## ESSB 6137 - H COMM AMD

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By Committee on Health Care & Wellness

- 1 Strike everything after the enacting clause and insert the 2 following:
- 3 "NEW SECTION. Sec. 1. The definitions in this section apply 4 throughout this chapter unless the context clearly requires otherwise.
- 5 (1) "Claim" means a request from a pharmacy or pharmacist to be 6 reimbursed for the cost of filling or refilling a prescription for a 7 drug or for providing a medical supply or service.
  - (2) "Insurer" has the same meaning as in RCW 48.01.050.
  - (3) "Pharmacist" has the same meaning as in RCW 18.64.011.
- 10 (4) "Pharmacy" has the same meaning as in RCW 18.64.011.
- 11 (5)(a) "Pharmacy benefit manager" means a person that contracts 12 with pharmacies on behalf of an insurer, a third-party payor, or the 13 prescription drug purchasing consortium established under RCW 70.14.060 14 to:
- 15 (i) Process claims for prescription drugs or medical supplies or 16 provide retail network management for pharmacies or pharmacists;
- 17 (ii) Pay pharmacies or pharmacists for prescription drugs or 18 medical supplies; or
- 19 (iii) Negotiate rebates with manufacturers for drugs paid for or 20 procured as described in this subsection.
- 21 (b) "Pharmacy benefit manager" does not include a health care 22 service contractor as defined in RCW 48.44.010.
- 23 (6) "Third-party payor" means a person licensed under RCW 24 48.39.005.
- NEW SECTION. Sec. 2. (1) To conduct business in this state, a pharmacy benefit manager must register with the department of revenue's business licensing service and annually renew the registration.
- 28 (2) To register under this section, a pharmacy benefit manager 29 must:

- 1 (a) Submit an application requiring the following information:
  - (i) The identity of the pharmacy benefit manager;
- 3 (ii) The name, business address, phone number, and contact person 4 for the pharmacy benefit manager; and
- 5 (iii) Where applicable, the federal tax employer identification 6 number for the entity; and
  - (b) Pay a registration fee of two hundred dollars.
- 8 (3) To renew a registration under this section, a pharmacy benefit 9 manager must pay a renewal fee of two hundred dollars.
- 10 (4) All receipts from registrations and renewals collected by the 11 department must be deposited into the business license account created 12 in RCW 19.02.210.

## 13 <u>NEW SECTION.</u> **Sec. 3.** As used in sections 3 through 9 of this act:

- 14 (1) "Audit" means an on-site or remote review of the records of a pharmacy by or on behalf of an entity.
  - (2) "Clerical error" means a minor error:
- 17 (a) In the keeping, recording, or transcribing of records or 18 documents or in the handling of electronic or hard copies of 19 correspondence;
  - (b) That does not result in financial harm to an entity; and
- 21 (c) That does not involve dispensing an incorrect dose, amount or 22 type of medication, or dispensing a prescription drug to the wrong 23 person.
  - (3) "Entity" includes:
  - (a) A pharmacy benefit manager;
- 26 (b) An insurer;

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- 27 (c) A third-party payor;
- 28 (d) A state agency; or
- 29 (e) A person that represents or is employed by one of the entities 30 described in this subsection.
- 31 (4) "Fraud" means knowingly and willfully executing or attempting 32 to execute a scheme, in connection with the delivery of or payment for 33 health care benefits, items, or services, that uses false or misleading 34 pretenses, representations, or promises to obtain any money or property 35 owned by or under the custody or control of any person.

