

**A-Engrossed**  
**House Bill 2123**

Ordered by the House April 3  
Including House Amendments dated April 3

Introduced and printed pursuant to House Rule 12.00. Pre-session filed (at the request of House Interim Committee on Health Care)

**SUMMARY**

The following summary is not prepared by the sponsors of the measure and is not a part of the body thereof subject to consideration by the Legislative Assembly. It is an editor's brief statement of the essential features of the measure.

Requires State Board of Pharmacy to license pharmacy benefit managers.

Imposes limits on audits of pharmacies by pharmacy benefit managers and other entities.

*[Requires pharmacy benefit managers that have contracted with provider of health care plan or that are under control of provider of health care plan to permit covered individuals to fill mail order prescriptions at retail community pharmacy in same manner and at similar price that individuals fill orders at mail order pharmacies.]*

Places restrictions on use of maximum allowable cost pricing index by pharmacy benefit managers.

**A BILL FOR AN ACT**

1  
2 Relating to prescription drugs.

3 **Be It Enacted by the People of the State of Oregon:**

4 **SECTION 1. Sections 2 to 4 of this 2013 Act are added to and made a part of ORS chapter**  
5 **689.**

6 **SECTION 2. (1) As used in this section and sections 3 and 4 of this 2013 Act:**

7 (a) "Insurer" has the meaning given that term in ORS 731.106.

8 (b)(A) "Pharmacy benefit manager" means a person that contracts with pharmacies on  
9 behalf of an insurer, a third party administrator or the Oregon Prescription Drug Program  
10 established in ORS 414.312 to:

11 (i) Process claims for prescription drugs or medical supplies or provide retail network  
12 management for pharmacies or pharmacists;

13 (ii) Pay pharmacies or pharmacists for prescription drugs or medical supplies;

14 (iii) Contract with pharmacies or pharmacists for the procurement of prescription drugs  
15 or medical supplies; or

16 (iv) Negotiate rebates with manufacturers for drugs paid for or procured as described in  
17 this paragraph.

18 (B) "Pharmacy benefit manager" does not include a health care service contractor as  
19 defined in ORS 750.005.

20 (c) "Third party administrator" means a person licensed under ORS 744.702.

21 (2) A person must obtain a license from the State Board of Pharmacy in order to act as  
22 a pharmacy benefit manager in this state. The license must be renewed annually. The board  
23 shall establish by rule the procedure and qualifications for obtaining and renewing a license  
24 under this section. The procedure must include a requirement to:

**NOTE:** Matter in **boldfaced** type in an amended section is new; matter *[italic and bracketed]* is existing law to be omitted. New sections are in **boldfaced** type.

1 (a) Submit an application, in a form prescribed by the board, that contains the name and  
2 address of an agent for the service of process;

3 (b) Pay a fee established by the board; and

4 (c) Verify that the applicant has obtained a surety bond.

5 (3) The board may refuse to issue or renew, or may suspend or revoke, a pharmacy  
6 benefit manager license if the applicant or licensee:

7 (a) Fails to comply with this section or section 3 or 4 of this 2013 Act;

8 (b) Engages in conduct likely to mislead, deceive or defraud the general public or the  
9 board;

10 (c) Engages in unfair or deceptive business practices; or

11 (d) Fails to pay fees or fines.

12 (4) The board shall deposit all moneys collected under this section into the State Board  
13 of Pharmacy Account established in ORS 689.139. Moneys collected under this section may  
14 be used only for the purpose of administering this section and sections 3 and 4 of this 2013  
15 Act.

16 **SECTION 3. (1) As used in this section:**

17 (a) "Audit" means an on-site or remote review of the records of a pharmacy by or on  
18 behalf of an entity.

19 (b) "Claim" means a request from a pharmacy or pharmacist to be reimbursed for the  
20 cost of filling or refilling a prescription for a drug or for providing a medical supply or ser-  
21 vice.

22 (c) "Clerical error" means a minor error:

23 (A) In the keeping, recording or transcribing of records or documents or in the handling  
24 of electronic or hard copies of correspondence;

25 (B) That does not result in financial harm to an entity; and

26 (C) That does not involve dispensing an incorrect dose, amount or type of medication or  
27 dispensing a prescription drug to the wrong person.

28 (d) "Entity" includes:

29 (A) A pharmacy benefit manager;

30 (B) An insurer;

31 (C) A third party administrator;

32 (D) A state agency; or

33 (E) A person that represents or is employed by one of the entities described in this par-  
34 agraph.

