

IN THE HOUSE OF REPRESENTATIVES

HOUSE BILL NO. 291

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 EXCEPT UNDER CERTAIN CIR-
6 CUMSTANCES; AND DECLARING AN EMERGENCY AND PROVIDING AN EFFECTIVE DATE.
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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;

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

26 (d) The audit is a financial examination conducted by a certified pub-
27 lic accountant according to generally accepted auditing standards.

28 41-6603. REQUIREMENTS AND PROHIBITIONS FOR PHARMACY AUDITS. (1) Any
29 person or entity conducting an audit of a pharmacy shall:

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

36 (b) Give written notice to the pharmacy and the pharmacy's contracting
37 agent at least fourteen (14) days prior to conducting the on-site au-
38 dit. For purposes of this subsection, the term "audit" means an audit
39 conducted on behalf of an auditing entity of any records of a pharmacy

1 for drugs dispensed by a pharmacy to a covered individual. The pharmacy
2 shall have the opportunity to reschedule any on-site audit no more than
3 seven (7) days from the date designated on the original audit notifica-
4 tion;

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

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

10 (e) Prior to leaving the pharmacy after the on-site portion of the phar-
11 macy audit, provide to the pharmacy a complete list of pharmacy records
12 reviewed;

13 (f) Not subject a pharmacy to a charge-back or recoupment for a cleri-
14 cal or recordkeeping error, such as a typographical error, scrivener's
15 error, or computer error, including but not limited to a miscalculated
16 day supply, an incorrectly billed prescription written date, or an in-
17 correct prescription origin code, unless the error resulted in overpay-
18 ment to the pharmacy. Prior to payment of the claim, the pharmacy shall
19 have the right to submit amended claims electronically to correct cler-
20 ical or recordkeeping errors in lieu of recoupment. A person shall not
21 be subject to criminal penalties for errors described in this paragraph
22 without proof of the intent required for conviction of the applicable
23 crime;

24 (g) Limit any fee, charge-back, recoupment, or other adjustment to the
25 actual overpayment associated with the dispensed product or portion of
26 the dispensed product or the actual underpayment or overpayment as set
27 forth in this subsection;

28 (h) Permit a pharmacy to use any valid prescription, including com-
29 puterized patterned medical records or the records of a hospital,
30 physician, or other authorized health care practitioner for drugs or
31 medicinal supplies written or transmitted by any means of communication
32 for purposes of validating the pharmacy record with respect to orders or
33 refills of a legend or other prescribed drug. Documentation of an oral
34 prescription order that has been verified by the prescribing health
35 care provider shall meet the provisions of this paragraph for the ini-
36 tial audit review;

37 (i) Permit a pharmacy to use authentic and verifiable statements
38 or records, including but not limited to medication administration
39 records of a nursing home, assisted living facility, hospital, or
40 health care provider with prescriptive authority, to validate the phar-
41 macy record and delivery;

42 (j) Not include the dispensing fee in the calculation of overpayment of
43 the prescription dispensed in a finding of an audit recoupment unless
44 a prescription was not actually dispensed or a physician denied autho-
45 rization of a dispensing order;

46 (k) Audit each pharmacy under standards, regularity, and parameters as
47 other similarly situated pharmacies in a pharmacy network contract in
48 this state. If the person or entity conducting the audit owns or man-
49 ages pharmacies, all audits of such pharmacies shall be conducted un-

1 der standards, regularity, and parameters as other similarly situated
2 pharmacies in a pharmacy network contract in this state;

3 (l) Not exceed fifteen (15) months from the date the claim was submitted
4 to or adjudicated by the person or entity conducting the audit;

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

7 (n) Disclose to any plan sponsor whose claims were included in the audit
8 any money recouped in the audit;

9 (o) Provide network pharmacies information on the adjudication process
10 for unit of use prescription products where the smallest unit either ex-
11 ceeds or does not maximize the benefit day's supply; and

12 (p) Permit a pharmacy to use a paper or electronic signature log that
13 documents the delivery of a prescription to the possession of the pa-
14 tient or the patient's agent.

