

SECOND EXTRAORDINARY SESSION  
**HOUSE BILL NO. 8**

**99TH GENERAL ASSEMBLY**

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INTRODUCED BY REPRESENTATIVE QUADE.

2455H.011

D. ADAM CRUMBLISS, Chief Clerk

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**AN ACT**

To repeal section 338.010, RSMo, and section 338.010 as truly agreed to and finally passed by senate bill no. 501, ninety-ninth general assembly, first regular session, and to enact in lieu thereof three new sections relating to contraceptives.

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

Section A. Section 338.010, RSMo, and section 338.010 as truly agreed to and finally passed by senate bill no. 501, ninety-ninth general assembly, first regular session, are repealed and three new sections enacted in lieu thereof, to be known as sections 338.010, 338.660, and 376.1240, to read as follows:

338.010. 1. The "practice of pharmacy" means the interpretation, implementation, and evaluation of medical prescription orders, including any legend drugs under 21 U.S.C. Section 353; receipt, transmission, or handling of such orders or facilitating the dispensing of such orders; the designing, initiating, implementing, and monitoring of a medication therapeutic plan as defined by the prescription order so long as the prescription order is specific to each patient for care by a pharmacist; the compounding, dispensing, labeling, and administration of drugs and devices pursuant to medical prescription orders and administration of viral influenza, pneumonia, shingles, hepatitis A, hepatitis B, diphtheria, tetanus, pertussis, and meningitis vaccines by written protocol authorized by a physician for persons twelve years of age or older as authorized by rule or the administration of pneumonia, shingles, hepatitis A, hepatitis B, diphtheria, tetanus, pertussis, and meningitis vaccines by written protocol authorized by a physician for a specific patient as authorized by rule; the participation in drug selection according to state law and participation in drug utilization reviews; the proper and safe storage of drugs and devices and the maintenance of proper records thereof; consultation with patients and other health care

EXPLANATION — Matter enclosed in bold-faced brackets [thus] in the above bill is not enacted and is intended to be omitted from the law. Matter in **bold-face** type in the above bill is proposed language.

15 practitioners, and veterinarians and their clients about legend drugs, about the safe and effective  
16 use of drugs and devices; **the prescribing and dispensing of self-administered oral hormonal**  
17 **contraceptives under section 338.660**; and the offering or performing of those acts, services,  
18 operations, or transactions necessary in the conduct, operation, management and control of a  
19 pharmacy. No person shall engage in the practice of pharmacy unless he is licensed under the  
20 provisions of this chapter. This chapter shall not be construed to prohibit the use of auxiliary  
21 personnel under the direct supervision of a pharmacist from assisting the pharmacist in any of  
22 his or her duties. This assistance in no way is intended to relieve the pharmacist from his or her  
23 responsibilities for compliance with this chapter and he or she will be responsible for the actions  
24 of the auxiliary personnel acting in his or her assistance. This chapter shall also not be construed  
25 to prohibit or interfere with any legally registered practitioner of medicine, dentistry, or podiatry,  
26 or veterinary medicine only for use in animals, or the practice of optometry in accordance with  
27 and as provided in sections 195.070 and 336.220 in the compounding, administering,  
28 prescribing, or dispensing of his or her own prescriptions.

29         2. Any pharmacist who accepts a prescription order for a medication therapeutic plan  
30 shall have a written protocol from the physician who refers the patient for medication therapy  
31 services. The written protocol and the prescription order for a medication therapeutic plan shall  
32 come from the physician only, and shall not come from a nurse engaged in a collaborative  
33 practice arrangement under section 334.104, or from a physician assistant engaged in a  
34 supervision agreement under section 334.735.

35         3. Nothing in this section shall be construed as to prevent any person, firm or corporation  
36 from owning a pharmacy regulated by sections 338.210 to 338.315, provided that a licensed  
37 pharmacist is in charge of such pharmacy.

38         4. Nothing in this section shall be construed to apply to or interfere with the sale of  
39 nonprescription drugs and the ordinary household remedies and such drugs or medicines as are  
40 normally sold by those engaged in the sale of general merchandise.

41         5. No health carrier as defined in chapter 376 shall require any physician with which they  
42 contract to enter into a written protocol with a pharmacist for medication therapeutic services.

43         6. This section shall not be construed to allow a pharmacist to diagnose or independently  
44 prescribe pharmaceuticals.

