

## **HOUSE BILL No. 4352**

February 28, 2013, Introduced by Reps. Lyons, Graves, Haines, Walsh and Lipton and referred to the Committee on Education.

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

by amending sections 17745, 17751, 17754, and 17757 (MCL 333.17745, 333.17751, 333.17754, and 333.17757), sections 17745 and 17757 as amended by 2011 PA 210 and sections 17751 and 17754 as amended by 2012 PA 209, and by adding section 17744a.

## THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

- 1 SEC. 17744A. (1) NOTWITHSTANDING ANY PROVISION OF THIS ACT TO
- 2 THE CONTRARY, A PRESCRIBER MAY ISSUE A PRESCRIPTION FOR AND A
- 3 DISPENSING PRESCRIBER OR PHARMACIST MAY DISPENSE AUTO-INJECTABLE
  - EPINEPHRINE TO A SCHOOL BOARD FOR THE PURPOSE OF MEETING THE
- REQUIREMENTS OF SECTION 1179A OF THE REVISED SCHOOL CODE, 1976 PA
- 6 451, MCL 380.1179A. WHEN ISSUING A PRESCRIPTION FOR OR DISPENSING
- 7 AUTO-INJECTABLE EPINEPHRINE TO A SCHOOL BOARD AS AUTHORIZED UNDER

House Bill No. 4352 as amended October 22, 2013 THIS SECTION, THE PRESCRIBER, DISPENSING PRESCRIBER, OR PHARMACIST, 1 AS APPROPRIATE, SHALL INSERT THE NAME OF THE SCHOOL BOARD AS THE 2 3 PROFESSIONAL NURSE OR WHO IS TRAINED IN THE ADMINISTRATION OF AN 4

NAME OF THE PATIENT. A SCHOOL EMPLOYEE WHO IS A LICENSED REGISTERED

EPINEPHRINE AUTO-INJECTOR UNDER SECTION 1179A OF THE REVISED SCHOOL 5

CODE, 1976 PA 451, MCL 380.1179A, MAY POSSESS AND ADMINISTER AN 6

EPINEPHRINE AUTO-INJECTOR DISPENSED TO A SCHOOL BOARD UNDER THIS

8 SECTION.

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[(2) A PRESCRIBER WHO ISSUES A PRESCRIPTION FOR OR A DISPENSING 9 PRESCRIBER OF PHARMACIST WHO DISPENSES AUTO-INJECTABLE EPINEPHRINE TO A 10 SCHOOL BOARD AS AUTHORIZED UNDER THIS SECTION IS NOT LIABLE IN A CIVIL 11 ACTION FOR A PROPERLY STORED AND DISPENSED EPINEPHRINE AUTO-INJECTOR 12 13 THAT WAS A PROXIMATE CAUSE OF INJURY OR DEATH TO AN INDIVIDUAL DUE TO 14 THE ADMINISTRATION OF OR FAILURE TO ADMINISTER THE EPINEPHRINE AUTO-15 INJECTOR.

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Sec. 17745. (1) Except as otherwise provided in this subsection, a prescriber who wishes to dispense prescription drugs shall obtain from the board a drug control license for each location in which the storage and dispensing of prescription drugs occur. A drug control license is not necessary if the dispensing occurs in the emergency department, emergency room, or trauma center of a hospital licensed under article 17 or if the dispensing involves only the issuance of complimentary starter dose drugs.

- 1 (2) A-EXCEPT AS OTHERWISE PROVIDED IN SECTION 17744A, A
- 2 dispensing prescriber shall dispense prescription drugs only to his
- 3 or her own patients.
- 4 (3) A dispensing prescriber shall include in a patient's chart
- 5 or clinical record a complete record, including prescription drug
- 6 names, dosages, and quantities, of all prescription drugs dispensed
- 7 directly by the dispensing prescriber or indirectly under his or
- 8 her delegatory authority. If prescription drugs are dispensed under
- 9 the prescriber's delegatory authority, the delegatee who dispenses
- 10 the prescription drugs shall initial the patient's chart, clinical
- 11 record, or log of prescription drugs dispensed. In a patient's
- 12 chart or clinical record, a dispensing prescriber shall distinguish
- 13 between prescription drugs dispensed to the patient, and
- 14 prescription drugs prescribed for the patient, AND PRESCRIPTION
- 15 DRUGS DISPENSED OR PRESCRIBED AS AUTHORIZED UNDER SECTION 17744A. A
- 16 dispensing prescriber shall retain information required under this
- 17 subsection for not less than 5 years after the information is
- 18 entered in the patient's chart or clinical record.
- 19 (4) A dispensing prescriber shall store prescription drugs
- 20 under conditions that will maintain their stability, integrity, and
- 21 effectiveness and will assure that the prescription drugs are free
- 22 of contamination, deterioration, and adulteration.
- 23 (5) A dispensing prescriber shall store prescription drugs in
- 24 a substantially constructed, securely lockable cabinet. Access to
- 25 the cabinet shall be limited to individuals authorized to dispense
- 26 prescription drugs in compliance with this part and article 7.
- 27 (6) Unless otherwise requested by a patient, a dispensing

