Division of Medical Services
Pharmacy Program

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ARKANSAS MEDICAID DUR BOARD
PHYSICIAN ADMINISTERED DRUGS (PAD) SUBCOMMITTEE

June 10, 2026 12:00 pm – 1:00 pm CST

VIRTUAL TEAMS MEETING LINK

Join the meeting now
Meeting ID: 233 781 903 264 22
Passcode: 3KF2cp7j

Dial in by phone
+1 501-244-3310,,979808470# United States, Little Rock
Phone conference ID: 979 808 470#

****TENTATIVE AGENDA IS SUBJECT TO CHANGE****

I. OUTSIDE SPEAKERS

Outside speakers with clinical or scientific credentials or patient experience pertinent to a product or topic that is posted on the upcoming PAD Subcommittee meeting agenda may request to speak on that product or topic. Public comments will not be permitted for topics not included on the agenda. Requests to speak at the PAD Subcommittee meeting must be made in writing to the DUR Chairperson (cinnamon.pearson@dhs.arkansas.gov) at least two (2) weeks before the PAD Subcommittee meeting date, and should include:

(1) The speaker's name, title, relevant credentials, and organization;
(2) Contact information for the speaker including address, telephone number, and email;
(3) The agenda item(s) which the speaker intends to address;
(4) Prepared comments which are not a reiteration of the FDA approved package insert; and
(5) Any educational materials for Board members in advance of the meeting.

This information shall be included in information sent to Subcommittee members two (2) weeks prior to the meeting. Presentations or public comments given at the PAD Subcommittee meeting are limited to a total of **three (3) minutes** per drug, which may be shared by multiple speakers. This time limit does not include responses to any questions raised by DUR Board Subcommittee members during the course of the meeting.

II. UNFINISHED BUSINESS

A. Vote on minutes from March 11, 2026 DUR PAD Subcommittee meeting
B. Implementation information from April 15, 2026 DUR Board meeting

III. NEW BUSINESS

A. GENERAL MONOGRAPH UPDATES NOT REQUIRING REVIEW

1. Expanded indications concerning age
2. Updated or added J-codes (HCPCS code)
3. Biosimilars covered as medical benefit will be added to the current reference product monograph

B. UPDATE CURRENT MONOGRAPHS

1. ENHERTU (fam-trastuzumab deruxtecan-nxki) injection
2. OPDIVO (nivolumab) injection
3. DARZALEX FASPRO (daratumumab and hyaluronidase-fihj) injection
4. PROLIA AND XGEVA (denosumab) injection
5. ENTYVIO (vedolizumab) injection
6. LEQVIO (inclisiran) injection
7. TECVAYLI (teclistamab-cqyv) injection
8. VABYSMO (faricimab-svoa) injection

C. NEW MONOGRAPHS

1. EXDENSUR (depemokimab-ulaa) injection
2. XIPERE (triamcinolone acetonide injectable suspension)
3. ILARIS (canakinumab) injection