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 383.

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,

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

- 1 THIS SECTION IF THE DISPENSING PRESCRIBER DISPENSES BUPRENORPHINE,
- 2 OR A DRUG CONTAINING BUPRENORPHINE AND METHADONE, IN A SUBSTANCE
- 3 USE DISORDER PROGRAM AND THE PATIENT PROVIDES CONSENT IN A MANNER
- 4 CONSISTENT WITH SECTION 262 OF THE MENTAL HEALTH CODE, 1974 PA 258,
- 5 MCL 330.1262, AND FEDERAL LAW, TO HAVE THE DATA REPORTED INTO THE
- 6 ELECTRONIC MONITORING SYSTEM FOR THE PURPOSES DESCRIBED IN THIS
- 7 SECTION. A DISPENSING PRESCRIBER WHO RECEIVES THE CONSENT DESCRIBED
- 8 IN THIS SUBDIVISION SHALL MAINTAIN THE PATIENT'S CONSENT FORM AND
- 9 MAKE IT AVAILABLE TO THE DEPARTMENT UPON THE DEPARTMENT'S REQUEST.
- 10 AS USED IN THIS SUBDIVISION:
- 11 (i) "APPROVED SERVICES PROGRAM" MEANS THAT TERM AS DEFINED IN
- 12 SECTION 100A OF THE MENTAL HEALTH CODE, 1974 PA 258, MCL 330.1100A.
- 13 (ii) "SUBSTANCE USE DISORDER PROGRAM" MEANS A PROGRAM AS THAT
- 14 TERM IS DEFINED IN SECTION 260 OF THE MENTAL HEALTH CODE, 1974 PA
- 15 258, MCL 330.1260, AN APPROVED SERVICE PROGRAM, A NONREGULATED
- 16 SUBSTANCE USE DISORDER SERVICES PROGRAM, A FEDERAL CERTIFIED
- 17 SUBSTANCE USE DISORDER SERVICES PROGRAM, OR A FEDERALLY REGULATED
- 18 SUBSTANCE USE DISORDER SERVICES PROGRAM.
- 19 (2) Notwithstanding any practitioner-patient privilege, the
- 20 director of the department may provide data obtained under this
- 21 section to all of the following:
- 22 (a) A designated representative of a board responsible for the
- 23 licensure, regulation, or discipline of a practitioner, pharmacist,
- 24 or other person that is authorized to prescribe, administer, or
- 25 dispense controlled substances.
- 26 (b) An employee or agent of the department.
- 27 (c) A state, federal, or municipal employee or agent whose

- 1 duty is to enforce the laws of this state or the United States
- 2 relating to drugs.
- 3 (d) A state-operated Medicaid program.
- 4 (e) A state, federal, or municipal employee who is the holder
- 5 of a search warrant or subpoena properly issued for the records.
- 6 (f) A practitioner or pharmacist who requests information and
- 7 certifies that the requested information is for the purpose of
- 8 providing medical or pharmaceutical treatment to a bona fide
- 9 current patient.
- 10 (g) An individual with whom the department has contracted
- 11 under subsection (7).
- 12 (h) A practitioner or other person that is authorized to
- 13 prescribe controlled substances for the purpose of determining if
- 14 prescriptions written by that practitioner or other person have
- 15 been dispensed.
- 16 (i) The health care payment or benefit provider for the
- 17 purposes of ensuring patient safety and investigating fraud and
- 18 abuse.
- 19 (3) Except as otherwise provided in this part, a person shall
- 20 use information submitted under this section only for bona fide
- 21 drug-related criminal investigatory or evidentiary purposes or for
- 22 the investigatory or evidentiary purposes in connection with the
- 23 functions of a disciplinary subcommittee or 1 or more of the
- 24 licensing or registration boards created in article 15.
- 25 (4) A person that receives data or any report under subsection
- 26 (2) containing any patient identifiers of the system from the
- 27 department shall not provide it to any other person except by order

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

- 1 (7) The department may enter into 1 or more contractual
- 2 agreements for the administration of this section.
- 3 (8) The department, all law enforcement officers, all officers
- 4 of the court, and all regulatory agencies and officers, in using
- 5 the data for investigative or prosecution purposes, shall consider
- 6 the nature of the prescriber's and dispenser's practice and the
- 7 condition for which the patient is being treated.
- 8 (9) The data and any report containing any patient identifiers
- 9 obtained from the data are not public records and are not subject
- 10 to DISCLOSURE UNDER the freedom of information act, 1976 PA 442,
- 11 MCL 15.231 to 15.246.
- 12 (10) The department may issue a written request to a health
- 13 care payment or benefit provider to determine if the provider has
- 14 accessed the electronic monitoring system as provided in subsection
- 15 (2)(i) in the previous calendar year and, if so, to determine the
- 16 number of inquiries the provider made in the previous calendar year
- 17 and any other information the department requests in relation to
- 18 the provider's access to the electronic monitoring system. A health
- 19 care payment or benefit provider shall respond to the written
- 20 request on or before the March 31 following the request. The
- 21 department shall collaborate with health care payment or benefit
- 22 providers to develop a reasonable request and reporting form for
- 23 use under this subsection.
- 24 (11) R 338.3162E OF THE MICHIGAN ADMINISTRATIVE CODE IS
- 25 RESCINDED.
- 26 (12) (11) As used in this section:
- 27 (a) "Department" means the department of licensing and

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