HOUSE SUBSTITUTE FOR SENATE BILL NO. 47

A bill to amend 1978 PA 368, entitled "Public health code," by amending section 7333a (MCL 333.7333a), as amended by 2016 PA

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 must provide an appropriate electronic format
- 7 for the reporting of data including, but not limited to, patient
- 8 identifiers, and the name of the controlled substance dispensed,
- 9 the date of dispensing, the quantity dispensed, the prescriber, and

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- 1 the dispenser. The department shall require a veterinarian,
- 2 pharmacist, or dispensing prescriber to utilize the electronic data
- 3 transmittal process developed by the department or the department's
- 4 contractor. The department shall not require a veterinarian,
- 5 pharmacist, or dispensing prescriber to pay a new fee dedicated to
- 6 the operation of the electronic monitoring system or to incur any
- 7 additional costs solely related to the transmission of data to the
- 8 department. The rules promulgated under this subsection must exempt
- 9 both of the following circumstances from the reporting
- 10 requirements:
- 11 (a) The administration of a controlled substance directly to a
- 12 patient.
- 13 (b) The dispensing from a health facility or agency licensed
- 14 under article 17 of a controlled substance by a dispensing
- 15 prescriber in a quantity adequate to treat a patient for not more
- 16 than 48 hours. THE DISPENSING OF A CONTROLLED SUBSTANCE IN ANY OF
- 17 THE FOLLOWING IS EXEMPT FROM THE REPORTING REQUIREMENTS:
- 18 (A) A HOSPITAL THAT IS LICENSED UNDER ARTICLE 17 THAT
- 19 ADMINISTERS THE CONTROLLED SUBSTANCE TO AN INDIVIDUAL WHO IS AN
- 20 INPATIENT.
- 21 (B) A HEALTH FACILITY OR AGENCY LICENSED UNDER ARTICLE 17 IF
- 22 THE CONTROLLED SUBSTANCE IS DISPENSED BY A DISPENSING PRESCRIBER IN
- 23 A QUANTITY ADEQUATE TO TREAT THE PATIENT FOR NOT MORE THAN 48
- 24 HOURS.
- 25 (C) A VETERINARY HOSPITAL OR CLINIC THAT ADMINISTERS THE
- 26 CONTROLLED SUBSTANCE TO AN ANIMAL THAT IS AN INPATIENT.
- 27 (2) Notwithstanding any practitioner-patient privilege, the

- 1 director of the department may provide data obtained under this
- 2 section to all of the following:
- 3 (a) A designated representative of a board responsible for the
- 4 licensure, regulation, or discipline of a practitioner, pharmacist,
- 5 or other person that is authorized to prescribe, administer, or
- 6 dispense controlled substances.
- 7 (b) An employee or agent of the department.
- 8 (c) A state, federal, or municipal employee or agent whose
- 9 duty is to enforce the laws of this state or the United States
- 10 relating to drugs.
- 11 (d) A state-operated Medicaid program.
- 12 (e) A state, federal, or municipal employee who is the holder
- 13 of a search warrant or subpoena properly issued for the records.
- 14 (f) A practitioner or pharmacist who requests information and
- 15 certifies that the requested information is for the purpose of
- 16 providing medical or pharmaceutical treatment to a bona fide
- 17 current patient.
- 18 (q) An individual with whom the department has contracted
- 19 under subsection (7).
- 20 (h) A practitioner or other person that is authorized to
- 21 prescribe controlled substances for the purpose of determining if
- 22 prescriptions written by that practitioner or other person have
- 23 been dispensed.
- 24 (i) The health care payment or benefit provider for the
- 25 purposes of ensuring patient safety and investigating fraud and
- 26 abuse.
- 27 (3) Except as otherwise provided in this part, a person shall

- 1 use information submitted under this section only for bona fide
- 2 drug-related criminal investigatory or evidentiary purposes or for
- 3 the investigatory or evidentiary purposes in connection with the
- 4 functions of a disciplinary subcommittee or 1 or more of the
- 5 licensing or registration boards created in article 15.
- 6 (4) A person that receives data or any report under subsection
- 7 (2) containing any patient identifiers of the system from the
- 8 department shall not provide it to any other person except by order
- 9 of a court of competent jurisdiction.
- 10 (5) Except as otherwise provided in this subsection, reporting
- 11 under subsection (1) is mandatory for a veterinarian, pharmacist,
- 12 and dispensing prescriber. However, the department may issue a
- 13 written waiver of the electronic reporting requirement to a
- 14 veterinarian, pharmacist, or dispensing prescriber who establishes
- 15 grounds that he or she is unable to use the electronic monitoring
- 16 system. The department shall require the applicant for the waiver
- 17 to report the required information in a manner approved by the
- 18 department.
- 19 (6) The department, in consultation with the Michigan board of
- 20 pharmacy, the Michigan board of medicine, the Michigan board of
- 21 osteopathic medicine and surgery, the department of state police,
- 22 and appropriate medical professional associations, shall examine
- 23 the need for and may promulgate rules for the production of a
- 24 prescription form on paper that minimizes the potential for
- 25 forgery. The rules must not include any requirement that sequential
- 26 numbers, bar codes, or symbols be affixed, printed, or written on a
- 27 prescription form or that the prescription form be a state produced

