SECOND REGULAR SESSION [PERFECTED]

SENATE SUBSTITUTE FOR

SENATE COMMITTEE SUBSTITUTE FOR

SENATE BILLS NOS. 865 & 866

98TH GENERAL ASSEMBLY

INTRODUCED BY SENATOR SATER.

Offered February 24, 2016.

Senate Substitute adopted, February 24, 2016.

Taken up for Perfection February 24, 2016. Bill declared Perfected and Ordered Printed, as amended.

5458S.03P

ADRIANE D. CROUSE, Secretary.

AN ACT

To repeal sections 338.270, 338.347, and 354.535, RSMo, and to enact in lieu thereof seven new sections relating to pharmacy.

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

Section A. Sections 338.270, 338.347, and 354.535, RSMo, are repealed

- 2 and seven new sections enacted in lieu thereof, to be known as sections 338.075,
- 3 338.270, 338.347, 354.535, 376.379, 376.387, and 376.388, to read as follows:

338.075. 1. All licensees, registrants, and permit holders of the

- 2 board shall report to the board:
- 3 (1) Any final adverse action taken by another licensing state,
- 4 jurisdiction, or government agency against any license, permit, or
- 5 authorization held by the person or entity to practice or operate as a
- 6 pharmacist, intern pharmacist, pharmacy technician, pharmacy, drug
- 7 distributor, drug manufacturer, or drug outsourcing facility. For
- 8 purposes of this section, "adverse action" shall include, but is not
- 9 limited to, revocation, suspension, censure, probation, disciplinary
- 10 reprimand, or disciplinary restriction of a license, permit, or other
- 11 authorization or a voluntary surrender of such license, permit, or other
- 12 authorization in lieu of discipline or adverse action;
- 13 (2) Any surrender of a license or authorization to practice or
- 14 operate as a pharmacist, intern pharmacist, pharmacy technician,

EXPLANATION—Matter enclosed in bold-faced brackets [thus] in this bill is not enacted and is intended to be omitted in the law.

6

8

- pharmacy, drug distributor, drug manufacturer, or drug outsourcing facility while under disciplinary investigation by another licensing state, jurisdiction, or governmental agency, and;
- 18 (3) Any exclusion to participate in any state or federally funded 19 health care program such as Medicare, Medicaid, or MO HealthNet for 20 fraud, abuse, or submission of any false or fraudulent claim, payment, 21 or reimbursement request.
 - 2. Reports shall be submitted as provided by the board by rule.
- 3. The board shall promulgate rules to implement the provisions 23 of this section. Any rule or portion of a rule, as that term is defined in 2425 section 536.010 that is created under the authority delegated in this section shall become effective only if it complies with and is subject to 26 27all of the provisions of chapter 536, and, if applicable, section 28 536.028. This section and chapter 536 are nonseverable and if any of the powers vested with the general assembly pursuant to chapter 536, 30 to review, to delay the effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking 31 authority and any rule proposed or adopted after August 28, 2016, shall 32be invalid and void. 33
- 338.270. 1. Application blanks for renewal permits shall be mailed to each permittee on or before the first day of the month in which the permit expires and, if application for renewal of permit is not made before the first day of the following month, the existing permit, or renewal thereof, shall lapse and become null and void upon the last day of that month.
 - 2. The board shall not renew a nonresident pharmacy license if the renewal applicant does not hold a current pharmacy license or its equivalent in the state in which the nonresident pharmacy is located.
- 338.347. 1. Application blanks for renewal of license shall be mailed to each licensee on or before the first day of the month in which the license expires and, if application for renewal of license with required fee is not made before the first day of the following month, the existing license, or renewal thereof, shall lapse and become null and void upon the last day of that month.
- 2. The board shall not renew an out-of-state wholesale drug distributor, out-of-state pharmacy distributor, or drug distributor license or registration if the renewal applicant does not hold a current distributor license or its equivalent in the state or jurisdiction in which the distribution facility is located, or, if a drug distributor registrant,

25

26

27

28

33

the entity is not authorized and in good standing to operate as a drug manufacturer with the Food and Drug Administration or within the

13 state or jurisdiction where the facility is located.