- 1 prescriber shall dispense a prescription drug in a safety closure
- 2 container that complies with the poison prevention packaging act of
- **3** 1970, 15 USC 1471 to 1477.
- 4 (7) A dispensing prescriber shall dispense a drug in a
- 5 container that bears a label containing all of the following
- 6 information:
- 7 (a) The name and address of the location from which the
- 8 prescription drug is dispensed.
- 9 (b) The EXCEPT AS OTHERWISE AUTHORIZED UNDER SECTION 17744A,
- 10 THE patient's name and record number.
- 11 (c) The date the prescription drug was dispensed.
- 12 (d) The prescriber's name or, if dispensed under the
- 13 prescriber's delegatory authority, shall list the name of the
- **14** delegatee.
- 15 (e) The directions for use.
- 16 (f) The name and strength of the prescription drug.
- 17 (g) The quantity dispensed.
- 18 (h) The expiration date of the prescription drug or the
- 19 statement required under section 17756.
- 20 (8) A dispensing prescriber who dispenses a complimentary
- 21 starter dose drug to a patient shall give the patient at least all
- 22 of the following information, either by dispensing the
- 23 complimentary starter dose drug to the patient in a container that
- 24 bears a label containing the information or by giving the patient a
- 25 written document which THAT may include, but is not limited to, a
- 26 preprinted insert that comes with the complimentary starter dose
- 27 drug, that contains ALL OF the FOLLOWING information:

- 1 (a) The name and strength of the complimentary starter dose
- 2 drug.
- 3 (b) Directions for the patient's use of the complimentary
- 4 starter dose drug.
- 5 (c) The expiration date of the complimentary starter dose drug
- 6 or the statement required under section 17756.
- 7 (9) The information required under subsection (8) is in
- 8 addition to, and does not supersede or modify, other state or
- 9 federal law regulating the labeling of prescription drugs.
- 10 (10) In addition to meeting the requirements of this part, a
- 11 dispensing prescriber who dispenses controlled substances shall
- 12 comply with section 7303a.
- 13 (11) The board may periodically inspect locations from which
- 14 prescription drugs are dispensed.
- 15 (12) The act, task, or function of dispensing prescription
- 16 drugs shall be delegated only as provided in this part and sections
- 17 16215, 17048, 17076, 17212, and 17548.
- 18 (13) A supervising physician may delegate in writing to a
- 19 pharmacist practicing in a hospital pharmacy within a hospital
- 20 licensed under article 17 the receipt of complimentary starter dose
- 21 drugs other than controlled substances as defined by article 7 or
- 22 federal law. When the delegated receipt of complimentary starter
- 23 dose drugs occurs, both the pharmacist's name and the supervising
- 24 physician's name shall be used, recorded, or otherwise indicated in
- 25 connection with each receipt. A pharmacist described in this
- 26 subsection may dispense a prescription for complimentary starter
- 27 dose drugs written or transmitted by facsimile, electronic