45         7. The state board of registration for the healing arts, under section 334.125, and the state  
46 board of pharmacy, under section 338.140, shall jointly promulgate rules regulating the use of  
47 protocols for prescription orders for medication therapy services and administration of viral  
48 influenza vaccines. Such rules shall require protocols to include provisions allowing for timely  
49 communication between the pharmacist and the referring physician, and any other patient  
50 protection provisions deemed appropriate by both boards. In order to take effect, such rules shall

51 be approved by a majority vote of a quorum of each board. Neither board shall separately  
52 promulgate rules regulating the use of protocols for prescription orders for medication therapy  
53 services and administration of viral influenza vaccines. Any rule or portion of a rule, as that term  
54 is defined in section 536.010, that is created under the authority delegated in this section shall  
55 become effective only if it complies with and is subject to all of the provisions of chapter 536  
56 and, if applicable, section 536.028. This section and chapter 536 are nonseverable and if any of  
57 the powers vested with the general assembly pursuant to chapter 536 to review, to delay the  
58 effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the  
59 grant of rulemaking authority and any rule proposed or adopted after August 28, 2007, shall be  
60 invalid and void.

61 8. The state board of pharmacy may grant a certificate of medication therapeutic plan  
62 authority to a licensed pharmacist who submits proof of successful completion of a  
63 board-approved course of academic clinical study beyond a bachelor of science in pharmacy,  
64 including but not limited to clinical assessment skills, from a nationally accredited college or  
65 university, or a certification of equivalence issued by a nationally recognized professional  
66 organization and approved by the board of pharmacy.

67 9. Any pharmacist who has received a certificate of medication therapeutic plan authority  
68 may engage in the designing, initiating, implementing, and monitoring of a medication  
69 therapeutic plan as defined by a prescription order from a physician that is specific to each  
70 patient for care by a pharmacist.

71 10. Nothing in this section shall be construed to allow a pharmacist to make a therapeutic  
72 substitution of a pharmaceutical prescribed by a physician unless authorized by the written  
73 protocol or the physician's prescription order.

74 11. "Veterinarian", "doctor of veterinary medicine", "practitioner of veterinary  
75 medicine", "DVM", "VMD", "BVSe", "BVMS", "BSe (Vet Science)", "VMB", "MRCVS", or  
76 an equivalent title means a person who has received a doctor's degree in veterinary medicine  
77 from an accredited school of veterinary medicine or holds an Educational Commission for  
78 Foreign Veterinary Graduates (EDFVG) certificate issued by the American Veterinary Medical  
79 Association (AVMA).

80 12. In addition to other requirements established by the joint promulgation of rules by  
81 the board of pharmacy and the state board of registration for the healing arts:

82 (1) A pharmacist shall administer vaccines **by protocol** in accordance with treatment  
83 guidelines established by the Centers for Disease Control and Prevention (CDC);

84 (2) A pharmacist who is administering a vaccine shall request a patient to remain in the  
85 pharmacy a safe amount of time after administering the vaccine to observe any adverse reactions.  
86 Such pharmacist shall have adopted emergency treatment protocols;

87 (3) In addition to other requirements by the board, a pharmacist shall receive additional  
88 training as required by the board and evidenced by receiving a certificate from the board upon  
89 completion, and shall display the certification in his or her pharmacy where vaccines are  
90 delivered.

91 13. A pharmacist shall provide a written report within fourteen days of administration  
92 of a vaccine to the patient's primary health care provider, if provided by the patient, containing:

- 93 (1) The identity of the patient;
- 94 (2) The identity of the vaccine or vaccines administered;
- 95 (3) The route of administration;
- 96 (4) The anatomic site of the administration;
- 97 (5) The dose administered; and
- 98 (6) The date of administration.

**338.660. 1. For purposes of this chapter, “self-administered oral hormonal  
2 contraceptive” shall mean a drug composed of a combination of hormones that is approved  
3 by the Food and Drug Administration to prevent pregnancy and that the patient to whom  
4 the drug is prescribed may take orally.**

**5 2. A pharmacist may prescribe and dispense self-administered oral hormonal  
6 contraceptives to a person who is:**

**7 (1) Eighteen years of age or older, regardless of whether the person has evidence  
8 of a previous prescription from a primary care practitioner or women’s health care  
9 practitioner for a self-administered oral hormonal contraceptive; or**

**10 (2) Under eighteen years of age, if the person has evidence of a previous  
11 prescription from a primary care practitioner or women’s health care practitioner for a  
12 self-administered oral hormonal contraceptive.**

**13 3. The board of pharmacy shall adopt rules, in consultation with the board of  
14 registration for the healing arts, board of nursing, and department of health and senior  
15 services, and in consideration of guidelines established by the American Congress of  
16 Obstetricians and Gynecologists, to establish standard procedures for the prescribing of  
17 self-administered oral hormonal contraceptives by pharmacists. The board of pharmacy  
18 shall adopt rules and regulations to implement the provisions of this section. Any rule or  
19 portion of a rule, as that term is defined in section 536.010, that is created under the  
20 authority delegated in this section shall become effective only if it complies with and is  
21 subject to all of the provisions of chapter 536 and, if applicable, section 536.028. This  
22 section and chapter 536 are nonseverable, and if any of the powers vested with the general  
23 assembly pursuant to chapter 536 to review, to delay the effective date, or to disapprove  
24 and annul a rule are subsequently held unconstitutional, then the grant of rulemaking**

25 authority and any rule proposed or adopted after the effective date of this section shall be  
26 invalid and void.

