



## Oregon Prescription Drug Affordability Board

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# Agenda

*This is a draft agenda and subject to change*

**Wednesday, January 21, 2026 – 8:00 a.m.**

Register for meeting: [Zoom link](#)

Purpose	Subject	Presenter
<i>Informational and vote</i>	Call to order and roll call	Chair Shelley Bailey
<i>Informational</i>	Board declarations of conflict of interest and meetings with entities or individuals related to board activities	Chair Shelley Bailey
<i>Review</i>	<b><u>Board review of 12/17/2025 minutes</u></b>	Chair Shelley Bailey
<i>Informational</i>	PDAB program update	Sarah Young
<i>Informational</i>	General public comment: limited to 3 minutes per speaker	Chair Shelley Bailey
<i>Discussion and vote</i>	<b><u>2025 drug review</u></b> : Votes to identify up to nine prescription drugs and at least one insulin product that may create affordability challenges for the health care systems or high out-of-pocket costs for patients in Oregon, pursuant to ORS 646A.694	Cortnee Whitlock
<i>Discussion</i>	Recommendations to include in the 2025 drug review report	Cortnee Whitlock
<i>Informational</i>	Executive session for legal advice pursuant to ORS 192.660(2)(f)	Oregon Department of Justice
<i>Discussion and review</i>	<b><u>2026 drug review</u></b> : Discussion and review of the prescription drugs and insulin products preliminary lists	Cortnee Whitlock
<i>Break</i>	The board will take a break around 10:30	Chair Shelley Bailey
<i>Informational</i>	Announcements	Chair Shelley Bailey
<i>Vote</i>	Adjournment	Chair Shelley Bailey

Accessibility: Anyone needing assistance due to a disability or language barrier can contact Melissa Stiles at least 48 hours ahead of the meeting at [pdab@dcbs.oregon.gov](mailto:pdab@dcbs.oregon.gov) or 971-374-3724. American sign language will be available during the Jan. 21 board meeting.



Oregon Prescription Drug  
Affordability Board

Oregon Prescription Drug Affordability Board Regular Meeting  
Wednesday, December 17, 2025  
Draft Minutes

**Web link to the meeting video:** <https://youtu.be/zTyUMuPgpOs>

**Web link to the meeting materials:** <https://dfr.oregon.gov/pdab/Documents/20251217-PDAB-document-package.pdf>

**Call to order:** Chair Shelley Bailey called the meeting to order at 8:03 a.m.

**Roll call:**

Present: Chair Shelley Bailey, Vice Chair Dan Hartung, Lauri Hoagland, Dan Kennedy, Michele Koder, and John Murray

Arrived late: Chris Laman (8:21 a.m.)

Absent: None

**Board declarations of conflict of interest and meetings with entities or individuals related to board activities:** John Murray provided a statement. View at video minute [00:01:02](#).

**Approval of board minutes:** Chair Bailey approved by consensus the Nov. 19, 2025 minutes as shown on [Pages 3-4](#) of the agenda materials. View at video minute [00:02:08](#).

**PDAB program update:** Sarah Young, PDAB executive director, provided a program update. View the video at minute [00:02:28](#).

**General public comment:** Public comment was received from the following individuals who signed up in advance: Dharia McGrew, PhRMA; Jessica McBride, Oregon Coalition for Affordable Prescriptions; Ranier Simons, Community Access National Network; and Lorren Sandt, Caring Ambassadors. The board received 7 written comments, which are posted on the [PDAB website](#). View at video minute [00:09:23](#).

**Board discussion and vote on annual report:** Board members discussed the 2025 annual report and proposed amendments. The board voted unanimously to approve the amended 2025 annual report and submit it to the Oregon Legislature. The approved [annual report](#) is posted on the PDAB website. View the discussion and vote at video minute [00:19:26](#).



**MOTION to approve the 2025 annual report as amended by the board today and deliver to the Oregon Legislature by Dec. 31, 2025, as required by ORS 646A.696.**

**Board Vote:**

Motion: Chris Laman

Second: John Murray

Yes: Lauri Hoagland, Dan Kennedy, Michele Koder, Chris Laman, John Murray, Vice Chair Dan Hartung, Chair Shelley Bailey

No: None

Absent: None

**MOTION PASSED 7-0**

**2026 drug review – presentation of the prescription drugs and insulin products preliminary lists:** The board received the draft preliminary lists of prescription drugs and insulin products for the 2026 drug reviews, based on commercial insurer reporting from calendar year 2024. The board began reviewing the list at the Dec. 17 meeting and will continue reviewing and narrowing the lists for further evaluation in 2026. The [preliminary list](#) of drugs is posted on the PDAB website. View at video minute [00:44:57](#).