35 (e) "Fraud" means knowingly and willfully executing or attempting to execute a scheme,  
36 in connection with the delivery of or payment for health care benefits, items or services, that  
37 uses false or misleading pretenses, representations or promises to obtain any money or  
38 property owned by or under the custody or control of any person.

39 (2) An entity that audits claims:

40 (a) Must establish, in writing, a procedure for a pharmacy to appeal the entity's findings  
41 with respect to a claim and must provide a pharmacy with a notice regarding the procedure,  
42 in writing or electronically, prior to conducting an audit of the pharmacy's claims;

43 (b) Must give at least 15 days' advance written notice of an audit to the pharmacy or  
44 corporate headquarters of the pharmacy;

45 (c) Must conduct the audit in consultation with a pharmacist if the audit involves clinical

1 or professional judgment;

2 (d) May not conduct an audit of a claim more than 24 months after the date the claim  
3 was adjudicated by the entity;

4 (e) May not conduct the audit during the first five days of any month without the  
5 pharmacy's consent;

6 (f) May not review more than 200 claims of a pharmacy in any 12-month period except  
7 in cases of alleged fraud;

8 (g) May not conduct more than one on-site audit of a pharmacy in any 12-month period;

9 (h) Must use the same standards and procedures for all pharmacies of a similar size and  
10 doing a similar volume of business;

11 (i) Must pay any outstanding claims of a pharmacy no more than 45 days after the earlier  
12 of the date all appeals are concluded or the date a final report is issued under subsection (8)  
13 of this section;

14 (j) May not include dispensing fees or interest in the amount of any overpayment as-  
15 sessed on a claim unless the overpaid claim was for a prescription that was not filled cor-  
16 rectly;

17 (k) May not recoup costs associated with:

18 (A) Clerical errors; or

19 (B) Other errors that do not result in financial harm to the entity or a consumer;

20 (L) May not charge a pharmacy for a denied or disputed claim until the audit and the  
21 appeals procedure established in paragraph (a) of this subsection are final;

22 (m) May not offset the amount of an overpayment against future remittances; and

23 (n) Must bill a pharmacy separately for the amount of the overpayment.

24 (3) An entity's finding that a claim was incorrectly presented or paid must be based on  
25 identified transactions and not based on probability sampling, extrapolation or other means  
26 that project an error using the number of patients served who have a similar diagnosis or  
27 the number of similar prescriptions or refills for similar drugs.

28 (4) An entity that contracts with an independent third party to conduct audits may not:

29 (a) Agree to compensate the independent third party based on a percentage of the  
30 amount of overpayments recovered; or

31 (b) Disclose information obtained during an audit except to the contracting entity, the  
32 pharmacy subject to the audit or the holder of the policy or certificate of insurance that paid  
33 the claim.

34 (5) For purposes of this section, an entity, or an independent third party that contracts  
35 with an entity to conduct audits, must accept as validation of a claim:

36 (a) An electronic or physical copy of a prescription that complies with this chapter if the  
37 prescribed drug was, within 14 days of the dispensing date:

38 (A) Picked up by the patient or the patient's designee;

39 (B) Delivered by the pharmacy to the patient; or

40 (C) Sent by the pharmacy to the patient using the United States Postal Service or other  
41 common carrier;

42 (b) Point of sale electronic register data showing purchase of the prescribed drug, med-  
43 ical supply or service by the patient or the patient's designee; or

44 (c) Electronic records, including electronic beneficiary signature logs, electronically  
45 scanned and stored patient records maintained at or accessible to the audited pharmacy's

1 central operations and any other reasonably clear and accurate electronic documentation  
2 that corresponds to a claim.

3 (6)(a) After conducting an audit, an entity must provide the pharmacy that is the subject  
4 of the audit with a preliminary report of the audit. The preliminary report must be received  
5 by the pharmacy no later than 30 days after the date on which the audit was completed and  
6 must be sent:

7 (A) By mail or common carrier with a return receipt requested; or

8 (B) Electronically with electronic receipt confirmation.

9 (b) An entity shall provide a pharmacy receiving a preliminary report under this sub-  
10 section no fewer than 45 days after receiving the report to contest the report or any findings  
11 in the report in accordance with the procedure established in subsection (2)(a) of this section  
12 and to provide additional documentation in support of the claim. The entity shall approve a  
13 reasonable request for an extension of time to submit documentation to contest the report  
14 or any findings in the report.

15 (7) If an audit results in a full or partial denial of a claim, the entity conducting the audit  
16 shall allow the pharmacy to resubmit the claim using any commercially reasonable method.

