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H.723

Introduced by Representatives Sullivan of Dorset, Batchelor of Derby,
Brennan of Colchester, Buckholz of Hartford, Canfield of
Fair Haven, Cupoli of Rutland City, Dunn of Essex, Fagan of
Rutland City, Gannon of Wilmington, Higley of Lowell,
Joseph of North Hero, Juskiewicz of Cambridge, LaClair of
Barre Town, Lawrence of Lyndon, Lewis of Berlin, Martel of
Waterford, McCoy of Poultney, McFaun of Barre Town,
Morrissey of Bennington, Nolan of Morristown, Noyes of
Wolcott, Savage of Swanton, Shaw of Pittsford, Smith of
New Haven, and Strong of Albany

Referred to Committee on

Date:

Subject: Health; opioids; Vermont Prescription Monitoring System; health care
professionals

Statement of purpose of bill as introduced: This bill proposes to specify
statutory limits on the prescription of opioids and create a private right of
action for injuries caused by prescriptions made in violation of those limits.

An act relating to limits on the prescription of opioids and creating a private
right of action for prescriptions in excess of those limits

1 It is hereby enacted by the General Assembly of the State of Vermont:

2 Sec. 1. 18 V.S.A. chapter 84, subchapter 4 is added to read:

3 Subchapter 4. Prescription of Opioids; Right of Action

4 § 4271. DEFINITIONS

5 As used in this subchapter:

6 (1) “Acute pain” means pain lasting fewer than 90 days that is a normal
7 and predicted physiological response to a traumatic injury, surgical procedure,
8 or specific disease.

9 (2) “Affiliate or employer of a prescriber” means any hospital, physician
10 practice group, or other health care facility or entity with which the prescriber
11 has an employment, ownership, or contractual relationship and that has, with
12 respect to any individual patient for whom the prescriber has prescribed a
13 controlled substance to treat acute or chronic pain, any professional or medical
14 responsibility.

15 (3) “Chronic pain” means pain caused by one or more diseases or
16 abnormal conditions and that continues for 90 days or longer.

17 (4) “Controlled substance” means a drug or other substance, or
18 immediate precursor, included in schedule I, II, III, IV, or V of the federal
19 Controlled Substances Act, 21 U.S.C. § 801 et seq.

20 (5) “Initial prescription for acute pain” means a prescriber’s first order
21 of a controlled substance as a palliative treatment for a patient who is

1 experiencing pain based on the occurrence of a traumatic injury or surgical
2 procedure or upon the recent onset of pain naturally caused by a specific
3 disease.

4 (6) “Morphine milligram equivalent” or “MME” means a value assigned
5 to opioids to enable comparison of relative potency using morphine as the
6 standard.

7 (7) “Prescriber” means a licensed health care professional with the
8 authority to prescribe controlled substances.

9 (8) “Subsequent or additional prescription for acute pain” means a
10 prescriber’s second and subsequent orders of a controlled substance as a
11 palliative treatment for a patient suffering acute pain for an event or condition
12 that caused any prescriber to write an initial prescription for acute pain.

13 § 4272. PRESCRIPTION OF OPIOIDS FOR PAIN

14 (a) A prescriber shall not:

15 (1) except in an inpatient hospital or a skilled or intermediate care
16 nursing facility setting, make an initial prescription of an opioid controlled
17 substance for acute pain having an average daily MME of 50 for a period
18 exceeding seven days for a patient 18 years of age or older or an average daily
19 MME of 24 for a period exceeding three days for a patient under 18 years
20 of age;

1 (2) in an inpatient hospital or a skilled or intermediate care nursing
2 facility setting, make an initial prescription of an opioid controlled substance
3 for acute pain for a period exceeding 10 days for a patient 18 years of age or
4 older or for a period exceeding five days for a patient under 18 years of age;

5 (3) except in an inpatient hospital or a skilled or intermediate care
6 nursing facility setting, for pain associated with significant or severe trauma,
7 complex surgical interventions, or postoperative complications, make a
8 subsequent or additional prescription of an opioid controlled substance for
9 acute pain or in an amount or duration that exceeds the prescription durations
10 and average daily MME opioid limits for adults and children set forth in rules
11 adopted by the Commissioner of Health in accordance with subsection 4289(e)
12 of this title;

13 (4) write a prescription for an opioid controlled substance as a palliative
14 treatment for chronic pain except pursuant to and in compliance with a written
15 plan that:

16 (A) identifies the diagnosis or diagnoses supporting the use of
17 opioids for chronic pain;

18 (B) requires periodic follow-up visits and evaluations at least once
19 every 90 days regarding the continued need for opioid use;

20 (C) in the case of a prescription for an opioid controlled substance
21 with an average daily MME equal to or exceeding 90, requires a co-

1 prescription for naloxone as well as the preparation of an initial written
2 analysis and subsequent 90-day written reanalyses of all relevant
3 considerations of opioid abuse and addiction risks, including evidence of any
4 aberrant behaviors, early refills of opioid controlled substance prescriptions,
5 and other known risks associated with misuse, abuse, diversion, addiction, and
6 overdose; or

7 (5) write a prescription for an opioid controlled substance as a palliative
8 treatment for chronic or acute pain, or write a prescription for an extended-
9 release hydrocodone or oxycodone in a non-abuse-deterrent formulation,
10 without first querying the Vermont Prescription Monitoring System.

11 (b) A prescriber shall comply with any additional requirements of rules
12 adopted by the Commissioner of Health in accordance with section 4287 and
13 subsection 4289(e) of this title, which shall be consistent with the provisions of
14 this section.

15 § 4273. PRIVATE CAUSE OF ACTION

16 (a) A person who is injured as a result of the development of an opioid
17 dependency or addiction caused by the failure of a prescriber to comply with
18 the requirements of subsection 4272(a) of this title may bring an action in
19 Superior Court against the prescriber for damages, reasonable costs, and
20 attorney's fees.

21 (b) In an action brought under subsection (a) of this section:

1 (1) The court may issue an award for the person's actual damages or
2 \$500.00 for a first violation or \$1,000.00 for each subsequent violation,
3 whichever is greater.

4 (2) Any affiliate or employer of the prescriber shall be jointly and
5 severally liable for all damages, costs, and fees awarded to the complainant
6 pursuant to this section.

7 (3) If the prescriber commits two or more violations of subsection
8 4272(e) of this title with respect to the complainant within any 12-month
9 period, the prescriber shall be subject to a rebuttable presumption that the
10 prescriber caused the opioid dependency and any injury suffered by the
11 complainant.

12 (4) If the prescriber commits three or more violations of subsection
13 4272(e) of this title with respect to the care and treatment of the complainant in
14 any 12-month period, the prescriber shall be subject to an award of punitive
15 damages.

16 (5) If the prescriber commits four or more violations of subsection
17 4272(e) of this title with respect to the care and treatment of the complainant in
18 any 12-month period, any manufacturer and distributor of an opioid controlled
19 substance prescribed to the complainant consistently throughout that period as
20 a palliative treatment for acute or chronic pain shall be strictly liable, jointly

1 and severally, for all damages, costs, and fees awarded to the complainant
2 pursuant to this section.

3 (c) This section shall not limit any other claims the person may have under
4 applicable law.

5 (d) An action brought under this section shall be commenced not later than
6 three years after the date of the last occurring violation of subsection 4272(a)
7 of this title.

8 Sec. 2. EFFECTIVE DATE

9 This act shall take effect on July 1, 2018.