SUBSTITUTE FOR HOUSE BILL NO. 5262

A bill to amend 1978 PA 368, entitled "Public health code," by amending sections 7104, 7107, and 7109 (MCL 333.7104, 333.7107, and 333.7109), section 7104 as amended by 1994 PA 38 and sections 7107 and 7109 as amended by 1993 PA 80.

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

- Sec. 7104. (1) "Bureau" means the drug enforcement adminis tration, United States department of justice, or its successor
 agency.
- 4 (2) "Controlled substance" means a drug, substance, or imme-
- ${f 5}$ diate precursor included in schedules 1 to 5 of part 72.
- **6** (3) "Controlled substance analogue" means a substance the
- 7 chemical structure of which is substantially similar to that of a
- 8 controlled substance in schedule 1 or 2 and that has a narcotic,
- 9 stimulant, depressant, or hallucinogenic effect on the central

- 1 nervous system substantially similar to or greater than the
- 2 narcotic, stimulant, depressant, or hallucinogenic effect on the

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- 3 central nervous system of a controlled substance included in
- 4 schedule 1 or 2 or, with respect to a particular individual, that
- 5 the individual represents or intends to have a narcotic, stimu-
- 6 lant, depressant, or hallucinogenic effect on the central nervous
- 7 system substantially similar to or greater than the narcotic,
- 8 stimulant, depressant, or hallucinogenic effect on the central
- 9 nervous system of a controlled substance included in schedule 1
- 10 or 2. Controlled substance analogue does not include any 1 OR
- 11 MORE of the following:
- 12 (a) A controlled substance.
- 13 (b) A substance for which there is an approved new drug
- 14 application.
- 15 (c) A substance with respect to which an exemption is in
- 16 effect for investigational use by a particular person under
- 17 section 505 of the federal food, drug and cosmetic act,
- **18** chapter 675, 52 Stat. 1040— 1052, 21 U.S.C. 355, to the extent
- 19 conduct with respect to the substance is pursuant to the
- 20 exemption.
- 21 (d) Any substance to the extent not intended for human con-
- 22 sumption before an exemption takes effect with respect to the
- 23 substance.
- (4) "Counterfeit prescription form" means a printed form
- 25 that is the same or similar to a prescription form or an offi-
- 26 cial prescription form, and that was manufactured, printed,
- 27 duplicated, forged, ELECTRONICALLY TRANSMITTED, or altered

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- 1 without the knowledge or permission of a licensed prescribing
- 2 practitioner, or, in the case of official prescription forms, the
- 3 department of commerce PRESCRIBER.
- 4 (5) "Counterfeit substance" means a controlled substance
- 5 which THAT, or the container or labeling of which, without
- 6 authorization, bears the trademark, trade name or other identify-
- 7 ing mark, imprint, number, or device, or any likeness thereof, of
- 8 a manufacturer, distributor, or dispenser other than the person
- 9 who in fact manufactured, distributed, or dispensed the
- 10 substance.
- 11 (6) "Deleterious drug" means a drug, other than a propri-
- 12 etary medicine, likely to be destructive to adult human life in
- 13 quantities of 3.88 grams or less.
- 14 (7) "ELECTRONIC SIGNATURE" MEANS AN ELECTRONIC SOUND,
- 15 SYMBOL, OR PROCESS ATTACHED TO OR LOGICALLY ASSOCIATED WITH A
- 16 RECORD AND EXECUTED OR ADOPTED BY A PERSON WITH THE INTENT TO
- 17 SIGN THE RECORD.
- 18 Sec. 7107. -(1) "Narcotic drug" means any 1 OR MORE of
- 19 the following, whether produced directly or indirectly by extrac-
- 20 tion from substances of vegetable origin, or independently by
- 21 means of chemical synthesis, or by a combination of extraction
- 22 and chemical synthesis:
- 23 (a) Opium and opiate, and any salt, compound, derivative, or
- 24 preparation of opium or opiate.
- 25 (b) Any salt, compound, isomer, derivative, or preparation
- 26 thereof which is chemically equivalent or identical with any of