17 (8) An entity must provide a pharmacy that is the subject of an audit with a final report  
18 of the audit no later than 60 days after the later of the date the preliminary report was re-  
19 ceived or the date the pharmacy contested the report using the procedure established in  
20 subsection (2)(a) of this section. The final report must include a final accounting of all  
21 moneys to be recovered by the entity.

22 (9) This section does not preclude an entity from instituting an action for fraud against  
23 a pharmacy.

24 (10) This section does not apply to any audit or investigation that follows a finding:

25 (a) Of fraud;

26 (b) That a claim was submitted for an item or service that was not provided;

27 (c) That a pharmacy deliberately submitted duplicate claims for an item or service and  
28 the duplicate claims did not result from a clerical error;

29 (d) That a pharmacy altered claim forms, electronic claim records or medical documen-  
30 tation for the purpose of receiving a greater amount of reimbursement;

31 (e) Of soliciting, offering or receiving a kickback or bribe;

32 (f) Of collusion between a pharmacy or pharmacist and a patient to defraud the entity;

33 (g) That a pharmacy misrepresented a date or description of items or services furnished  
34 or the identity of the provider or recipient of items or services;

35 (h) That a claim for a prescription was submitted without a prescription's being on file  
36 or was submitted for an over-the-counter item;

37 (i) That a pharmacy filled a prescription using an expired product;

38 (j) That a claim was submitted using an incorrect national drug code number or claiming  
39 reimbursement for a brand name drug when a generic drug was dispensed;

40 (k) That a pharmacy failed to credit the entity for a prescription or a portion of a pre-  
41 scription that was obtained by a patient more than 14 days after the drug was dispensed,  
42 unless good cause exists for the delay; or

43 (L) That a pharmacy submitted a claim without proof that the item or service was pur-  
44 chased.

45 (11) This section does not apply to a state agency that is conducting audits or a person

1 that has contracted with a state agency to conduct audits of pharmacy records for pre-  
2 scription drugs paid for by the state medical assistance program.

3 **SECTION 4.** (1) As used in this section:

4 (a) "List" means the list of drugs for which a third party administrator has established  
5 maximum allowable costs.

6 (b) "Maximum allowable cost" means the maximum amount that a pharmacy benefit  
7 manager will reimburse a pharmacy for the cost of a drug.

8 (c) "Multiple source drug" means a therapeutically equivalent drug that is available from  
9 at least two manufacturers.

10 (d) "Network pharmacy" means a retail drug outlet registered under ORS 689.305 that  
11 contracts with a pharmacy benefit manager.

12 (e) "Therapeutically equivalent" has the meaning given that term in ORS 689.515.

13 (2) A pharmacy benefit manager may not place a drug on a list unless:

14 (a) There are at least two therapeutically equivalent, multiple source drugs, or at least  
15 one generic drug available from only one manufacturer, generally available for purchase by  
16 network pharmacies from national or regional wholesalers; and

17 (b) The drug is not obsolete.

18 (3) A pharmacy benefit manager:

19 (a) Shall provide to each network pharmacy at the beginning of the term of the contract,  
20 and upon each renewal of the contract, notice of the sources used by the pharmacy benefit  
21 manager to determine maximum allowable costs and the lists that apply to the network  
22 pharmacy;

23 (b) Shall make its lists available to a network pharmacy in a format that is readily ac-  
24 cessible to and usable by the pharmacy;

25 (c) Shall update the lists at least once every seven business days and promptly notify  
26 network pharmacies of any changes;

27 (d) May not set a maximum allowable cost below the cost set by the sources described  
28 in paragraph (a) of this subsection; and

29 (e) May not include dispensing fees in the calculation of the maximum allowable cost.

30 (4)(a) A pharmacy benefit manager must establish a reasonable administrative process  
31 for a network pharmacy to request an adjustment of a maximum allowable cost.

32 (b) A pharmacy benefit manager must make a determination on a request for adjustment  
33 no later than seven business days after the pharmacy makes the request.

34 (c) If the pharmacy benefit manager makes an adjustment in response to a request by a  
35 network pharmacy under this subsection, the pharmacy benefit manager shall apply the ad-  
36 justment to all network pharmacies retroactive to the date of the determination under par-  
37 agraph (b) of this subsection.

38 (5) This section does not apply to the state medical assistance program.

39 **SECTION 5.** Section 4 of this 2013 Act applies to contracts between pharmacies and  
40 pharmacy benefit managers that are entered into, renewed or extended on or after the ef-  
41 fective date of this 2013 Act.