

IN THE HOUSE OF REPRESENTATIVES

HOUSE BILL NO. 270

BY HEALTH AND WELFARE COMMITTEE

AN ACT

1 RELATING TO THE FAIR PHARMACY AUDITS ACT; AMENDING TITLE 41, IDAHO CODE,  
2 BY THE ADDITION OF A NEW CHAPTER 66, TITLE 41, IDAHO CODE, TO PROVIDE A  
3 SHORT TITLE, TO PROVIDE PURPOSE AND APPLICABILITY, TO PROVIDE REQUIRE-  
4 MENTS AND PROHIBITIONS FOR PHARMACY AUDITS, TO PROVIDE FOR AN APPEALS  
5 PROCESS, AND TO PROHIBIT EXTRAPOLATION AUDITS; AND DECLARING AN EMER-  
6 GENCY AND PROVIDING AN EFFECTIVE DATE.  
7

8 Be It Enacted by the Legislature of the State of Idaho:

9 SECTION 1. That Title 41, Idaho Code, be, and the same is hereby amended  
10 by the addition thereto of a NEW CHAPTER, to be known and designated as Chap-  
11 ter 66, Title 41, Idaho Code, and to read as follows:

12 CHAPTER 66  
13 FAIR PHARMACY AUDITS ACT

14 41-6601. SHORT TITLE. This chapter shall be known and may be cited as  
15 the "Fair Pharmacy Audits Act."

16 41-6602. PURPOSE AND APPLICABILITY. (1) The purpose of this chapter  
17 is to establish minimum and uniform standards and criteria for the audit of  
18 pharmacies.

19 (2) The provisions of this chapter shall apply to any audit of a phar-  
20 macy conducted on or after July 1, 2023, unless:

21 (a) Contrary provisions for a specific type of audit are provided in  
22 federal or state law, rule, or procedure;

23 (b) The audit relates to medicaid payments; or

24 (c) The audit is an investigative audit based on reasonable suspicion  
25 of willful misrepresentation, abuse, or fraud.

26 41-6603. REQUIREMENTS AND PROHIBITIONS FOR PHARMACY AUDITS. (1) Any  
27 person or entity conducting an audit of a pharmacy shall:

28 (a) If performing the audit pursuant to a contract, identify and  
29 specifically describe the contract provisions authorizing the audit,  
30 including provisions relating to audit appeals. No contract may re-  
31 quire prescription claim documentation or recordkeeping requirements  
32 that exceed requirements set forth in applicable federal or state law,  
33 regulation, or rule;

34 (b) Give written notice to the pharmacy and the pharmacy's contracting  
35 agent, including identification of specific prescription numbers and  
36 fill dates to be audited, at least fourteen (14) days prior to conduct-  
37 ing the audit. For purposes of this subsection, the term "audit" in-  
38 cludes but is not limited to an on-site audit, a desk audit, a wholesale  
39 purchase audit, a request for documentation related to the dispensing

1 of a prescription drug, and a request for documentation related to any  
2 reimbursed activity by a pharmacy provider. Notice of a wholesale pur-  
3 chase audit shall be given at least thirty (30) days prior to the audit.  
4 The pharmacy shall have the opportunity to reschedule any audit no more  
5 than seven (7) days from the date designated on the original audit noti-  
6 fication;

7 (c) Not interfere with the delivery of pharmacist services to a patient  
8 and use every reasonable effort to minimize inconvenience and interrup-  
9 tion to pharmacy operations during the audit process;

10 (d) Conduct any audit involving clinical or professional judgment by  
11 means of or in consultation with a licensed pharmacist;

12 (e) Not consider as fraud any clerical or recordkeeping error, such as  
13 a typographical error, scrivener's error, or computer error, includ-  
14 ing but not limited to a miscalculated day supply, an incorrectly billed  
15 prescription written date, or an incorrect prescription origin code.  
16 Such errors shall not be subject to recoupment. The pharmacy shall have  
17 the right to submit amended claims electronically to correct clerical  
18 or recordkeeping errors in lieu of recoupment. To the extent that an au-  
19 dit results in the identification of any clerical or recordkeeping er-  
20 rors in a required document or record, the pharmacy shall not be subject  
21 to recoupment of funds unless there is proof of intent to commit fraud.  
22 A person shall not be subject to criminal penalties for errors provided  
23 for in this paragraph without proof of the intent required for convic-  
24 tion of the applicable crime;