354.535. 1. If a pharmacy, operated by or contracted with by a health 2 maintenance organization, is closed or is unable to provide health care services 3 to an enrollee in an emergency, a pharmacist may take an assignment of such 4 enrollee's right to reimbursement, if the policy or contract provides for such 5 reimbursement, for those goods or services provided to an enrollee of a health 6 maintenance organization. No health maintenance organization shall refuse to 7 pay the pharmacist any payment due the enrollee under the terms of the policy 8 or contract.

- 2. No health maintenance organization, conducting business in the state of Missouri, shall contract with a pharmacy, pharmacy distributor or wholesale drug distributor, nonresident or otherwise, unless such pharmacy or distributor has been granted a permit or license from the Missouri board of pharmacy to operate in this state.
- 3. Every health maintenance organization shall apply the same 14 coinsurance, co-payment and deductible factors to all drug prescriptions filled by 15 a pharmacy provider who participates in the health maintenance organization's 16 network if the provider meets the contract's explicit product cost determination. 17 If any such contract is rejected by any pharmacy provider, the health 18 maintenance organization may offer other contracts necessary to comply with any 19 20 network adequacy provisions of this act. However, nothing in this section shall be construed to prohibit the health maintenance organization from applying 2122 different coinsurance, co-payment and deductible factors between generic and 23 brand name drugs.
 - 4. If the co-payment applied by a health maintenance organization exceeds the usual and customary retail price of the prescription drug, enrollees shall only be required to pay the usual and customary retail price of the prescription drug, and no further charge to the enrollee or plan sponsor shall be incurred on such prescription.
- 5. Health maintenance organizations shall not set a limit on the quantity of drugs which an enrollee may obtain at any one time with a prescription, unless such limit is applied uniformly to all pharmacy providers in the health maintenance organization's network.
 - [5.] **6.** Health maintenance organizations shall not insist or mandate any

physician or other licensed health care practitioner to change an enrollee's 35 maintenance drug unless the provider and enrollee agree to such change. For the purposes of this provision, a maintenance drug shall mean a drug prescribed by 36 a practitioner who is licensed to prescribe drugs, used to treat a medical condition 37for a period greater than thirty days. Violations of this provision shall be subject 38 to the penalties provided in section 354.444. Notwithstanding other provisions 39 of law to the contrary, health maintenance organizations that change an 40 enrollee's maintenance drug without the consent of the provider and enrollee 41 42 shall be liable for any damages resulting from such change. Nothing in this 43 subsection, however, shall apply to the dispensing of generically equivalent products for prescribed brand name maintenance drugs as set forth in section 45 338.056.

- 376.379. 1. A health carrier or managed care plan offering a health benefit plan in this state that provides prescription drug coverage shall offer, as part of the plan, medication synchronization services developed by the health carrier or managed care plan that allow for the alignment of refill dates for an enrollee's prescription drugs that are covered benefits.
- 7 2. Under its medication synchronization services, a health 8 carrier or managed care plan shall:
- 9 (1) Not charge an amount in excess of the otherwise applicable 10 copayment amount under the health benefit plan for dispensing a 11 prescription drug in a quantity that is less than the prescribed amount 12 if:
- (a) The pharmacy dispenses the prescription drug in accordance
 with the medication synchronization services offered under the health
 benefit plan; and
 - (b) A participating provider dispenses the prescription drug;
- 17 (2) Provide a full dispensing fee to the pharmacy that dispenses 18 the prescription drug to the covered person.
- 3. For the purposes of this section the terms "health carrier", "managed care plan", "health benefit plan", "enrollee", and "participating provider" shall have the same meaning as defined in section 376.1350.

376.387. If the co-payment for prescription drugs applied by a 2 health insurer or health carrier, as defined in section 376.1350, exceeds 3 the usual and customary retail price of the prescription drug, enrollees 4 shall only be required to pay the usual and customary retail price of

30

31 32

33

34

the prescription drug, and no further charge to the enrollee or plan sponsor shall be incurred on such prescription.