15 (2) Except as otherwise provided by federal or state law, an auditing
16 entity that audits wholesale invoices during an audit of a pharmacy may not
17 audit the pharmacy claims of another health benefit plan or pharmacy benefit
18 manager.

19 (3) Any person or entity conducting a wholesale invoice audit shall not
20 identify or label a prescription claim as an audit discrepancy when:

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

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

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

29 (4) Any person or entity conducting a wholesale invoice audit shall ac-
30 cept as evidence, subject to validation, to support the validity of a phar-
31 macy claim related to a dispensed drug:

32 (a) Supplier invoices issued before the date the drug was dispensed in
33 the pharmacist's or pharmacy's possession; or

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

37 (5) Any person or entity conducting a wholesale invoice audit shall
38 provide, no later than five (5) business days after the date of a request
39 by the pharmacist or pharmacy, all supporting documents the pharmacist's
40 or pharmacy's purchase suppliers provided to the person or entity on whose
41 behalf the audit is being conducted.

42 (6) Any person or entity conducting an audit shall not audit more than
43 two hundred fifty (250) prescriptions, based on date of service, per calen-
44 dar year. The annual limit to the number of prescription claims audited
45 shall be inclusive of all audits, including any prescription-related docu-
46 mentation requests from the person or entity conducting the audit or the per-
47 son or entity on whose behalf the audit is being conducted during a calendar
48 year.

49 (7) If paper copies of records are requested by the person or entity
50 conducting an audit, the person or entity shall pay twenty-five cents (25¢)

1 per page to cover the costs incurred by the pharmacy. The person or entity
2 conducting the audit shall provide the pharmacy with accurate instructions,
3 including any required form for obtaining reimbursement for the copied
4 records.

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

6 (a) Deliver a preliminary audit findings report to the pharmacy and
7 the pharmacy's contracting agent within sixty (60) calendar days of
8 conducting the audit. The preliminary report shall include contact
9 information for the auditing entity that conducted the pharmacy audit
10 and an appropriate and accessible point of contact, including telephone
11 number, facsimile number, electronic mail address, and auditing firm
12 name and address so that audit results, procedures, and discrepancies
13 may be reviewed. The preliminary audit report shall include but is not
14 limited to claim level information for any discrepancy found and total
15 dollar amounts of claims subject to recoupment;

16 (b) Allow the pharmacy at least sixty (60) calendar days following re-
17 ceipt of the preliminary audit findings report in which to produce docu-
18 mentation to address any discrepancy found during the audit. A pharmacy
19 may request an extension, not to exceed an additional thirty (30) calen-
20 dar days;

21 (c) Deliver a final audit findings report to the pharmacy and the phar-
22 macy's contracting agent signed by the auditor within thirty (30) cal-
23 endar days after receipt of documentation or evidence provided by the
24 pharmacy, as provided for in section 41-6604, Idaho Code;

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

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

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

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

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

45 (a) The plan sponsor and the person or entity conducting the audit have
46 a contract that explicitly states the percentage charge or assessment
47 to the plan sponsor; and

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

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

8 41-6604. APPEALS PROCESS. (1) Each person or entity conducting an au-
9 dit shall establish a written appeals process under which a pharmacy may ap-
10 peal an unfavorable preliminary audit report or final audit report to the
11 person or entity. The pharmacy must submit documentation or other evidence
12 to support its appeal.

13 (2) Following an appeal, if the person or entity finds that an unfavor-
14 able audit report is unsubstantiated, the person or entity shall dismiss the
15 unsubstantiated portion of the audit report.

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

20 41-6605. EXTRAPOLATION AUDIT PROHIBITED. (1) As used in this section,
21 "extrapolation audit" means an audit of a sample of prescription drug bene-
22 fit claims submitted by a pharmacy to the person or entity conducting an au-
23 dit that is then used to estimate audit results for a larger batch or group of
24 claims not reviewed by the auditor.

25 (2) No person or entity may conduct an extrapolation audit unless oth-
26 erwise required by federal law or federal plans. A person or entity conduct-
27 ing an audit shall not use the accounting practice of extrapolation in calcu-
28 lating recoupments or penalties for audits.

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