**2025 drug review – continued review of the prescription drugs and insulin products:** The board continued its review of the prescription drug and insulin products. The board reviewed 23 prescription drugs and insulin products during its meetings in July, August, September, and October, with additional discussions in November and December. The board will vote in January on a list of up to nine prescription drugs and at least one insulin product for inclusion in a report to the Oregon Legislature in March. Drug reviews are conducted according to ORS 646A.694 and OAR 925-200-0020. Read the detailed [reports for each drug](#) posted on the PDAB website. View at video minute [01:39:25](#).

**Announcements:** Chair Bailey announced the next board meeting will be Jan. 21, 2026 at 8 a.m. View at video minute [03:31:43](#).

**Adjournment:** Chair Bailey adjourned the meeting at 12 p.m. with all board members in agreement. View at video minute [03:31:57](#).



## Web links to final reports for the 2025 drug review

**Agenda item:** 2025 drug review: Votes to identify up to nine prescription drugs and at least one insulin product that may create affordability challenges for the health care systems or high out-of-pocket costs for patients in Oregon, pursuant to ORS 646A.694.

In Table 1 below, click on the prescription drug name to read the final version of the drug review document. This table of information is also available on the [Prescription Drug Affordability Board drug review page](#).

**Table 1: 2025 Oregon PDAB drug review documents, videos, and minutes web links**

Drug review documents	Review date	Meeting video	Meeting minutes
<ul style="list-style-type: none"><li>• <a href="#">Vraylar</a></li><li>• <a href="#">Entresto</a></li><li>• <a href="#">Ajoyv</a></li><li>• <a href="#">Emgality</a></li><li>• <a href="#">Nurtec</a></li><li>• <a href="#">Ubrelvy</a></li></ul>	July 16, 2025	<a href="#">July 16, 2025</a>	<a href="#">July 16, 2025</a>
<ul style="list-style-type: none"><li>• <a href="#">Trelegy</a></li><li>• <a href="#">Eliquis</a></li><li>• <a href="#">Xarelto</a></li><li>• <a href="#">Cosentyx</a></li><li>• <a href="#">Creon</a></li></ul>	Aug. 20, 2025	<a href="#">Aug. 20, 2025</a>	<a href="#">Aug. 20, 2025</a>
<ul style="list-style-type: none"><li>• <a href="#">Jardiance</a></li><li>• <a href="#">Mounjaro</a></li><li>• <a href="#">Ozempic</a></li><li>• <a href="#">Rybelsus</a></li><li>• <a href="#">Trulicity</a></li></ul>	Sept. 17, 2025	<a href="#">Sept. 17, 2025</a>	<a href="#">Sept. 17, 2025</a>
<ul style="list-style-type: none"><li>• <a href="#">Insulin products:</a> Basaglar KwikPen, Insulin Glargine-yfgn, Lantus, Lantus SoloStar, Semglee, Toujeo Max SoloStar Toujeo SoloStar</li></ul>	Oct. 15, 2025	<a href="#">Oct. 15, 2025</a>	<a href="#">Oct. 15, 2025</a>

PDAB scoring rubric for 2025 prescription drug cost reviews.  
 Scoring rubric is a decision support tool and not the  
 determinative of drug review outcomes.

	Review Drug	Vraylar	Entresto	Ajovy	Emgality	Nurtec	Ubrelvy
<b>Rubris Criteria</b>							
Utilization		Severe	Severe	High	High	High	High
Price evaluation		Moderate	Severe	High	High	High	Severe
Price concessions		Moderate	Low	High	Low	Moderate	Moderate
System and payer costs		High	High	Low	Low	High	Moderate
Enrollee burden		Severe	Moderate	Moderate	High	Moderate	Moderate
Equity impact		Yes	Yes	Yes	Yes	Yes	Yes
Access restrictions		Yes	No	Yes	Yes	Yes	Yes
Less expensive therapeutic alternatives available		Yes	Yes	Yes	Yes	Yes	Yes
Stakeholder input identify		Yes	No	Yes	Yes	Yes	Yes
Patent expirations more than 18 months		No	<b>No</b>	Yes	Yes	No	Yes
Excluded from MFP		No	No	Yes	Yes	Yes	Yes
Scoring totals		14	11	13	12	13	14

