

Division of Medical Services Pharmacy Program



P.O. Box 1437, Slot S415 · Little Rock, AR 72203-1437
501-683-4120 · Fax: 501-683-4124 or 1-800-424-5851



ARKANSAS MEDICAID DUR BOARD PHYSICIAN ADMINISTERED DRUGS (PAD) SUBCOMMITTEE

December 9, 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 September 9, 2026 DUR PAD Subcommittee meeting
- B. Implementation information from October 21, 2026 DUR Board meeting

III. NEW BUSINESS

A. GENERAL MONOGRAPH UPDATES NOT REQUIRING REVIEW

1. ENZEEVU (aflibercept abzv)—Biosimilar added to EYLEA monograph
2. IMULDOSA (ustekinumab-srlf)—Biosimilar added to STELARA monograph
3. STEQEYMA (ustekinumab-stba)—Biosimilar added to STELARA monograph

B. UPDATE CURRENT MONOGRAPHS

1. GAZYVA (obinutuzumab) injection
2. TECENTRIQ (atezolizumab) injection
3. BRIUMVI (ublituximab) injection
4. PADCEV (enfortumab vedotin) injection
5. UPLIZNA (inebilizumab) injection
6. SARCLISA (isatuximab) and SARCLISA ESCENA (isatuximab-irfc) injection
7. SKYRIZI (risankizumab-rzaa) injection
8. TREMFYA (guselkumab) injection
9. CIMZIA (certolizumab pegol) injection
10. OMVOH (mirikizumab-mrkz) injection
11. PLUVICTO (lutetium lu 177 vipivotide tetraxetan) injection
12. ULTOMIRIS (ravulizumab) solution

C. NEW MONOGRAPHS

1. ISEMBYLD (apitegromab-mstn) injection
2. IDELVION (coagulation factor ix recombinant)
3. CASGEVY (exagamglogene autotemcel) injection