HOUSE BILL No. 4369

March 2, 2011, Introduced by Reps. Liss, Segal, Barnett and Haugh and referred to the Committee on Health Policy.

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

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

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

- 1 Sec. 7333a. (1) The department shall establish, by rule, an
- 2 electronic system for monitoring schedule 2, 3, 4, and 5 controlled
- 3 substances dispensed in this state by veterinarians, and by
- 4 pharmacists and dispensing prescribers licensed under part 177 or
- 5 dispensed to an address in this state by a pharmacy licensed in
- 6 this state. The rules shall provide an appropriate electronic
- 7 format for the reporting of data including, but not limited to,
- patient identifiers, the name of the controlled substance
- 9 dispensed, date of dispensing, quantity dispensed, prescriber, and
- 10 dispenser. The department shall require a veterinarian, pharmacist,

- 1 or dispensing prescriber to utilize the electronic data transmittal
- 2 process developed by the department or the department's contractor.
- 3 A veterinarian, pharmacist, or dispensing prescriber shall not be
- 4 required to pay a new fee dedicated to the operation of the
- 5 electronic monitoring system and shall not incur any additional
- 6 costs solely related to the transmission of data to the department.
- 7 The rules promulgated under this subsection shall exempt both of
- 8 the following circumstances from the reporting requirements:
- 9 (a) The administration of a controlled substance directly to a
- 10 patient.
- 11 (b) The dispensing from a health facility or agency licensed
- 12 under article 17 of a controlled substance by a dispensing
- 13 prescriber in a quantity adequate to treat a patient for not more
- 14 than 48 hours.
- 15 (2) Notwithstanding any practitioner-patient privilege, the
- 16 director of the department may provide data obtained under this
- 17 section to all of the following:
- 18 (a) A designated representative of a board responsible for the
- 19 licensure, regulation, or discipline of a practitioner, pharmacist,
- 20 or other person who is authorized to prescribe, administer, or
- 21 dispense controlled substances.
- (b) An employee or agent of the department.
- 23 (c) A state, federal, or municipal employee or agent whose
- 24 duty is to enforce the laws of this state or the United States
- 25 relating to drugs.
- 26 (d) A state-operated medicaid program.
- (e) A state, federal, or municipal employee who is the holder

- 1 of a search warrant or subpoena properly issued for the records.
- 2 (f) A practitioner or pharmacist who requests information and
- 3 certifies that the requested information is for the purpose of
- 4 providing medical or pharmaceutical treatment to a bona fide
- 5 current patient.
- 6 (g) An individual with whom the department has contracted
- 7 under subsection (9) (8).
- 8 (H) THE HEALTH CARE PAYMENT OR BENEFIT PROVIDER FOR THE
- 9 PURPOSES OF ENSURING PATIENT SAFETY AND INVESTIGATING FRAUD AND
- 10 ABUSE. AS USED IN THIS SUBDIVISION, "HEALTH CARE PAYMENT OR BENEFIT
- 11 PROVIDER" MEANS A PERSON THAT PROVIDES HEALTH BENEFITS, COVERAGE,
- 12 OR INSURANCE IN THIS STATE, INCLUDING A HEALTH INSURANCE COMPANY, A
- 13 NONPROFIT HEALTH CARE CORPORATION, A HEALTH MAINTENANCE
- 14 ORGANIZATION, A MULTIPLE EMPLOYER WELFARE ARRANGEMENT, A MEDICAID
- 15 CONTRACTED HEALTH PLAN, OR ANY OTHER PERSON PROVIDING A PLAN OF
- 16 HEALTH BENEFITS, COVERAGE, OR INSURANCE SUBJECT TO STATE INSURANCE
- 17 REGULATION.
- 18 (3) Except as otherwise provided in this part, information
- 19 submitted under this section shall be used only for bona fide drug-
- 20 related criminal investigatory or evidentiary purposes or for the
- 21 investigatory or evidentiary purposes in connection with the
- 22 functions of a disciplinary subcommittee or 1 or more of the
- 23 licensing or registration boards created in article 15.
- 24 (4) A person who receives data or any report under subsection
- 25 (2) containing any patient identifiers of the system from the
- 26 department shall not provide it to any other person or entity
- 27 except by order of a court of competent jurisdiction.

- 1 (5) Except as otherwise provided in this subsection, reporting
- 2 under subsection (1) is mandatory for a veterinarian, pharmacist,
- 3 and dispensing prescriber. However, the department may issue a
- 4 written waiver of the electronic reporting requirement to a
- 5 veterinarian, pharmacist, or dispensing prescriber who establishes
- 6 grounds that he or she is unable to use the electronic monitoring
- 7 system. The department shall require the applicant for the waiver
- 8 to report the required information in a manner approved by the
- 9 department.
- 10 (6) In addition to the information required to be reported
- 11 annually under section 7112(3), the controlled substances advisory
- 12 commission shall include in the report information on the
- 13 implementation and effectiveness of the electronic monitoring
- 14 system.
- 15 (7) The department, in consultation with the controlled
- 16 substances advisory commission, the Michigan board of pharmacy, the
- 17 Michigan board of medicine, the Michigan board of osteopathic
- 18 medicine and surgery, the Michigan state police, and appropriate
- 19 medical professional associations, shall examine the need for and
- 20 may promulgate rules for the production of a prescription form on
- 21 paper that minimizes the potential for forgery. The rules shall not
- 22 include any requirement that sequential numbers, bar codes, or
- 23 symbols be affixed, printed, or written on a prescription form or
- 24 that the prescription form be a state produced prescription form.
- 25 In examining the need for rules for the production of a
- 26 prescription form on paper that minimizes the potential for
- 27 forgery, the department shall consider and identify the following:

- 1 (a) Cost, benefits, and barriers.
- 2 (b) Overall cost-benefit analysis.
- 3 (c) Compatibility with the electronic monitoring system
- 4 required under this section.
- 5 (8) The department shall report its findings under subsection
- 6 (7) to the members of the house and senate standing committees
- 7 having jurisdiction over health policy issues not later than
- 8 October 1, 2002, and before the electronic monitoring system
- 9 required under this section becomes operational.
- 10 (8) (9) The department may enter into 1 or more contractual
- 11 agreements for the administration of this section.
- 12 (9) (10) The department, all law enforcement officers, all
- 13 officers of the court, and all regulatory agencies and officers, in
- 14 using the data for investigative or prosecution purposes, shall
- 15 consider the nature of the prescriber's and dispenser's practice
- 16 and the condition for which the patient is being treated.
- 17 (10) (11) The data and any report containing any patient
- 18 identifiers obtained therefrom is not a public record, and is not
- 19 subject to the freedom of information act, 1976 PA 442, MCL 15.231
- 20 to 15.246.
- 21 (11) (12) As used in this section, "department" means the
- 22 department of consumer and industry services COMMUNITY HEALTH.