25 (f) Permit a pharmacy to use computerized patterned medical records or  
26 the records of a hospital, physician, or other authorized health care  
27 practitioner for drugs or medicinal supplies written or transmitted  
28 by any means of communication for purposes of validating the pharmacy  
29 record with respect to orders or refills of a legend or other prescribed  
30 drug;

31 (g) Not include the dispensing fee amount or the actual invoice cost of  
32 the prescription dispensed in a finding of an audit recoupment unless  
33 a prescription was not actually dispensed or a physician denied autho-  
34 rization of a dispensing order;

35 (h) Audit each pharmacy under identical standards, regularity, and pa-  
36 rameters as other similarly situated pharmacies. If the person or en-  
37 tity conducting the audit owns or manages pharmacies, all audits of such  
38 pharmacies shall be conducted under identical standards, regularity,  
39 and parameters;

40 (i) Not exceed six (6) months from the date the claim was submitted to or  
41 adjudicated by the person or entity conducting the audit;

42 (j) Not schedule or initiate an audit during the first seven (7) calen-  
43 dar days of any month unless otherwise consented to by the pharmacy;

44 (k) Disclose to any plan sponsor whose claims were included in the audit  
45 any money recouped in the audit;

46 (l) Not require pharmacists to break open packaging labeled "for  
47 single-patient-use only." Packaging labeled "for single-patient-use  
48 only" shall be deemed to be the smallest package size available; and

1 (m) Permit a pharmacy to use the written statement of a patient or the  
2 patient's agent for purposes of validating the delivery of a prescrip-  
3 tion to the possession of the patient or the patient's agent.

4 (2) Any person or entity that conducts wholesale purchase review dur-  
5 ing an audit of a pharmacist or pharmacy shall not require the pharmacist or  
6 pharmacy to provide a full dispensing report. Wholesaler invoice reviews  
7 shall be limited to verification of purchase inventory specific to the phar-  
8 macy claims paid by the person or entity conducting the audit or the person or  
9 entity on whose behalf the audit is being conducted.

10 (3) Any person or entity conducting an audit shall not identify or label  
11 a prescription claim as an audit discrepancy when:

12 (a) The national drug code for the dispensed drug is in a quantity that  
13 is a subunit or multiple of the drug purchased by the pharmacist or phar-  
14 macy as supported by a wholesale invoice;

15 (b) The pharmacist or pharmacy dispensed the correct quantity of the  
16 drug according to the prescription; and

17 (c) The drug dispensed by the pharmacist or pharmacy shares all but the  
18 last two (2) digits of the national drug code of the drug reflected on  
19 the supplier invoice.

20 (4) Any person or entity conducting an audit shall accept as evidence,  
21 subject to validation, to support the validity of a pharmacy claim related to  
22 a dispensed drug:

23 (a) Redacted copies of supplier invoices in the pharmacist's or phar-  
24 macy's possession; or

25 (b) Invoices and any supporting documents from any supplier as autho-  
26 rized by federal or state law to transfer ownership of the drug acquired  
27 by the pharmacist or pharmacy.

28 (5) Any person or entity conducting an audit shall provide, no later  
29 than five (5) business days after the date of a request by the pharmacist or  
30 pharmacy, all supporting documents the pharmacist's or pharmacy's purchase  
31 suppliers provided to the person or entity on whose behalf the audit is being  
32 conducted.

33 (6) Any person or entity conducting an audit shall not audit more than  
34 fifty (50) prescriptions, based on date of service, per calendar year. The  
35 annual limit to the number of prescription claims audited shall be inclusive  
36 of all audits, including any prescription-related documentation requests  
37 from the person or entity conducting the audit or the person or entity on  
38 whose behalf the audit is being conducted during a calendar year.

39 (7) If paper copies of records are requested by the person or en-  
40 tity conducting an audit, the person or entity shall pay twenty-five cents  
41 (\$0.25) per page to cover the costs incurred by the pharmacy. The person or  
42 entity conducting the audit shall provide the pharmacy with accurate in-  
43 structions, including any required form for obtaining reimbursement for the  
44 copied records.

