

FIRST REGULAR SESSION

SENATE BILL NO. 353

100TH GENERAL ASSEMBLY

INTRODUCED BY SENATOR EMERY.

Read 1st time February 6, 2019, and ordered printed.

ADRIANE D. CROUSE, Secretary.

1412S.01I

AN ACT

To repeal section 195.080, RSMo, and to enact in lieu thereof one new section relating to opioid controlled substance prescriptions.

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

Section A. Section 195.080, RSMo, is repealed and one new section
2 enacted in lieu thereof, to be known as section 195.080, to read as follows:

195.080. 1. Except as otherwise provided in this chapter and chapter 579,
2 this chapter and chapter 579 shall not apply to the following cases: prescribing,
3 administering, dispensing or selling at retail of liniments, ointments, and other
4 preparations that are susceptible of external use only and that contain controlled
5 substances in such combinations of drugs as to prevent the drugs from being
6 readily extracted from such liniments, ointments, or preparations, except that
7 this chapter and chapter 579 shall apply to all liniments, ointments, and other
8 preparations that contain coca leaves in any quantity or combination.

9 2. Unless otherwise provided in sections 334.037, 334.104, and 334.747,
10 a practitioner, other than a veterinarian, shall not issue an initial prescription
11 for more than a seven-day supply of any opioid controlled substance upon the
12 initial consultation and treatment of a patient for acute pain. Upon any
13 subsequent consultation for the same pain, the practitioner may issue any
14 appropriate renewal, refill, or new prescription in compliance with the general
15 provisions of this chapter and chapter 579. Prior to issuing an initial prescription
16 for an opioid controlled substance, a practitioner shall consult with the patient
17 regarding the quantity of the opioid and the patient's option to fill the
18 prescription in a lesser quantity and shall inform the patient of the risks
19 associated with the opioid prescribed, **as provided for in subsection 3 of this**

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

20 **section.** If, in the professional medical judgment of the practitioner, more than
21 a seven-day supply is required to treat the patient's acute pain, the practitioner
22 may issue a prescription for the quantity needed to treat the patient; provided,
23 that the practitioner shall document in the patient's medical record the condition
24 triggering the necessity for more than a seven-day supply and that a nonopioid
25 alternative was not appropriate to address the patient's condition. The provisions
26 of this subsection shall not apply to prescriptions for opioid controlled substances
27 for a patient who is currently undergoing treatment for cancer, is receiving
28 hospice care from a hospice certified under chapter 197 or palliative care, is a
29 resident of a long-term care facility licensed under chapter 198, or is receiving
30 treatment for substance abuse or opioid dependence.

31 **3. Prior to issuing an opioid controlled substance under**
32 **subsection 2 of this section and prior to issuing a third prescription of**
33 **such opioid in a course of treatment, the practitioner shall consult with**
34 **the patient, or the patient's parent or guardian if the patient is under**
35 **eighteen years of age and unemancipated, regarding the risks**
36 **associated with the opioid prescribed, including, but not limited to, the**
37 **following:**

38 **(1) The risks of addiction and overdose associated with opioid**
39 **drugs and the dangers of taking opioid drugs with alcohol,**
40 **benzodiazepines, and other central nervous system depressants;**

41 **(2) The reasons why the prescription is necessary;**

42 **(3) Alternative treatments that may be available; and**

43 **(4) The risks associated with the opioid being prescribed;**
44 **specifically, that opioids are highly addictive, even when taken as**
45 **prescribed, that there is a risk of developing a physical or**
46 **psychological dependence on the opioid, and that the risks of taking**
47 **more opioids than prescribed, or mixing sedatives, benzodiazepines, or**
48 **alcohol with opioids, can result in fatal respiratory depression.**

49 **The practitioner shall document in the patient's medical record that**
50 **the patient, or the patient's parent or guardian, has discussed with the**
51 **practitioner the risks of developing a physical or psychological**
52 **dependence on the opioid and alternative treatments that may be**
53 **available. The provisions of this subsection shall not apply to**
54 **prescriptions for opioid controlled substances for a patient who is**
55 **currently undergoing treatment for cancer, is receiving hospice care**
56 **from a hospice certified under chapter 197 or palliative care, is a**

57 **resident of a long-term care facility licensed under chapter 198, or is**
58 **receiving treatment for substance abuse or opioid dependence. A**
59 **practitioner who violates the provisions of this subsection may be**
60 **subject to disciplinary action by his or her respective professional**
61 **licensing board or action by the department under section 195.040.**

62 4. A pharmacist or pharmacy shall not be subject to disciplinary action
63 or other civil or criminal liability for dispensing or refusing to dispense
64 medication in good faith pursuant to an otherwise valid prescription that exceeds
65 the prescribing limits established by subsection 2 of this section.

66 [4.] 5. Unless otherwise provided in this section, the quantity of Schedule
67 II controlled substances prescribed or dispensed at any one time shall be limited
68 to a thirty-day supply. The quantity of Schedule III, IV or V controlled
69 substances prescribed or dispensed at any one time shall be limited to a
70 ninety-day supply and shall be prescribed and dispensed in compliance with the
71 general provisions of this chapter and chapter 579. The supply limitations
72 provided in this subsection may be increased up to three months if the physician
73 describes on the prescription form or indicates via telephone, fax, or electronic
74 communication to the pharmacy to be entered on or attached to the prescription
75 form the medical reason for requiring the larger supply. The supply limitations
76 provided in this subsection shall not apply if:

77 (1) The prescription is issued by a practitioner located in another state
78 according to and in compliance with the applicable laws of that state and the
79 United States and dispensed to a patient located in another state; or

80 (2) The prescription is dispensed directly to a member of the United
81 States Armed Forces serving outside the United States.

82 [5.] 6. The partial filling of a prescription for a Schedule II substance is
83 permissible as defined by regulation by the department of health and senior
84 services.

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