\***Bolded words indicate a change in scoring**

\*score update  
12/23/25

	Review Drug	Trelegy	Eliquis	Xarelto	Cosentyx	Creon
<b>Rubris Criteria</b>						
Utilization		Severe	Severe	Severe	High	High
Price evaluation		Moderate	Severe	High	Severe	Severe
Price concessions		High	Low	High	High	Severe
System and payer costs		High	Severe	Severe	Severe	High
Enrollee burden		Moderate	Moderate	Moderate	Severe	Moderate
Equity impact		Yes	Yes	Yes	Yes	Yes
Access restrictions		No	No	No	Yes	no
Less expensive therapeutic alternatives available		Yes	Yes	Yes	Yes	Yes
Stakeholder input identify		No	Yes	Yes	Yes	Yes
Patent expirations more than 18 months		Yes	Yes	Yes	Yes	Yes
Excluded from MFP		No	No	No	Yes	Yes
Scoring totals		12	14	15	19	16

\***Bolded words indicate a change in scoring**

	Review Drug	Jardiance	Mounjaro	Ozempic	Rybelsus	Trulicity
<b>Rubris Criteria</b>						
Utilization		Severe	High	Severe	High	Severe
Price evaluation		High	High	High	Moderate	Severe
Price concessions		Moderate	High	high	Moderate	High
System and payer costs		Severe	High	Severe	High	Severe
Enrollee burden		High	Moderate	Moderate	Moderate	Moderate
Equity impact		Yes	Yes	Yes	Yes	Yes
Access restrictions		no	Yes	Yes	Yes	Yes
Less expensive therapeutic alternatives available		Yes	Yes	Yes	Yes	Yes
Stakeholder input identify		Yes	Yes	Yes	No	Yes
Patent expirations more than 18 months		No	Yes	Yes	Yes	Yes
Excluded from MFP		No	<b>Yes</b>	No	No	Yes

Scoring totals

\***Bolded words indicate a change in scoring**

14	15	16	11	18
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\*score update  
12/23/25



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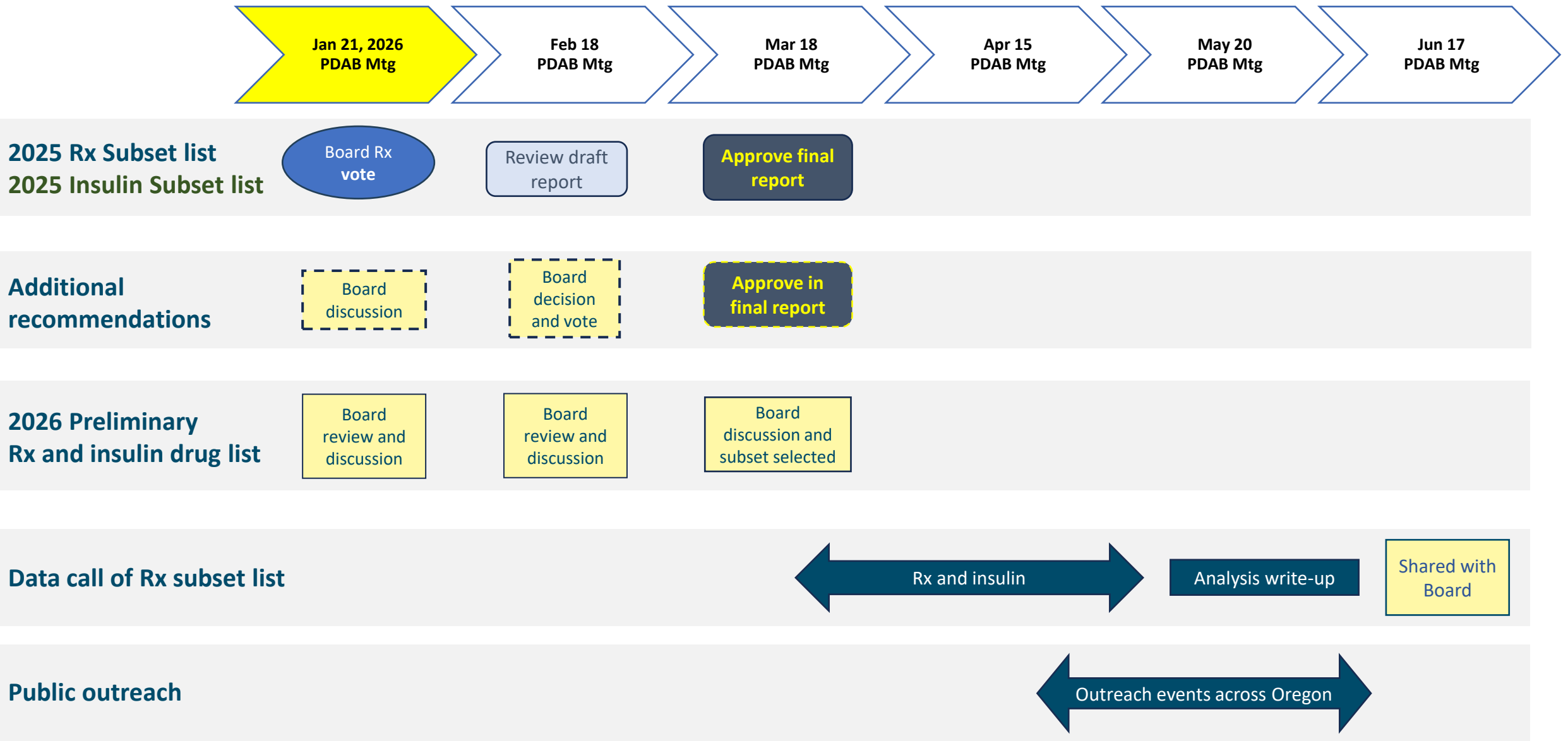


