1	SENATE BILL 394
2	54TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2019
3	INTRODUCED BY
4	Clemente Sanchez
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10	AN ACT
11	RELATING TO PHARMACIES; PROVIDING FOR CHANGES TO THE PHARMACY
12	AUDIT PROCESS; EXCEPTING CERTAIN AUDIT FINDINGS FROM FORMING
13	THE BASIS FOR RECOUPMENT; ADDING A PHARMACY BENEFITS MANAGER OR
14	ITS SUBCONTRACTOR AS AN AUDITING ENTITY.
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16	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:
17	SECTION 1. Section 61-11-18.2 NMSA 1978 (being Laws 2007,
18	Chapter 15, Section 1) is amended to read:
19	"61-11-18.2. AUDIT OF PHARMACY RECORDS
20	[A. As used in this section, "entity" means a
21	managed care company, insurance company, third-party payor or
22	the representative of the managed care company, insurance
23	company or third-party payor.
24	B.] A. An audit of the records of a pharmacy by an
25	entity shall be conducted in accordance with the following
	.212241.1

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criteria:

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2	(1) the entity conducting the initial on-site
3	audit shall give the pharmacy notice at least two weeks prior
4	to conducting the initial on-site audit for each audit cycle;
5	(2) an audit that involves clinical or
6	professional judgment shall be conducted by or in consultation
7	with a pharmacist;
8	(3) a clerical or [record-keeping]
9	recordkeeping error, regarding a required document or record,
10	shall not necessarily constitute fraud, [but such a claim] <u>and</u>
11	that error:
12	(a) [may be subject to recoupment] <u>shall</u>
13	not be the basis for recoupment unless the error results in
14	demonstrable financial harm; and
15	(b) shall not be subject to criminal
16	penalties without proof of intent to commit fraud;
17	(4) a pharmacy may use the records of a
18	hospital, physician or other authorized practitioner of the
19	healing arts for drugs or medicinal supplies written or
20	transmitted by any means of communication for purposes of
21	validating the pharmacy record with respect to orders or
22	refills of a dangerous drug or controlled substance;
23	(5) a finding of an overpayment or
24	underpayment shall [not be a projection based on the number of
25	patients served having a similar diagnosis or on the number of
	.212241.1 - 2 -

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1 similar orders or refills for similar drugs and recoupment of 2 claims shall be based on the actual overpayment or underpayment [unless the entity demonstrates a statistically 3 justifiable method of projection or the projection for 4 overpayment or underpayment is part of a settlement as agreed 5 to by the pharmacy] of a specific individual claim; 6 7 (6) each pharmacy shall be audited under the same standards and parameters as other similarly situated 8 9 pharmacies audited by the entity; a pharmacy shall be allowed at least 10 (7) twenty-one business days, with reasonable extensions allowed, 11 12 following receipt of the preliminary audit report in which to produce documentation to address any discrepancy found during 13 14 an audit; the period covered by an audit shall not (8) 15 exceed [two years, unless otherwise provided by contractual 16 agreement] one year from the date the claim was submitted to or 17 adjudicated by an entity or unless it conflicts with state or 18 19 federal law; 20 (9) an audit shall not be initiated or scheduled during the first five calendar days of a month [due 21 to the high volume of prescriptions filled during that time 22 unless otherwise consented to by the pharmacy]; 23 (10) the preliminary audit report shall be 24 delivered to the pharmacy within one hundred twenty days, with 25 .212241.1 - 3 -

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1 reasonable extensions allowed, after conclusion of the audit, 2 and the final report shall be delivered to the pharmacy within 3 six months after receipt of the preliminary audit report or final appeal, as provided for in Subsection [Θ] <u>B</u> of this 4 5 section, whichever is later; [(11) the audit criteria set forth in this 6 7 subsection shall apply only to audits of claims submitted for payment after July 1, 2007; and 8 9 (12)] (11) notwithstanding any other provision in this [subsection] section, the entity conducting the audit 10 shall not use the accounting practice of extrapolation in 11 12 calculating recoupments or penalties for audits; (12) a person performing an on-site audit or a 13 desk audit shall not directly or indirectly receive 14 compensation based on the result of the audit; 15 (13) an entity shall not charge a fee for 16 conducting an on-site or a desk audit unless there is a finding 17 of actual fraud; 18 (14) after an audit is initiated, a pharmacist 19 or pharmacy may resubmit a claim to correct clerical or 20 recordkeeping errors; 21 (15) requirements for a valid prescription or 22 a pharmacy's operational standards shall not be more stringent 23 than federal or state requirements; 24 (16) a pharmacy or pharmacist may satisfy 25 .212241.1