376.388. 1. As used in this section, unless the context requires 2 otherwise, the following terms shall mean:

- (1) "Contracted pharmacy" or "pharmacy", a pharmacy located in
 Missouri participating in the network of a pharmacy benefits manager
 through a direct or indirect contract;
- (2) "Health carrier", an entity subject to the insurance laws and regulations of this state that contracts or offers to contract to provide, deliver, arrange for, pay for, or reimburse any of the costs of health care services, including a sickness and accident insurance company, a health maintenance organization, a nonprofit hospital and health service corporation, or any other entity providing a plan of health insurance, health benefits, or health services, except that such plan shall not include any coverage pursuant to a liability insurance policy, workers' compensation insurance policy, or medical payments insurance issued as a supplement to a liability policy;
- 16 (3) "Maximum allowable cost", the per unit amount that a 17 pharmacy benefits manager reimburses a pharmacist for a prescription 18 drug, excluding a dispensing or professional fee;
- 19 (4) "Maximum allowable cost list" or "MAC list", a listing of drug 20 products that meet the standard described in this section;
 - (5) "Pharmacy", as such term is defined in chapter 338;
- 22 (6) "Pharmacy benefits manager", an entity that contracts with 23 pharmacies on behalf of health carriers or any health plan sponsored 24 by the state or a political subdivision of the state.
- 2. Upon each contract execution or renewal between a pharmacy benefits manager and a pharmacy or between a pharmacy benefits manager and a pharmacy's contracting representative or agent, such as a pharmacy services administrative organization, a pharmacy benefits manager shall, with respect to such contract or renewal:
 - (1) Include in such contract or renewal the sources utilized to determine maximum allowable cost and update such pricing information at least every seven days; and
 - (2) Maintain a procedure to eliminate products from the maximum allowable cost list of drugs subject to such pricing or modify maximum allowable cost pricing at least every seven days if such drugs

49

52

53

56

58

59

60

61

62

63

64 65

67

68

36 do not meet the standards and requirements of this section in order to 37 remain consistent with pricing changes in the marketplace.

- 38 3. A pharmacy benefits manager shall reimburse pharmacies for 39 drugs subject to maximum allowable cost pricing which has been 40 updated to reflect market pricing at least every seven days as set forth 41 in subdivision (1) of subsection 2 of this section.
- 42 4. A pharmacy benefits manager shall not place a drug on a 43 maximum allowable cost list unless there are at least two 44 therapeutically equivalent multi-source generic drugs, or at least one 45 generic drug available from at least one manufacturer, generally 46 available for purchase by network pharmacies from national or 47 regional wholesalers.
 - 5. All contracts between a pharmacy benefits manager and a contracted pharmacy or between a pharmacy benefits manager and a pharmacy's contracting representative or agent, such as a pharmacy services administrative organization, shall include a process to internally appeal, investigate, and resolve disputes regarding maximum allowable cost pricing. The process shall include the following:
- 54 (1) The right to appeal shall be limited to fourteen calendar days 55 following the reimbursement of the initial claim; and
 - (2) A requirement that the pharmacy benefits manager shall respond to an appeal described in this subsection no later than fourteen calendar days after the date the appeal was received by such pharmacy benefits manager.
 - 6. For appeals that are denied, the pharmacy benefits manager shall provide the reason for the denial and identify the national drug code of a drug product that may be purchased by contracted pharmacies at a price at or below the maximum allowable cost, and when applicable, may be substituted lawfully.
- 7. If the appeal is successful, the pharmacy benefits manager shall:
 - (1) Adjust the maximum allowable cost price that is the subject of the appeal effective on the day after the date the appeal is decided;
- 69 (2) Apply the adjusted maximum allowable cost price to all 70 similarly situated pharmacies as determined by the pharmacy benefits 71 manager; and
- 72 (3) Allow the pharmacy that succeeded in the appeal to reverse

73 and rebill the pharmacy benefits claim giving rise to the appeal.

- 8. Appeals shall be upheld if:
- 75 (1) The pharmacy being reimbursed for the drug subject to the 76 maximum allowable cost pricing in question was not reimbursed as 77 required in subsection 3 of this section; or
- 78 (2) The drug subject to the maximum allowable cost pricing in 79 question does not meet the requirements set forth in subsection 4 of 80 this section.

Unofficial

Bill

Copy