# Prescription Drug Affordability Board

## 2026 drug review roadmap

Jan. 21, 2026

# Draft 2025-2026 drug review (DR) & annual report calendar





Oregon Prescription Drug  
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# Oregon PDAB authority to review drugs with orphan-drug designations

Cortnee Whitlock, senior policy analyst

January 21, 2026

# Purpose of briefing

- Clarify how federal orphan-drug designation (rare disease or condition) interacts with Oregon PDAB authority
- Explain what ORS 646A.694 exempts — and what it does *not exempt*
- Show how OAR 925-200-0020(2)(n) guides board decisions
- Provide an interpretation for reviewing drugs with multiple indications



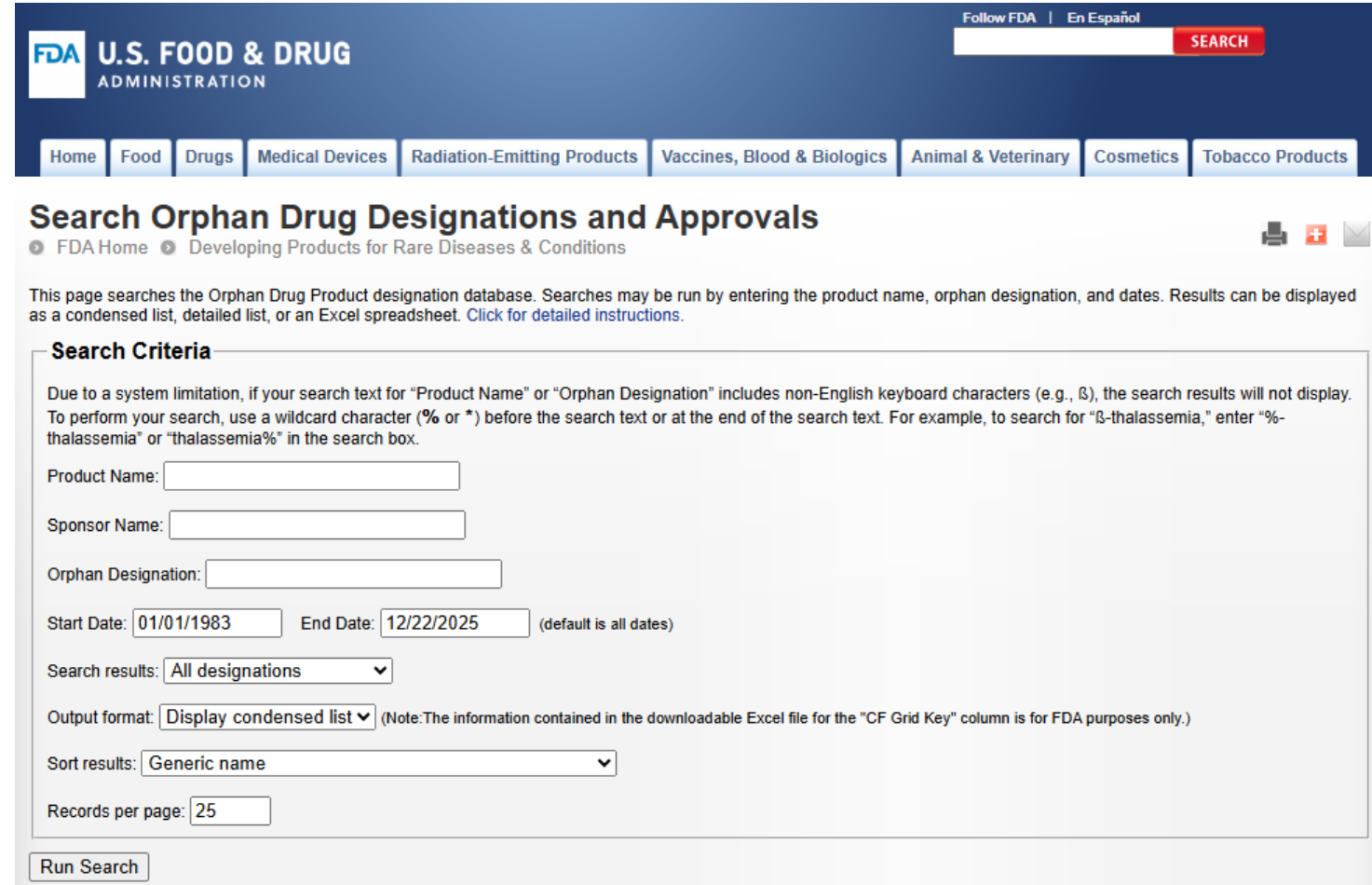
# Federal Law: What 21 U.S.C. § 360bb does