- 1 <u>NEW SECTION.</u> **Sec. 4.** An entity that audits claims or an independent third party that contracts with an entity to audit claims:
  - (1) Must establish, in writing, a procedure for a pharmacy to appeal the entity's findings with respect to a claim and must provide a pharmacy with a notice regarding the procedure, in writing or electronically, prior to conducting an audit of the pharmacy's claims;
  - (2) May not conduct an audit of a claim more than twenty-four months after the date the claim was adjudicated by the entity;
  - (3) Must give at least fifteen days' advance written notice of an on-site audit to the pharmacy or corporate headquarters of the pharmacy;
  - (4) May not conduct an on-site audit during the first five days of any month without the pharmacy's consent;
  - (5) Must conduct the audit in consultation with a pharmacist who is licensed by this or another state if the audit involves clinical or professional judgment;
  - (6) May not conduct an on-site audit of more than two hundred fifty unique prescriptions of a pharmacy in any twelve-month period except in cases of alleged fraud;
  - (7) May not conduct more than one on-site audit of a pharmacy in any twelve-month period;
  - (8) Must audit each pharmacy under the same standards and parameters that the entity uses to audit other similarly situated pharmacies;
  - (9) Must pay any outstanding claims of a pharmacy no more than forty-five days after the earlier of the date all appeals are concluded or the date a final report is issued under section 8(3) of this act;
  - (10) May not include dispensing fees or interest in the amount of any overpayment assessed on a claim unless the overpaid claim was for a prescription that was not filled correctly;
    - (11) May not recoup costs associated with:
  - (a) Clerical errors; or

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- 33 (b) Other errors that do not result in financial harm to the entity 34 or a consumer; and
- 35 (12) May not charge a pharmacy for a denied or disputed claim until 36 the audit and the appeals procedure established under subsection (1) of 37 this section are final.

- NEW SECTION. Sec. 5. An entity's finding that a claim was incorrectly presented or paid must be based on identified transactions and not based on probability sampling, extrapolation, or other means that project an error using the number of patients served who have a similar diagnosis or the number of similar prescriptions or refills for similar drugs.
- NEW SECTION. Sec. 6. An entity that contracts with an independent third party to conduct audits may not:
  - (1) Agree to compensate the independent third party based on a percentage of the amount of overpayments recovered; or
- 11 (2) Disclose information obtained during an audit except to the 12 contracting entity, the pharmacy subject to the audit, or the holder of 13 the policy or certificate of insurance that paid the claim.
- NEW SECTION. Sec. 7. For purposes of sections 3 through 9 of this act, an entity, or an independent third party that contracts with an entity to conduct audits, must allow as evidence of validation of a claim:
- 18 (1) An electronic or physical copy of a valid prescription if the 19 prescribed drug was, within fourteen days of the dispensing date:
  - (a) Picked up by the patient or the patient's designee;
  - (b) Delivered by the pharmacy to the patient; or

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- 22 (c) Sent by the pharmacy to the patient using the United States 23 postal service or other common carrier;
  - (2) Point of sale electronic register data showing purchase of the prescribed drug, medical supply, or service by the patient or the patient's designee; or
  - (3) Electronic records, including electronic beneficiary signature logs, electronically scanned and stored patient records maintained at or accessible to the audited pharmacy's central operations, and any other reasonably clear and accurate electronic documentation that corresponds to a claim.
- NEW SECTION. Sec. 8. (1)(a) After conducting an audit, an entity must provide the pharmacy that is the subject of the audit with a preliminary report of the audit. The preliminary report must be

received by the pharmacy no later than forty-five days after the date 1 2 on which the audit was completed and must be sent:

- (i) By mail or common carrier with a return receipt requested; or
- (ii) Electronically with electronic receipt confirmation.

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- (b) An entity shall provide a pharmacy receiving a preliminary report under this subsection no fewer than forty-five days after receiving the report to contest the report or any findings in the report in accordance with the appeals procedure established under section 4(1) of this act and to provide additional documentation in support of the claim. The entity shall consider a reasonable request for an extension of time to submit documentation to contest the report or any findings in the report.
- (2) If an audit results in the dispute or denial of a claim, the entity conducting the audit shall allow the pharmacy to resubmit the claim using any commercially reasonable method, including facsimile, mail, or electronic mail.
- (3) An entity must provide a pharmacy that is the subject of an audit with a final report of the audit no later than sixty days after the later of the date the preliminary report was received or the date the pharmacy contested the report using the appeals procedure established under section 4(1) of this act. The final report must include a final accounting of all moneys to be recovered by the entity.
- (4) Recoupment of disputed funds from a pharmacy by an entity or repayment of funds to an entity by a pharmacy, unless otherwise agreed to by the entity and the pharmacy, shall occur after the audit and the appeals procedure established under section 4(1) of this act are final. If the identified discrepancy for an individual audit exceeds forty thousand dollars, any future payments to the pharmacy may be withheld by the entity until the audit and the appeals procedure established under section 4(1) of this act are final.