27 4. The rules adopted under this section shall require a pharmacist to:

28 (1) Complete a training program approved by the board of pharmacy that is  
29 related to prescribing self-administered oral hormonal contraceptives;

30 (2) Provide a self-screening risk assessment tool that the patient shall use prior to  
31 the pharmacist's prescribing the self-administered oral hormonal contraceptive;

32 (3) Refer the patient to the patient's primary care practitioner or women's health  
33 care practitioner upon prescribing and dispensing the self-administered oral hormonal  
34 contraceptive;

35 (4) Provide the patient with a written record of the self-administered oral hormonal  
36 contraceptive prescribed and dispensed and advise the patient to consult with a primary  
37 care practitioner or women's health care practitioner; and

38 (5) Dispense the self-administered oral hormonal contraceptive to the patient as  
39 soon as practicable after the pharmacist issues the prescription.

40 5. The rules adopted under this section shall prohibit a pharmacist from requiring  
41 a patient to schedule an appointment with the pharmacist for the prescribing or dispensing  
42 of a self-administered oral hormonal contraceptive.

43 6. All state and federal laws governing insurance coverage of contraceptive drugs,  
44 devices, products, and services shall apply to self-administered oral hormonal  
45 contraceptives prescribed by a pharmacist under this section.

376.1240. 1. For purposes of this section, the terms "health carrier" and "health  
2 benefit plan" shall have the same meaning as defined in section 376.1350. The term  
3 "prescription contraceptive" shall mean a drug or device that requires a prescription and  
4 is approved by the Food and Drug Administration to prevent pregnancy.

5 2. Each health carrier or health benefit plan that offers or issues health benefit  
6 plans which are delivered, issued for delivery, continued, or renewed in this state on or  
7 after January 1, 2018, and that provides coverage for prescription contraceptives shall  
8 provide coverage to reimburse a health care provider or dispensing entity for a dispensing  
9 of prescription contraceptives intended to last for a:

10 (1) Three-month period for the first dispensing of the prescription contraceptive  
11 to an insured; and

12 (2) Three-month period for subsequent dispensations of the same contraceptive to  
13 the insured regardless of whether the insured was enrolled in the health benefit plan or  
14 policy at the time of the first dispensing.

15           **3. The coverage required by this section shall not be subject to any greater**  
16 **deductible or co-payment than other similar health care services provided by the health**  
17 **benefit plan.**

18           **4. The provisions of this section shall not apply to a supplemental insurance policy**  
19 **including a life care contract, accident-only policy, specified disease policy, hospital policy**  
20 **providing a fixed daily benefit only, Medicare supplement policy, long-term care policy,**  
21 **short-term major medical policies of six months' or less duration, or any other**  
22 **supplemental policy as determined by the director of the department of insurance,**  
23 **financial institutions and professional registration.**

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3 ~~implementation, and evaluation of medical prescription orders, including any~~  
4 ~~legend drugs under 21 U.S.C. Section 353; receipt, transmission, or handling of~~  
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7 ~~prescription order so long as the prescription order is specific to each patient for~~  
8 ~~care by a pharmacist; the compounding, dispensing, labeling, and administration~~  
9 ~~of drugs and devices pursuant to medical prescription orders and administration~~  
10 ~~of viral influenza, pneumonia, shingles, hepatitis A, hepatitis B, diphtheria,~~  
11 ~~tetanus, pertussis, and meningitis vaccines by written protocol authorized by a~~  
12 ~~physician for persons twelve years of age or older as authorized by rule or the~~  
13 ~~administration of pneumonia, shingles, hepatitis A, hepatitis B, diphtheria,~~  
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15 ~~physician for a specific patient as authorized by rule; the participation in drug~~  
16 ~~selection according to state law and participation in drug utilization reviews; the~~  
17 ~~proper and safe storage of drugs and devices and the maintenance of proper~~  
18 ~~records thereof; consultation with patients and other health care practitioners, and~~  
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20 ~~of drugs and devices, and the offering or performing of those acts, services,~~  
21 ~~operations, or transactions necessary in the conduct, operation, management and~~  
22 ~~control of a pharmacy. No person shall engage in the practice of pharmacy unless~~  
23 ~~he is licensed under the provisions of this chapter. This chapter shall not be~~  
24 ~~construed to prohibit the use of auxiliary personnel under the direct supervision~~  
25 ~~of a pharmacist from assisting the pharmacist in any of his or her duties. This~~  
26 ~~assistance in no way is intended to relieve the pharmacist from his or her~~  
27 ~~responsibilities for compliance with this chapter and he or she will be responsible~~  
28 ~~for the actions of the auxiliary personnel acting in his or her assistance. This~~  
29 ~~chapter shall also not be construed to prohibit or interfere with any legally~~  
30 ~~registered practitioner of medicine, dentistry, or podiatry, or veterinary medicine~~  
31 ~~only for use in animals, or the practice of optometry in accordance with and as~~  
32 ~~provided in sections 195.070 and 336.220 in the compounding, administering,~~  
~~prescribing, or dispensing of his or her own prescriptions.~~