- Defines how a **drug** receives a designation for a rare disease or condition (often called orphan-drug designation)
- Does **not** regulate pricing, affordability, or state review
- Does **not** preempt state PDAB authority



# FDA's orphan-designation program is built on 21 U.S.C. § 360bb and 21 C.F.R. Part 316

- FDA guidance on how the agency grants orphan designation to drugs intended to treat rare diseases or conditions.
- That authority comes from the Orphan Drug Act, which includes **21 U.S.C. § 360bb**.



The screenshot shows the FDA's website for searching orphan drug designations. The header includes the FDA logo and navigation links for various product categories. The main heading is "Search Orphan Drug Designations and Approvals". Below this, there is a brief description of the search function and a "Search Criteria" section. The search criteria section includes input fields for Product Name, Sponsor Name, and Orphan Designation. It also features date pickers for Start Date (01/01/1983) and End Date (12/22/2025), a dropdown for Search results (All designations), a dropdown for Output format (Display condensed list), a dropdown for Sort results (Generic name), and a text input for Records per page (25). A "Run Search" button is located at the bottom of the form.

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### Search Orphan Drug Designations and Approvals

FDA Home Developing Products for Rare Diseases & Conditions

This page searches the Orphan Drug Product designation database. Searches may be run by entering the product name, orphan designation, and dates. Results can be displayed as a condensed list, detailed list, or an Excel spreadsheet. [Click for detailed instructions.](#)

#### Search Criteria

Due to a system limitation, if your search text for "Product Name" or "Orphan Designation" includes non-English keyboard characters (e.g., ß), the search results will not display. To perform your search, use a wildcard character (%) or (\*) before the search text or at the end of the search text. For example, to search for "ß-thalassemia," enter "%-thalassemia" or "thalassemia%" in the search box.

Product Name:

Sponsor Name:

Orphan Designation:

Start Date:  End Date:  (default is all dates)

Search results:

Output format:  (Note: The information contained in the downloadable Excel file for the "CF Grid Key" column is for FDA purposes only.)

Sort results:

Records per page:

Run Search



# ORS 646A.694: Oregon's statutory drug review framework

ORS 646A.694 (2) states:

“A drug that is designated... under 21 U.S.C. 360bb, as a drug for a rare disease or condition is **not subject to review under subsection (1) of this section.**”

What this means:

- The **orphan indication** is exempt for review
- ORS 646A.694 (2) does **not** exempt the entire drug
- ORS 646A.694 (2) does **not** prohibit review of **non-orphan indications**



## What ORS 646A.694 (2) *does not say*

- That all uses of drug with orphan-designation are exempt from review
- That PDAB cannot review the drug for other indications
- **Important:** The exemption is **orphan indication-specific**, not **drug-wide**.



# Administrative Rule: OAR 925-200-0020(2)(n)

A drug used for other indications, in addition to a rare disease or condition, is not exempt from an affordability review for those other indications.