## 31 NEW SECTION. Sec. 9. Sections 3 through 9 of this act do not:

- 32 (1) Preclude an entity from instituting an action for fraud against 33 a pharmacy;
- 34 (2) Apply to an audit of pharmacy records when fraud or other 35 intentional and willful misrepresentation is indicated by physical 36 review, review of claims data or statements, or other investigative 37 methods; or

1 (3) Apply to a state agency that is conducting audits or a person 2 that has contracted with a state agency to conduct audits of pharmacy 3 records for prescription drugs paid for by the state medical assistance 4 program.

## NEW SECTION. Sec. 10. (1) As used in this section:

- 6 (a) "List" means the list of drugs for which maximum allowable 7 costs have been established.
- 8 (b) "Maximum allowable cost" means the maximum amount that a 9 pharmacy benefit manager will reimburse a pharmacy for the cost of a drug.
- 11 (c) "Multiple source drug" means a therapeutically equivalent drug 12 that is available from at least two manufacturers.
- 13 (d) "Network pharmacy" means a retail drug outlet licensed as a 14 pharmacy under RCW 18.64.043 that contracts with a pharmacy benefit 15 manager.
- 16 (e) "Therapeutically equivalent" has the same meaning as in RCW 17 69.41.110.
  - (2) A pharmacy benefit manager:

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- (a) May not place a drug on a list unless are is at least two therapeutically equivalent multiple source drugs, or at least one generic drug available from only one manufacturer, generally available for purchase by network pharmacies from national or regional wholesalers;
- (b) Shall ensure that all drugs on a list are generally available for purchase by pharmacies in this state from national or regional wholesalers;
  - (c) Shall ensure that all drugs on a list are not obsolete;
- (d) Shall make available to each network pharmacy at the beginning of the term of a contract, and upon renewal of a contract, the sources utilized to determine the maximum allowable cost pricing of the pharmacy benefit manager;
- (e) Shall make a list available to a network pharmacy upon request in a format that is readily accessible to and usable by the network pharmacy;
- 35 (f) Shall update each list maintained by the pharmacy benefit 36 manager every seven business days and make the updated lists, including

all changes in the price of drugs, available to network pharmacies in 1 2 a readily accessible and usable format;

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- (g) Shall ensure that dispensing fees are not included in the calculation of maximum allowable cost.
- (3) A pharmacy benefit manager must establish a process by which a network pharmacy may appeal its reimbursement for a drug subject to maximum allowable cost pricing. A network pharmacy may appeal a maximum allowable cost if the reimbursement for the drug is less than the net amount that the network pharmacy paid to the supplier of the drug. An appeal requested under this section must be completed within thirty calendar days of the pharmacy making the claim for which an appeal has been requested.
- (4) A pharmacy benefit manager must provide as part of the appeals process established under subsection (3) of this section:
- (a) A telephone number at which a network pharmacy may contact the pharmacy benefit manager and speak with an individual who responsible for processing appeals;
- (b) A final response to an appeal of a maximum allowable cost within seven business days; and
- (c) If the appeal is denied, the reason for the denial and the national drug code of a drug that may be purchased by similarly situated pharmacies at a price that is equal to or less than the maximum allowable cost.
- (5)(a) If an appeal is upheld under this section, the pharmacy benefit manager shall make an adjustment on a date no later than one day after the date of determination. The pharmacy benefit manager shall make the adjustment effective for all similarly situated pharmacies in this state that are within the network.
- (b) If the request for an adjustment has come from a critical access pharmacy, as defined by the state health care authority by rule for purposes related to the prescription drug purchasing consortium established under RCW 70.14.060, the adjustment approved under (a) of this subsection shall apply only to critical access pharmacies.
- 34 (6) This section does not apply to the state medical assistance 35 program.
- Sections 1 through 10 of this act 36 NEW SECTION. Sec. 11. 37 constitute a new chapter in Title 19 RCW."

Establishes the registration fee for pharmacy benefit managers at \$200 rather than an amount up to \$200 as established by the Department of Revenue. Specifies that pharmacy benefit managers must register with the Department of Revenue's Business Licensing Service. Removes the limitation on the types of information that must be submitted on an application.

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