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1	state and federal requirements for a valid prescription by
2	affixing or writing additional information on the front or back
3	of a prescription or if the required information is
4	electronically recorded on a patient's profile and is readily
5	<u>retrievable;</u>
6	(17) the days' supply for unit-of-use items,
7	such as topicals, drops, vials and inhalants, shall not be
8	limited beyond manufacturer recommendations;
9	(18) if the only commercially available
10	package size exceeds an entity's maximum days' supply, the
11	dispensing of such package size must be accepted by the entity
12	and shall not be the basis for recoupment;
13	(19) if the only commercially available
14	package size exceeds an entity's maximum days' supply and the
15	entity accepts the refill of such prescription, the entity
16	shall not recoup such claim as an early refill; and
17	(20) the failure of a pharmacy to collect a
18	copayment shall not be the basis for recoupment if the pharmacy
19	provides documentation of billing of the claim and an attempt
20	to collect the copayment.
21	[C.] <u>B.</u> Recoupment of any disputed funds shall
22	occur after final internal disposition of the audit, including
23	the appeals process set forth in Subsection [$ embed{D}$] <u>C</u> of this
24	section. Should the identified discrepancy for an individual
25	audit exceed twenty-five thousand dollars (\$25,000), future
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payments to the pharmacy may be withheld pending finalization
 of the audit.

3 [D.] C. Each entity conducting an audit shall establish an appeals process under which a pharmacy may appeal 4 an unfavorable preliminary audit report to the entity. If, 5 following the appeal, the entity finds that an unfavorable 6 7 audit report or any portion of the audit is unsubstantiated, the entity shall dismiss the audit report or the 8 9 unsubstantiated portion of the report of the audit without the necessity of any further proceedings. 10 [E. This section does not apply to any 11 12 investigative audit that involves probable or potential fraud, willful misrepresentation.] 13 D. In a wholesale invoice audit conducted by an 14 15 entity: (1) an entity shall not audit the claims of 16 another entity; 17 (2) the following shall not form the basis for 18 19 recoupment: 20 (a) the national drug code for the dispensed drug is in a quantity that is a sub-unit or multiple 21 of the purchased drug as reflected on a supporting wholesale 22 invoice; 23 (b) the correct quantity dispensed is 24 reflected on the audited pharmacy claim; 25 .212241.1

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1	(c) the drug dispensed by the pharmacy
2	on an audited pharmacy claim is identical to the strength and
3	dosage form of the drug purchased; or
4	(d) discrepancies in wholesale invoice
5	purchasing where the audited claims are otherwise supported by
6	records of receipt by the patient or patient's agent of the
7	dispensed drug underlying the audited claim;
8	(3) the entity shall accept as evidence:
9	(a) supplier invoices issued prior to
10	the date of dispensing the drug underlying the audited claim;
11	(b) invoices from any supplier
12	authorized by law to transfer ownership of the drug acquired by
13	the audited pharmacy;
14	(c) copies of supplier invoices in the
15	possession of the audited pharmacy; and
16	(d) reports required by any state board
17	or agency; and
18	(4) within five business days of request by
19	the audited pharmacy, the entity shall provide supporting
20	documentation provided to the entity by the audited pharmacy's
21	suppliers.
22	E. The provisions of this section may not be
23	waived, voided or nullified by contract.
24	F. As used in this section:
25	(1) "entity" means a managed care company,
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	1	insurance company or third-party payor, or representative of a
	2	managed care company, insurance company or third-party payor,
	3	or a pharmacy benefits manager or a subcontractor of a pharmacy
	4	benefits manager; and
	5	(2) "extrapolation" means a mathematical
	6	process or technique used to estimate audit results or findings
	7	for a larger batch or group of claims not reviewed."
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