SECOND REGULAR SESSION HOUSE COMMITTEE SUBSTITUTE FOR HOUSE BILL NO. 1332

94TH GENERAL ASSEMBLY

Reported from the Special Committee on Healthcare Transformation February 14, 2008 with recommendation that House Committee Substitute for House Bill No. 1332 Do Pass. Referred to the Committee on Rules pursuant to Rule 25(21)(f).

D. ADAM CRUMBLISS, Chief Clerk

3027L.05C

AN ACT

To amend chapter 338, RSMo, by adding thereto one new section relating to pharmacists and pharmacies.

Be it enacted by the General Assembly of the state of Missouri, as follows:

Section A. Chapter 338, RSMo, is amended by adding thereto one new section, to be 2 known as section 338.600, to read as follows:

338.600. 1. Notwithstanding any other provision of law to the contrary, when an audit of the records of a pharmacy licensed in this state is conducted by a managed care company, insurance company, third-party payor, the department of insurance, financial institutions and professional registration, the board of pharmacy, or any entity that represents such companies, groups, department, or board, such audit shall be conducted in accordance with the following: (1) The entity conducting the initial on-site audit shall provide the pharmacy with notice at least one week prior to conducting the initial on-site audit for each audit cycle;

9 (2) Any audit which involves clinical judgment shall be conducted by or in 10 consultation with a licensed pharmacist;

(3) Any clerical or recordkeeping error, such as a typographical error, scriveners
 error, or computer error, regarding a required document or record shall not in and of
 itself constitute fraud; except that, such claims shall be subject to recoupment or payment

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of any discovered underpayment. No such claim shall be subject to criminal penalties
without proof of intent to commit fraud;

(4) A pharmacy may use the records of a hospital, physician, or other authorized
 practitioner of the healing arts involving drugs or medicinal supplies written or
 transmitted by any means of communication for purposes of validating the pharmacy
 record with respect to orders or refills of a legend or narcotic drug;

(5) A finding of an overpayment or underpayment may be a projection based on the number of patients served and having a similar diagnosis or on the number of similar orders or refills for similar drugs; except that, recoupment of claims shall be based on the actual overpayment or underpayment unless the projection for overpayment or underpayment is part of a settlement as agreed to by the pharmacy;

(6) Retail, hospital, and mail order pharmacies shall be audited under the same
 standards and parameters as other pharmacies of the same class audited by the entity;

(7) A pharmacy shall be allowed at least thirty days following receipt of the
preliminary audit report in which to produce documentation to address any discrepancy
found during an audit;

30 (8) The period covered by the audit shall not exceed a two-year period beginning 31 two years prior to the initial date of the on-site portion of the audit. The audit shall only 32 review claims that, during the same audit period, were submitted to or adjudicated by the 33 managed care company, insurance company, third-party payor, the state of Missouri, or 34 any entity that represents such company or group conducting the audit;

35 (9) An audit shall not be initiated or scheduled during the first five business days
36 of any month due to the high volume of prescriptions filled during such time unless
37 otherwise consented to by the pharmacy;

(10) The preliminary audit report shall be delivered to the pharmacy within one
hundred twenty days after conclusion of the audit, with reasonable extensions permitted.
A final audit report shall be delivered to the pharmacy within six months of receipt by the
pharmacy of the preliminary audit report or final appeal, as provided for in subsection 3
of this section, whichever is later;

(11) Notwithstanding any other provision in this subsection, the entity conducting
the audit shall not use the accounting practice of extrapolation in calculating recoupments
or penalties for audits, except as otherwise authorized under subdivision (5) of this
subsection.

47 2. Recoupments of any disputed moneys shall only occur after final internal
48 disposition of the audit, including the appeals process set forth in subsection 4 of this
49 section.

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3. Each entity conducting an audit shall establish an appeals process, lasting no longer than six months, under which a licensed pharmacy may appeal an unfavorable preliminary audit report to the entity. If, following such appeal, the entity finds that an unfavorable audit report or any portion thereof is unsubstantiated, the entity shall dismiss the audit report or such portion without the necessity of any further proceedings.

- 4. Each entity conducting an audit shall provide a copy of the final audit report,
 after completion of any appeal process, to the plan sponsor.
- 57 5. This section shall not apply to any audit conducted as a part of an investigation 58 regarding alleged criminal wrongdoing, willful misrepresentation, or abuse.
- 596. Unless required by federal law, no contract entered into or renewed after the60effective date of this section shall contain audit criteria provisions that are more restrictive
- 61 than the audit criteria provisions contained in this section.

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