This rule:

- Interprets ORS 646A.694 (2) consistently with legislative intent
- Confirms PDAB ability to review non-orphan uses
- Prevents shielding high-cost drugs with a single orphan indication



# Legal hierarchy

1. **Federal law 21 U.S.C. § 360bb** → Defines orphan designation; does **not** restrict state affordability review
2. **State statute ORS 646A.694 (2)** → Exempts only the orphan indication
3. **Administrative rule OAR 925-200-0020** → Clarifies that non-orphan indications remain reviewable

**Result:** PDAB may review any drug for its **non-orphan indications**, even if it has an orphan designation.



# Practical application for PDAB

When a drug has:

- **Only an orphan indication:** → Exempt from review
- **Orphan + non-orphan indications:** → PDAB may review the drug for non-orphan uses
- **Off-label use:** → PDAB may consider off-label utilization when assessing affordability



# Why this matters

- Many drugs with orphan designations have large commercial markets
- Oregon's statute and rule prevents inappropriate exemptions
- Ensures PDAB can address affordability challenges affecting Oregon patients and its healthcare system
- Aligns with PDAB mission and best practices



# Summary

- **21 U.S.C. § 360bb** defines how a **drug** receives orphan designation, and **does not** restrict PDAB authority to review drugs that treat other conditions besides their rare disease or condition designation
- **ORS 646A.694 (2)** exempts only the **orphan-drug designation**
- **OAR 925-200-0020(2)(n)** confirms PDAB may review non-orphan indications





## Web links to preliminary drug lists for 2026 drug review

**Agenda item:** 2026 drug review: Discussion and review of the prescription drugs and insulin products preliminary lists.

Below, click on the file name below to view the prescription drugs and insulin products preliminary lists in Excel spreadsheets. The lists are for the 2026 drug review using data information from 2024. This information is also available on the [Prescription Drug Affordability Board drug review page](#).

- **Prescription drugs preliminary list:**  
[https://dfr.oregon.gov/pdab/Documents/2026\\_Drug\\_Review\\_Preliminary\\_List\\_v02.xlsx](https://dfr.oregon.gov/pdab/Documents/2026_Drug_Review_Preliminary_List_v02.xlsx)
- **Insulin products preliminary list:**  
[https://dfr.oregon.gov/pdab/Documents/2026\\_Drug\\_Review\\_Insulin\\_Preliminary\\_List\\_v02.xlsx](https://dfr.oregon.gov/pdab/Documents/2026_Drug_Review_Insulin_Preliminary_List_v02.xlsx)

# DO YOU THINK YOUR PRESCRIPTION DRUGS COST TOO MUCH?

## Learn why drug costs are so high

Oregon Prescription Drug Affordability Board helps to protect Oregonians and the health care system from the high cost of prescription drugs. The board wants to reduce financial burdens for patients.

## Help us identify solutions to high drug costs

Please come share your story at a community forum with board staff about how prescription drug prices and medication costs have affected you.



### In-person forums

**Salem** – Tuesday, May 5, 5 to 6:30 p.m.  
Salem Public Library  
585 Liberty St. SE  
Salem

**Redmond** – Thursday, May 7, 5 to 6:30 p.m.  
Redmond Public Library  
827 SW Deschutes Ave.  
Redmond

**Portland** – Tuesday, May 12, 6 to 7:30 p.m.  
Asian Health Center  
9035 SE Foster Road  
Portland

**Beaverton** – Monday, May 18, 5 to 6:30 p.m.  
Beaverton Public Library  
12375 SW Fifth St.  
Beaverton



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### Online forums

Join any ZoomGov meeting at the scheduled time at [zoomgov.com/join](https://zoomgov.com/join).

Enter the meeting ID and passcode

**Passcode: OregonPDAB**  
**Dial in by phone: 669-254-5252**

**Tuesday, April 28**, 7 to 8:30 p.m.  
Join ZoomGov Meeting  
Meeting ID: 161 635 9753

**Monday, May 11**, noon to 1:30 p.m.  
Join ZoomGov Meeting  
Meeting ID: 160 556 2737

**Wednesday, May 13**, 6:30 to 8:30 p.m.  
Join ZoomGov Meeting  
Meeting ID: 161 233 0328  
[En Español](#)

**PDAB board meeting:**  
**Wednesday, May 20, 8 a.m. to noon**  
Join ZoomGov Meeting  
Meeting ID: 161 233 0328



Share your story in the online feedback form:  
[dfr.oregon.gov/pdab/](https://dfr.oregon.gov/pdab/)