- 1 prescription form. In examining the need for rules for the
- 2 production of a prescription form on paper that minimizes the
- 3 potential for forgery, the department shall consider and identify
- 4 the following:
- 5 (a) Cost, benefits, and barriers.
- 6 (b) Overall cost-benefit analysis.
- 7 (c) Compatibility with the electronic monitoring system
- 8 required under this section.
- 9 (7) The department may enter into 1 or more contractual
- 10 agreements for the administration of this section.
- 11 (8) The department, all law enforcement officers, all officers
- 12 of the court, and all regulatory agencies and officers, in using
- 13 the data for investigative or prosecution purposes, shall consider
- 14 the nature of the prescriber's and dispenser's practice and the
- 15 condition for which the patient is being treated.
- 16 (9) The data and any report containing any patient identifiers
- 17 obtained from the data are not public records and are not subject
- 18 to DISCLOSURE UNDER the freedom of information act, 1976 PA 442,
- **19** MCL 15.231 to 15.246.
- 20 (10) The department may issue a written request to a health
- 21 care payment or benefit provider to determine if the provider has
- 22 accessed the electronic monitoring system as provided in subsection
- 23 (2)(i) in the previous calendar year and, if so, to determine the
- 24 number of inquiries the provider made in the previous calendar year
- 25 and any other information the department requests in relation to
- 26 the provider's access to the electronic monitoring system. A health
- 27 care payment or benefit provider shall respond to the written

- 1 request on or before the March 31 following the request. The
- 2 department shall collaborate with health care payment or benefit
- 3 providers to develop a reasonable request and reporting form for
- 4 use under this subsection.
- 5 (11) BEFORE DISPENSING OR PRESCRIBING BUPRENORPHINE, OR A DRUG
- 6 CONTAINING BUPRENORPHINE OR METHADONE, TO A PATIENT IN A SUBSTANCE
- 7 USE DISORDER PROGRAM, A PRESCRIBER SHALL OBTAIN AND REVIEW DATA
- 8 CONCERNING THAT PATIENT FROM THE DEPARTMENT UNDER SUBSECTION (2). A
- 9 PRESCRIBER DISPENSING BUPRENORPHINE, OR A DRUG CONTAINING
- 10 BUPRENORPHINE OR METHADONE, TO A PATIENT IN A SUBSTANCE USE
- 11 DISORDER PROGRAM SHALL ALSO REPORT THE DATA REQUIRED IN SUBSECTION
- 12 (1), IF FEDERAL LAW DOES NOT PROHIBIT THE REPORTING OF DATA
- 13 CONCERNING THE PATIENT, TO THE DEPARTMENT. AS USED IN THIS
- 14 SUBSECTION:
- 15 (A) "APPROVED SERVICE PROGRAM" MEANS THAT TERM AS DEFINED IN
- 16 SECTION 100A OF THE MENTAL HEALTH CODE, 1974 PA 258, MCL 330.1100A.
- 17 (B) "SUBSTANCE USE DISORDER PROGRAM" MEANS A PROGRAM AS THAT
- 18 TERM IS DEFINED IN SECTION 260 OF THE MENTAL HEALTH CODE, 1974 PA
- 19 258, MCL 330.1260, AN APPROVED SERVICE PROGRAM, A NONREGULATED
- 20 SUBSTANCE USE DISORDER SERVICES PROGRAM, A FEDERAL CERTIFIED
- 21 SUBSTANCE USE DISORDER SERVICES PROGRAM, OR A FEDERALLY REGULATED
- 22 SUBSTANCE USE DISORDER SERVICES PROGRAM.
- 23 (12) R 338.3162E OF THE MICHIGAN ADMINISTRATIVE CODE IS
- 24 RESCINDED.
- 25 (13) $\frac{(11)}{}$ As used in this section:
- 26 (a) "Department" means the department of licensing and
- 27 regulatory affairs.

- 1 (b) "Health care payment or benefit provider" means a person
- 2 that provides health benefits, coverage, or insurance in this
- 3 state, including a health insurance company, a nonprofit health
- 4 care corporation, a health maintenance organization, a multiple
- 5 employer welfare arrangement, a Medicaid contracted health plan, or
- 6 any other person providing a plan of health benefits, coverage, or
- 7 insurance subject to state insurance regulation.
- 8 Enacting section 1. This amendatory act takes effect 90 days
- 9 after the date it is enacted into law.