1 the substances referred to in subdivision (a), but not including

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- 2 the isoquinoline alkaloids of opium.
- 3 (2) "Official prescription form" means a prescription form
- 4 that meets the requirements of section 7334 and is issued to
- 5 practitioners by the department of commerce.
- 6 Sec. 7109. (1) "Person" means a person as defined in sec-
- 7 tion 1106 or a governmental entity.
- 8 (2) "Poppy straw" means all parts, except the seeds, of the
- 9 opium poppy, after mowing.
- 10 (3) "Practitioner" means:
- 11 (a) A prescriber or pharmacist, a scientific investigator as
- 12 defined by rule of the administrator, or other person licensed,
- 13 registered, or otherwise permitted to distribute, dispense, con-
- 14 duct research with respect to, or administer a controlled sub-
- 15 stance in the course of professional practice or research in this
- 16 state, including an individual in charge of a dog pound or animal
- 17 shelter licensed or registered by the department of agriculture
- 18 pursuant to Act No. 287 of the Public Acts of 1969, being sec-
- 19 tions 287.331 to 287.340 of the Michigan Compiled Laws 1969
- 20 PA 287, MCL 287.331 TO 287.340, or a class B dealer licensed by
- 21 the United States department of agriculture pursuant to the
- 22 animal welfare act, Public Law 89-544, 7 U.S.C. 2131 to 2147,
- 23 2149, and 2151 to 2159 and the department of agriculture pursuant
- 24 to Act No. 224 of the Public Acts of 1969, being
- 25 sections 287.381 to 287.395 of the Michigan Compiled Laws 1969
- 26 PA 224, MCL 287.381 TO 287.395, for the limited purpose of
- 27 buying, possessing, and administering a commercially prepared,

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HB5262, As Passed House, November 1, 2001
Sub. HB 5262 (H-3) as amended November 1, 2001
 1 premixed solution of sodium pentobarbital to practice euthanasia
 2 on animals.
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        (b) A pharmacy, hospital, or other institution or place of
 4 professional practice licensed, registered, or otherwise permit-
 5 ted to distribute, prescribe, dispense, conduct research with
 6 respect to, or administer a controlled substance in the course of
 7 professional practice or research in this state.
        (4) "PRESCRIBER" MEANS THAT TERM AS DEFINED IN SECTION
 9 17708.
        (5) (4) "Prescription form" means a printed form, which
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11 THAT is authorized and intended for use by a prescribing practi-
12 tioner to prescribe controlled substances or OTHER prescription
13 drugs and which THAT meets the requirements of rules promul-
14 gated by the administrator, -- AND ALL OF THE FOLLOWING
15 REQUIREMENTS:
16
        (A) BEARS THE PREPRINTED, STAMPED, TYPED, OR MANUALLY
17 PRINTED NAME, ADDRESS, AND TELEPHONE NUMBER OR PAGER NUMBER OF
18 THE PRESCRIBING PRACTITIONER [.
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20
        (B) INCLUDES THE MANUALLY PRINTED NAME OF THE PATIENT, THE
21 ADDRESS OF THE PATIENT, THE PRESCRIBING PRACTITIONER'S SIGNATURE,
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- 22 AND THE PRESCRIBING PRACTITIONER'S DRUG ENFORCEMENT ADMINISTRA-
- 23 TION REGISTRATION NUMBER.
- 24 (C) THE QUANTITY OF THE PRESCRIPTION DRUG PRESCRIBED, IN
- 25 BOTH WRITTEN AND NUMERICAL TERMS.
 - [(D) INCLUDES THE DATE THE PRESCRIPTION DRUG WAS PRESCRIBED. (E) ANY RULES PROMULGATED BY THE DEPARTMENT PURSUANT TO SECTION 7333A(5).]
- (6) $\frac{(5)}{(5)}$ "Production" means the manufacture, planting, 26
- 27 cultivation, growing, or harvesting of a controlled substance.

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- (7) "SIGN" MEANS TO AFFIX ONE'S SIGNATURE MANUALLY TO A 1
- 2 DOCUMENT OR TO USE AN ELECTRONIC SIGNATURE.
- (8) $\overline{\text{(6)}}$ "Ultimate user" means an individual who lawfully 3
- 4 possesses a controlled substance for personal use or for the use
- 5 of a member of the individual's household, or for administering
- 6 to an animal owned by the individual or by a member of the
- 7 individual's household.
- Enacting section 1. Sections 7104, 7107, and 7109 of the
- 9 public health code, 1978 PA 368, MCL 333.7104, 333.7107, and
- 10 333.7109, as amended by this amendatory act, take effect upon the
- 11 promulgation of the rules required under section 7333a and
- 12 receipt by the secretary of state of written notice from the
- 13 director of the department of consumer and industry services that
- 14 the electronic monitoring system required by section 7333a is
- 15 operational. The notice to the secretary of state shall include
- 16 a statement that the department of consumer and industry services
- 17 is able to receive data from at least 80% of those required to
- 18 report under section 7333a and is able to respond to requests for
- 19 data from persons authorized to make such requests and to review
- 20 and utilize the data.
- 21 Enacting section 2. This amendatory act does not take
- 22 effect unless House Bill No. 5260 of the 91st Legislature is
- 23 enacted into law.