HOUSE BILL No. 5103

August 24, 2005, Introduced by Rep. Caswell and referred to the Committee on Health Policy.

A bill to amend 1978 PA 368, entitled "Public health code,"

by amending section 7333 (MCL 333.7333), as amended by 2001 PA 231.

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

- 1 Sec. 7333. (1) As used in this section, "good faith" means the
- 2 prescribing or dispensing of a controlled substance by a
- 3 practitioner licensed under section 7303 in the regular course of
- 4 professional treatment to or for an individual who is under
- 5 treatment by the practitioner for a pathology or condition other
- 6 than that individual's physical or psychological dependence upon or
 - addiction to a controlled substance, except as provided in this
- article. Application of good faith to a pharmacist means the

- 1 dispensing of a controlled substance pursuant to a prescriber's
- 2 order which, in the professional judgment of the pharmacist, is
- 3 lawful. The pharmacist shall be guided by nationally accepted
- 4 professional standards including, but not limited to, all of the
- 5 following, in making the judgment:
- 6 (a) Lack of consistency in the doctor-patient relationship.
- 7 (b) Frequency of prescriptions for the same drug by 1
- 8 prescriber for larger numbers of patients.
- 9 (c) Quantities beyond those normally prescribed for the same
- 10 drug.
- 11 (d) Unusual dosages.
- 12 (e) Unusual geographic distances between patient, pharmacist,
- 13 and prescriber.
- 14 (2) Except as otherwise provided in this section, a
- 15 practitioner, in good faith, may dispense a controlled substance
- 16 included in schedule 2 upon receipt of a prescription of a
- 17 practitioner licensed under section 7303 on a prescription form. A
- 18 practitioner shall not issue more than 1 prescription for a
- 19 controlled substance included in schedule 2 on a single
- 20 prescription form.
- 21 (3) In an emergency situation, as described in R $\frac{-338.3165}{}$
- 22 338.3164 of the Michigan administrative code, a controlled
- 23 substance included in schedule 2 may be dispensed upon the oral
- 24 prescription of a practitioner if, the prescribing practitioner
- 25 promptly fills out a prescription form and forwards the
- 26 prescription form to the dispensing pharmacy within 7 days after
- 27 the oral prescription is issued.

- 1 (4) EXCEPT AS OTHERWISE PROVIDED UNDER SUBSECTION (5), IF A
- 2 DISPENSING PHARMACIST IS UNABLE TO SUPPLY THE FULL QUANTITY CALLED
- 3 FOR IN A WRITTEN OR AN EMERGENCY ORAL PRESCRIPTION FOR A CONTROLLED
- 4 SUBSTANCE INCLUDED IN SCHEDULE 2, HE OR SHE MAY PARTIALLY FILL THAT
- 5 PRESCRIPTION AS LONG AS HE OR SHE MAKES A NOTATION OF THE QUANTITY
- 6 SUPPLIED ON THE FACE OF THE WRITTEN PRESCRIPTION OR WRITTEN RECORD
- 7 OF THE EMERGENCY ORAL PRESCRIPTION. THE REMAINDER OF THE
- 8 PRESCRIPTION MAY BE DISPENSED WITHIN 72 HOURS AFTER THE FIRST
- 9 PARTIAL DISPENSING. IF THE REMAINDER OF THE PRESCRIPTION IS NOT OR
- 10 CANNOT BE DISPENSED WITHIN THE 72 HOURS, THE PHARMACIST SHALL
- 11 NOTIFY THE PRESCRIBER. AFTER THE 72-HOUR PERIOD, A PHARMACIST SHALL
- 12 NOT DISPENSE A FURTHER QUANTITY WITHOUT A NEW PRESCRIPTION.
- 13 (5) Except for a terminally ill patient whose terminal illness
- 14 the pharmacist documents pursuant to rules promulgated by the
- 15 administrator, OR FOR A PATIENT IN A LONG-TERM CARE FACILITY WHOSE
- 16 STATUS AS A LONG-TERM CARE FACILITY PATIENT IS DOCUMENTED BY THE
- 17 PHARMACIST PURSUANT TO RULES PROMULGATED BY THE ADMINISTRATOR, a
- 18 prescription for a controlled substance included in schedule 2
- 19 shall not be filled more than 60 days after the date on which the
- 20 prescription was issued. A prescription for a controlled substance
- 21 included in schedule 2 for a terminally ill patient whose terminal
- 22 illness the pharmacist documents pursuant to rules promulgated by
- 23 the administrator OR FOR A PATIENT IN A LONG-TERM CARE FACILITY
- 24 WHOSE STATUS AS A LONG-TERM CARE FACILITY PATIENT IS DOCUMENTED BY
- 25 THE PHARMACIST PURSUANT TO RULES PROMULGATED BY THE ADMINISTRATOR,
- 26 may be partially filled in increments, INCLUDING INDIVIDUAL DOSAGE
- 27 UNITS, for not more than 60 days after the date on which the

