS	HYDEOXYUREA	Exclude, Code 13
(13) BLINCYTO	BLINATUMOMAB	Cover w/PA for relapsed/refractory indication only (medical drug)

Dr. Kirtley made a motion to approve the recommendations as presented and add the opportunity to go out to bid for the SGLT2 class. Dr. Pace seconded. All were in favor.

Motion Approved.

- b. Ticagrelor (Brilinta): 60, 90mg Tablets
 - i. EBRx Decision 4/22/21 regarding EBD plans: Exclude Ticagrelor, code 13 (alternative is clopidogrel)

Dr. Kirtley made a motion to accept the recommendation to exclude Ticagrelor (Brilinta) as presented. Dr. Pace seconded. All were in favor.

Motion Approved.

Dr. Kirtley made a motion to adjourn the meeting. Dr. Pace seconded. All were in favor.

Meeting Adjourned.

*New Drug Code Key:

Lacks meaningful clinical endpoint data; has shown efficacy for surrogate endpoints only. 2 Drug's best support is from single arm trial data 3 No information in recognized information sources (PubMed or Drug Facts & Comparisons or Lexicomp) Convenience Kit Policy - As new drugs are released to the market through Medispan, those drugs described as "kits will not be considered for inclusion in the plan and will therefore be excluded products unless the product is available solely as a kit. Kits typically contain, in addition to a pre-packaged quantity of the featured drug(s), items that may be associated with the administration of the drug (rubber gloves, sponges, etc.) and/or additional convenience items (lotion, skin cleanser, etc.). In most cases, the cost of the "kit" is greater than the individual items purchased separately. Medical Food Policy - Medical foods will be excluded from the plan unless two sources of peer-reviewed, published medical literature supports the use in reducing a medically necessary clinical endpoint. A medical food is defined below: A medical food, as defined in section 5(b)(3) of the Orphan Drug Act (21 U.S.C. 360ee(b)(3)), is "a food which is formulated to be consumed or administered eternally under the supervision of a physician and which is intended for the specific dietary management of a disease or condition for which distinctive nutritional requirements, based on recognized scientific principles, are established by medical evaluation." FDA considers the statutory definition of medical foods to narrowly constrain the types of products that fit within this category of food. Medical foods are distinguished from the broader category of foods for special dietary use and from foods that make health claims by the requirement that medical foods be intended to meet distinctive nutritional requirements of a disease or condition, used under medical supervision, and intended for the specific dietary management of a disease or condition. Medical foods are not those simply recommended by a physician as part of an overall diet to manage the symptoms or reduce the risk of a disease or condition, and all foods fed to sick patients are not medical foods. Instead, medical foods are foods that are specially formulated and processed (as opposed to a naturally occurring foodstuff used in a natural state) for a patient who is seriously ill or who requires use of the product as a major component of a disease or condition's specific dietary management. Cough & Cold Policy - As new cough and cold products enter the market, they are often simply re-formulations or new combinations of existing products already in the marketplace. Many of these existing products are available in generic form and are relatively inexpensive. The new cough and cold products are branded products and are generally considerably more expensive than existing products. The policy of the ASE/PSE prescription drug program will be to default all new cough and cold products to "excluded" unless the DUEC determines the product offers a distinct advantage over existing products. If so determined, the product will be reviewed at the next regularly scheduled DUEC meeting. Multivitamin Policy - As new vitamin products enter the market, they are often simply re-formulations or new combinations of vitamins/multivitamins in similar amounts already in the marketplace. Many of these existing products are available in generic form and are relatively inexpensive. The new vitamins are branded products and are generally considerably more expensive than existing products. The policy of the ASE/PSE prescription drug program will be to default all new vitamin/multivitamin products to "excluded" unless the DUEC determines the product offers a distinct advantage over existing products. If so determined, the product will be reviewed at the next regularly scheduled DUEC meeting. 7 Drug has limited medical benefit &/or lack of overall survival data or has overall survival data showing 8 minimal benefit Not medically necessary Peer -reviewed, published cost effectiveness studies support the drug lacks value to the plan. 10 Oral Contraceptives Policy - OCs which are new to the market may be covered by the plan with a zero dollar, tier 1, 2, or 3 copay, or may be excluded. If a new-to-market OC provides an alternative product not similarly achieved by other OCs currently covered by the plan, the DUEC will consider it as a new drug. IF the drug does not offer a novel alternative or offers only the advantage of convenience, it may not be considered for inclusion in the plan. Other 12 Insufficient clinical benefit OR alternative agent(s) available