45 (8) The person or entity conducting an audit shall:

46 (a) Deliver a preliminary audit findings report to the pharmacy and the  
47 pharmacy's contracting agent within forty-five (45) calendar days of  
48 conducting the audit;

49 (b) Allow the pharmacy at least ninety (90) calendar days following re-  
50 ceipt of the preliminary audit findings report in which to produce docu-

1           mentation to address any discrepancy found during the audit. A pharmacy  
2           may request an extension, not to exceed an additional forty-five (45)  
3           calendar days;

4           (c) Deliver a final audit findings report to the pharmacy and the phar-  
5           macy's contracting agent signed by the auditor within ten (10) calendar  
6           days after receipt of documentation or evidence provided by the phar-  
7           macy, as provided for in section 41-6604, Idaho Code;

8           (d) Allow the pharmacy to reverse and resubmit claims electronically  
9           within thirty (30) days of receipt of the final audit report in lieu of  
10          the auditing entity recouping discrepant claim amounts from the phar-  
11          macy;

12          (e) Not recoup any disputed funds until after final disposition of the  
13          audit findings, including the appeals process as provided for in sec-  
14          tion 41-6604, Idaho Code; and

15          (f) Not accrue interest during the audit and appeal period.

16          (9) Each person or entity conducting an audit shall provide a copy of  
17          the final audit results, and a final audit report upon request, after comple-  
18          tion of any review process to any plan sponsor whose claims were included in  
19          the audit.

20          (10) The full amount of any recoupment on an audit shall be refunded to  
21          the plan sponsor whose claims were included in the audit and to whom the re-  
22          coupment is owing. Except as otherwise provided for in this subsection, a  
23          charge or assessment for an audit shall not be based, directly or indirectly,  
24          on amounts recouped. This subsection does not prevent the person or entity  
25          conducting the audit from charging or assessing the responsible party, di-  
26          rectly or indirectly, based on amounts recouped if both of the following con-  
27          ditions are met:

28          (a) The plan sponsor and the person or entity conducting the audit have  
29          a contract that explicitly states the percentage charge or assessment  
30          to the plan sponsor; and

31          (b) A commission to an agent or employee of the person or entity con-  
32          ducting the audit is not based, directly or indirectly, on amounts re-  
33          couped.

34          (11) Unless the provisions of this subsection are superseded by state or  
35          federal law, auditors shall have access to previous audit reports on a par-  
36          ticular pharmacy only when the previous audits were conducted by the audit-  
37          ing person or entity for the same person or entity on whose behalf the audit  
38          is being conducted. An auditing vendor contracting with multiple persons or  
39          entities shall not use audit reports or other information gained from an au-  
40          dit on a pharmacy to conduct another audit for another person or entity.

41          41-6604. APPEALS PROCESS. (1) Each person or entity conducting an au-  
42          dit shall establish a written appeals process under which a pharmacy may ap-  
43          peal an unfavorable preliminary audit report or final audit report to the  
44          person or entity. The pharmacy may submit documentation or other evidence to  
45          support its appeal.

46          (2) Following an appeal, if the person or entity finds that an unfavor-  
47          able audit report or any portion of such report is unsubstantiated, the per-  
48          son or entity shall dismiss the audit report or the unsubstantiated portion  
49          of the audit report without any further action.

1           (3) Any final audit report, following the final audit appeal period,  
2 with a finding of potential criminal conduct shall be referred to the pros-  
3 ecuting attorney having proper jurisdiction upon completion of the appeals  
4 process.

5           41-6605.   EXTRAPOLATION AUDIT PROHIBITED. (1) As used in this section,  
6 "extrapolation audit" means an audit of a sample of prescription drug bene-  
7 fit claims submitted by a pharmacy to the person or entity conducting an au-  
8 dit that is then used to estimate audit results for a larger batch or group of  
9 claims not reviewed by the auditor.

10           (2) No person or entity may conduct an extrapolation audit. A person or  
11 entity conducting an audit shall not use the accounting practice of extrap-  
12 olation in calculating recoupments or penalties for audits.

13           SECTION 2. An emergency existing therefor, which emergency is hereby  
14 declared to exist, this act shall be in full force and effect on and after  
15 July 1, 2023.