

2025 PREFERRED DRUG LIST MEETING SCHEDULE

State of Montana
Department of Public Health & Human
Services

Montana Medicaid Drug Use Review (DUR) Board/Formulary Committee Meeting

The State of Montana Medicaid DUR Board/Formulary Committee will hold a meeting on:

Date: April 16, 2025 (Wednesday)

Time: 1:00 pm – 5:00 pm Mountain Time

Location: This meeting will be conducted by teleconference only. Register in advance for this meeting. After registering, you will receive a confirmation email containing information about joining the meeting. Those wishing to provide public comment must follow the additional instructions provided on page 2 of this agenda. Please note that ALL public comment, including manufacturer presentations, will be in the same public comment period at the beginning of the meeting
[https://events.teams.microsoft.com/event/2a554955-0200-49d0-a813-ef3759d55573@1f053f7a-e47d-43fd-9182-8507c9ff10c7\[events.teams.microsoft.com\]](https://events.teams.microsoft.com/event/2a554955-0200-49d0-a813-ef3759d55573@1f053f7a-e47d-43fd-9182-8507c9ff10c7[events.teams.microsoft.com])

At this time the Montana Medicaid DUR Board/Formulary Committee will review the following drug classes for Preferred Drug List (PDL) review:

All drugs reviewed pertain to oral drugs unless otherwise indicated

NI- New information, ND- New Drug, NG-New Generic

The Department will review **GROUP 3** as **NEW** information is known to exist:

- ANTI-ALLERGENS, ORAL – NI - Palforzia
- ANTIVIRALS, ORAL – ND – Paxlovid, NI-Oseltamivir
- BLADDER RELAXANTS – NI - Gemtesa
- COPD AGENTS – ND - Ohtuvayre
- CYTOKINES & CAM ANTAGONISTS – NI - Bimzelx, Kevzara, Omvoh, Otezla, Rinvoq, Skyrizi, Spevigo, Tremfya, biosimilars adalimumab, ustekinumab, tocilizumab
- EPINEPHRINE, SELF-INJECTED – ND - Neffy
- IMMUNOMODULATORS, ASTHMA – NI - Xolair, Fasenra
- IMMUNOMODULATORS, ATOPIC DERMATITIS – ND - Ebglyss, Nemluvio, NI - Adbry, Dupixent, Zoryve
- METHOTREXATE AGENTS – NI - Jylamvo

The Department will validate **GROUP 2** Formulary Committee's clinical recommendations unless manufacturers submit **NEW relevant clinical** information prior to the deadline noted below.

• ACNE AGENTS	• FLUOROQUINOLONES, ORAL
• ANTIBIOTICS, GI	• GLUCOCORTICOIDS, INHALED
• ANTIBIOTICS, INHALED	• GLUCOCORTICOIDS, ORAL
• ANTIBIOTICS, TOPICAL	• HEPATITIS C AGENTS
• ANTIBIOTICS, VAGINAL	• IDIOPATHIC PULMONARY FIBROSIS
• ANTIFUNGALS, ORAL	• IMMUNOMODULATORS, TOPICAL
• ANTIFUNGALS, TOPICAL	• IMMUNOSUPPRESSANTS
• ANTIHISTAMINES- MIN SEDATING	• INTRANASAL RHINITIS AGENTS
• ANTINEOPLASTIC AGENTS, TOPICAL	• LEUKOTRIENE MODIFIERS
• ANTIPARASITICS, TOPICAL	• MACROLIDES & KETOLIDES
• ANTIPSORIATICS, TOPICAL	• ROSACEA AGENTS, TOPICAL
• ANTIVIRALS, TOPICAL	• SMOKING CESSATION
• BPH AGENTS	• STEROIDS, TOPICAL
• BRONCHODILATORS, BETA AGONISTS	• TETRACYCLINES
• CEPHALOSPORINS & RELATED AGENTS	

There are no **GROUP 1** agents

Montana Medicaid
Department of Public Health and Human Services
DUR Board/Formulary Committee Meeting
General Procedures for Public Comment

Public Comment will be permitted on items over which the DUR Board has jurisdiction. The public comment period will be limited to 15 minutes. All persons wishing to speak through the video conferencing platform must notify the Department by submitting their name and topic they wish to speak on via <https://forms.office.com/g/EkPyKbJYTv> by **noon** the day prior to the meeting. Individual comment will be limited to a maximum of 3 minutes per person but may be subject to further limitations depending on the number of speakers (15 minutes divided by total speakers).

Due to time constraints, all persons are encouraged to provide written comment, especially those providing clinical information, to allow the board adequate time to review and research. Please email written comments and/or clinical information, with a statement indicating that it is public comment, to PDL@mt.gov at least 7 days prior to the meeting. Please limit clinical information to 2 pages and to new information since last review. New peer-reviewed randomized comparative controlled trials or randomized controlled trials with true health outcomes are most helpful. Please do not provide any pricing information. Materials submitted less than 7 days prior to the meeting, in excess of 2 pages, and/or including pricing information will not be included in the board members' meeting packets.

Note: These procedures may be revised at the discretion of the Department.