- 1 prescription was issued. THE TOTAL QUANTITY OF SCHEDULE 2
- 2 CONTROLLED SUBSTANCES DISPENSED IN ALL PARTIAL FILLINGS UNDER THIS
- 3 SUBSECTION AND SUBSECTION (4) SHALL NOT BE MORE THAN THE TOTAL
- 4 QUANTITY PRESCRIBED. FOR EACH PARTIAL FILLING UNDER THIS
- 5 SUBSECTION, THE PHARMACIST SHALL RECORD, ON THE BACK OF THE
- 6 PRESCRIPTION OR ON ANOTHER APPROPRIATE RECORD THAT IS UNIFORMLY
- 7 MAINTAINED AND READILY RETRIEVABLE, ALL OF THE FOLLOWING
- 8 INFORMATION:
- 9 (A) DATE OF THE PARTIAL FILLING.
- 10 (B) QUANTITY DISPENSED.
- 11 (C) REMAINING QUANTITY AUTHORIZED TO BE DISPENSED.
- 12 (D) IDENTIFICATION OF THE DISPENSING PHARMACIST.
- 13 (6) -(4) A practitioner, in good faith, may dispense a
- 14 controlled substance included in schedule 3, 4, or 5 that is a
- 15 prescription drug as determined under section 503(b) of the federal
- 16 food, drug, and cosmetic act, -chapter 675, 52 Stat. 1051, 21
- 17 U.S.C. USC 353, or section 17708, upon receipt of a prescription
- 18 on a prescription form or an oral prescription of a practitioner. A
- 19 prescription for a controlled substance included in schedule 3 or 4
- 20 shall not be filled or refilled without specific refill
- 21 instructions noted by the prescriber. A prescription for a
- 22 controlled substance included in schedule 3 or 4 shall not be
- 23 filled or refilled later than 6 months after the date of the
- 24 prescription or be refilled more than 5 times, unless renewed by
- 25 the prescriber in accordance with rules promulgated by the
- 26 administrator.
- 27 (7) -(5) A controlled substance included in schedule 5 shall

- 1 not be distributed or dispensed other than for a medical purpose,
- 2 or in any manner except in accordance with rules promulgated by the
- 3 administrator.
- 4 (8) —(6)— If a prescription is required under this section,
- 5 the prescription shall contain the quantity of the controlled
- 6 substance prescribed in both written and numerical terms. A
- 7 prescription is in compliance with this subsection if, in addition
- 8 to containing the quantity of the controlled substance prescribed
- 9 in written terms, it contains preprinted numbers representative of
- 10 the quantity of the controlled substance prescribed next to which
- 11 is a box or line the prescriber may check.
- 12 (9) -(7) A prescribing practitioner shall not use a
- 13 prescription form for a purpose other than prescribing. A
- 14 prescribing practitioner shall not postdate a prescription form
- 15 that contains a prescription for a controlled substance. A
- 16 prescriber may transmit a prescription by facsimile of a printed
- 17 prescription form and by electronic transmission of a printed
- 18 prescription form, if not prohibited by federal law. If, with the
- 19 patient's consent, a prescription is electronically transmitted, it
- 20 shall be transmitted directly to a pharmacy of the patient's choice
- 21 by the prescriber or the prescriber's authorized agent, and the
- 22 data shall not be altered, modified, or extracted in the
- 23 transmission process.
- 24 (10) $\frac{(8)}{(8)}$ Notwithstanding subsections (1) to $\frac{(5)}{(7)}$, a dog
- 25 pound or animal shelter licensed or registered by the department of
- 26 agriculture pursuant to 1969 PA 287, MCL 287.331 to 287.340, or a
- 27 class B dealer may acquire a limited permit only for the purpose of