33 ~~2. Any pharmacist who accepts a prescription order for a medication~~  
34 ~~therapeutic plan shall have a written protocol from the physician who refers the~~  
35 ~~patient for medication therapy services. The written protocol and the prescription~~  
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37 ~~shall not come from a nurse engaged in a collaborative practice arrangement~~  
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39 ~~agreement under section 334.735.~~

40 ~~3. Nothing in this section shall be construed as to prevent any person,~~  
41 ~~firm or corporation from owning a pharmacy regulated by sections 338.210 to~~  
42 ~~338.315, provided that a licensed pharmacist is in charge of such pharmacy.~~

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44 ~~the sale of nonprescription drugs and the ordinary household remedies and such~~  
45 ~~drugs or medicines as are normally sold by those engaged in the sale of general~~  
46 ~~merchandise.~~

47 ~~5. No health carrier as defined in chapter 376 shall require any physician~~  
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66 ~~are nonseverable and if any of the powers vested with the general assembly~~  
67 ~~pursuant to chapter 536 to review, to delay the effective date, or to disapprove~~  
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69 ~~rulemaking authority and any rule proposed or adopted after August 28, 2007,~~  
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71 ~~8. The state board of pharmacy may grant a certificate of medication~~  
72 ~~therapeutic plan authority to a licensed pharmacist who submits proof of~~  
73 ~~successful completion of a board-approved course of academic clinical study~~  
74 ~~beyond a bachelor of science in pharmacy, including but not limited to clinical~~  
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77 organization and approved by the board of pharmacy.

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80 and monitoring of a medication therapeutic plan as defined by a prescription  
81 order from a physician that is specific to each patient for care by a pharmacist.~~

82 ~~10. Nothing in this section shall be construed to allow a pharmacist to  
83 make a therapeutic substitution of a pharmaceutical prescribed by a physician  
84 unless authorized by the written protocol or the physician's prescription order.~~

85 ~~11. "Veterinarian", "doctor of veterinary medicine", "practitioner of  
86 veterinary medicine", "DVM", "VMD", "BVSc", "BVMS", "BSc (Vet Science)",  
87 "VMB", "MRCVS", or an equivalent title means a person who has received a  
88 doctor's degree in veterinary medicine from an accredited school of veterinary  
89 medicine or holds an Educational Commission for Foreign Veterinary Graduates  
90 (EDFVG) certificate issued by the American Veterinary Medical Association  
91 (AVMA).~~

92 ~~12. In addition to other requirements established by the joint  
93 promulgation of rules by the board of pharmacy and the state board of registration  
94 for the healing arts:~~

95 ~~(1) A pharmacist shall administer vaccines by protocol in accordance  
96 with treatment guidelines established by the Centers for Disease Control and  
97 Prevention (CDC);~~

98 ~~(2) A pharmacist who is administering a vaccine shall request a patient  
99 to remain in the pharmacy a safe amount of time after administering the vaccine  
100 to observe any adverse reactions. Such pharmacist shall have adopted emergency  
101 treatment protocols;~~

102 ~~(3) In addition to other requirements by the board, a pharmacist shall  
103 receive additional training as required by the board and evidenced by receiving  
104 a certificate from the board upon completion, and shall display the certification  
105 in his or her pharmacy where vaccines are delivered.~~

106 ~~13. A pharmacist shall provide a written report within fourteen days of  
107 administration of a vaccine to the patient's primary health care provider, if  
108 provided by the patient, containing:~~

109 ~~(1) The identity of the patient;~~

110 ~~(2) The identity of the vaccine or vaccines administered;~~

111 ~~(3) The route of administration;~~

112 ~~(4) The anatomic site of the administration;~~

113 ~~(5) The dose administered; and~~

114 ~~(6) The date of administration.]~~

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