- 1 buying, possessing, and administering a commercially prepared,
- 2 premixed solution of sodium pentobarbital to practice euthanasia on
- 3 injured, sick, homeless, or unwanted domestic pets and other
- 4 animals, if the dog pound or animal shelter or class B dealer does
- 5 all of the following:
- 6 (a) Applies to the administrator for a permit in accordance
- 7 with rules promulgated under this part. The application shall
- 8 contain the name of the individual in charge of the day to day
- 9 operations of the dog pound or animal shelter or class B dealer's
- 10 facilities and the name of the individual responsible for
- 11 designating employees who will be practicing euthanasia on animals
- 12 pursuant to this act.
- 13 (b) Complies with the rules promulgated by the administrator
- 14 for the storage, handling, and use of commercially prepared,
- 15 premixed solution of sodium pentobarbital to practice euthanasia on
- 16 animals. A record of use shall be maintained and shall be available
- 17 for inspection.
- (c) Certifies that an employee of the dog pound or animal
- 19 shelter or class B dealer has received, and can document completion
- 20 of, a minimum of 8 hours of training given by a licensed
- 21 veterinarian in the use of sodium pentobarbital to practice
- 22 euthanasia on animals pursuant to rules promulgated by the
- 23 administrator, in consultation with the Michigan board of
- 24 veterinary medicine as these rules relate to this training, and
- 25 that only an individual described in this subdivision or an
- 26 individual otherwise permitted to use a controlled substance
- 27 pursuant to this article will administer the commercially prepared,

- 1 premixed solution of sodium pentobarbital according to written
- 2 procedures established by the dog pound or animal shelter or class
- **3** B dealer.
- 4 (11) -(9) The application described in subsection -(8) (10)
- 5 shall include the names and addresses of all individuals employed
- 6 by the dog pound or animal shelter or class B dealer who have been
- 7 trained as described in subsection $\frac{(8)(c)}{(10)(c)}$ and the name of
- 8 the veterinarian who trained them. The list of names and addresses
- 9 shall be updated every 6 months.
- 10 (12) $\frac{(10)}{(10)}$ If a dog pound or animal shelter or class B dealer
- 11 issued a permit pursuant to subsection $\frac{(8)}{(10)}$ does not have in
- 12 its employ an individual trained as described in subsection $\frac{(8)(c)}{c}$
- 13 (10)(C), the dog pound or animal shelter or class B dealer shall
- 14 immediately notify the administrator and shall cease to administer
- 15 any commercially prepared, premixed solution of sodium
- 16 pentobarbital until the administrator is notified that 1 of the
- 17 following has occurred:
- 18 (a) An individual trained as described in subsection $\frac{(8)(c)}{(c)}$
- 19 (10)(C) has been hired by the dog pound or animal shelter or class
- 20 B dealer.
- 21 (b) An employee of the dog pound or animal shelter or class B
- 22 dealer has been trained as described in subsection $\frac{(8)(c)}{(10)(c)}$.
- 23 (13) -(11) A veterinarian, including a veterinarian who
- 24 trains individuals as described in subsection $\frac{(8)(c)}{(10)(C)}$, is
- 25 not civilly or criminally liable for the use of a commercially
- 26 prepared, premixed solution of sodium pentobarbital by a dog pound
- 27 or animal shelter or class B dealer unless the veterinarian is

- 1 employed by or under contract with the dog pound or animal shelter
- 2 or class B dealer and the terms of the veterinarian's employment or
- 3 the contract require the veterinarian to be responsible for the use
- 4 or administration of the commercially prepared, premixed solution
- 5 of sodium pentobarbital.
- 6 (14) $\frac{(12)}{(12)}$ A person shall not knowingly use or permit the use
- 7 of a commercially prepared, premixed solution of sodium
- 8 pentobarbital in violation of this section.
- 9 (15) -(13)— This section does not require that a veterinarian
- 10 be employed by or under contract with a dog pound or animal shelter
- 11 or class B dealer to obtain, possess, or administer a commercially
- 12 prepared, premixed solution of sodium pentobarbital pursuant to
- 13 this section.
- 14 (16) $\overline{(14)}$ As used in this section, "class B dealer" means a
- 15 class B dealer licensed by the United States department of
- 16 agriculture pursuant to the animal welfare act, Public Law 89-544,
- 17 7 U.S.C. 2131 to 2147, 2149, and 2151 7 USC 2131 to 2159 and the
- 18 department of agriculture pursuant to 1969 PA 224, MCL 287.381 to
- **19** 287.395.

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