- 1 transmission, or other means of communication by a prescriber.
- 2 (14) As used in this section, "complimentary starter dose"
- 3 means a prescription drug packaged, dispensed, and distributed in
- 4 accordance with state and federal law that is provided to a
- 5 dispensing prescriber free of charge by a manufacturer or
- 6 distributor and dispensed free of charge by the dispensing
- 7 prescriber to his or her patients.
- 8 Sec. 17751. (1) A pharmacist shall not dispense a drug
- 9 requiring a prescription under the federal act or a law of this
- 10 state except under authority of an original prescription or an
- 11 equivalent record of an original prescription approved by the
- 12 board.
- 13 (2) Subject to subsection (5), a pharmacist may dispense a
- 14 prescription written and signed; written or created in an
- 15 electronic format, signed, and transmitted by facsimile; or
- 16 transmitted electronically or by other means of communication by a
- 17 physician prescriber or dentist prescriber in a state other than
- 18 Michigan, but not including a prescription for a controlled
- 19 substance as defined in section 7104 except under circumstances
- 20 described in section 17763(e), only if the pharmacist in the
- 21 exercise of his or her professional judgment determines all of the
- 22 following:
- 23 (a) That EXCEPT AS OTHERWISE AUTHORIZED UNDER SECTION 17744A,
- 24 THAT the prescription was issued pursuant to an existing physician-
- 25 patient or dentist-patient relationship.
- 26 (b) That the prescription is authentic.
- (c) That the prescribed drug is appropriate and necessary for

- 1 the treatment of an acute, chronic, or recurrent condition.
- 2 (3) A pharmacist or a prescriber shall dispense a prescription
- 3 only if the prescription falls within the scope of practice of the
- 4 prescriber.
- 5 (4) A pharmacist shall not knowingly dispense a prescription
- 6 after the death of the prescriber or patient.
- 7 (5) A pharmacist shall not dispense a drug or device under a
- 8 prescription transmitted by facsimile or created in electronic
- 9 format and printed out for use by the patient unless the document
- 10 is manually signed by the prescriber. This subsection does not
- 11 apply to a prescription that is transmitted by a computer to a
- 12 facsimile machine if that prescription complies with section 17754.
- 13 (6) After consultation with and agreement from the prescriber,
- 14 a pharmacist may add or change a patient's address, dosage form,
- 15 drug strength, drug quantity, directions for use, or issue date
- 16 with regard to a prescription. A pharmacist shall note the details
- 17 of the consultation and agreement required under this subsection on
- 18 the prescription and shall maintain that documentation with the
- 19 prescription as required in section 17752. A pharmacist shall not
- 20 change the patient's name, controlled substance prescribed unless
- 21 authorized to dispense a lower cost generically equivalent drug
- 22 product under section 17755, or the prescriber's signature with
- 23 regard to a prescription.
- 24 (7) A prescription that is contained within a patient's chart
- 25 in a health facility or agency licensed under article 17 or other
- 26 medical institution and that is transmitted to a pharmacy under
- 27 section 17744 is the original prescription. If all other

- 1 requirements of this part are met, a pharmacist shall dispense a
- 2 drug or device under a prescription described in this subsection. A
- 3 pharmacist may dispense a drug or device under a prescription
- 4 described in this subsection even if the prescription does not
- 5 contain the quantity ordered. If a prescription described in this
- 6 subsection does not contain the quantity ordered, the pharmacist
- 7 shall consult with the prescriber to determine an agreed-upon
- 8 quantity. The pharmacist shall record the quantity dispensed on the
- 9 prescription and shall maintain that documentation with the
- 10 prescription as required in section 17752.
- 11 Sec. 17754. (1) Except as otherwise provided under article 7
- 12 and the federal act, a prescription may be transmitted
- 13 electronically as long as IF the prescription is transmitted in
- 14 compliance with the health insurance portability and accountability
- 15 act of 1996, Public Law 104-191, or regulations promulgated under
- 16 that act, 45 CFR parts 160 and 164, by a prescriber or his or her
- 17 agent and the data are not altered or modified in the transmission
- 18 process. The electronically transmitted prescription shall include
- 19 all of the following information:
- (a) The name, address, and telephone number of the prescriber.
- 21 (b) The EXCEPT AS OTHERWISE AUTHORIZED UNDER SECTION 17744A,
- 22 THE full name of the patient for whom the prescription is issued.
- 23 (c) An electronic signature or other identifier that
- 24 specifically identifies and authenticates the prescriber or his or
- 25 her agent.
- 26 (d) The time and date of the transmission.
- (e) The identity of the pharmacy intended to receive the

- 1 transmission.
- 2 (f) Any other information required by the federal act or state
- 3 law.
- 4 (2) The electronic equipment or system utilized in the
- 5 transmission and communication of prescriptions shall provide
- 6 adequate confidentiality safeguards and be maintained to protect
- 7 patient confidentiality as required under any applicable federal
- 8 and state law and to ensure against unauthorized access. The
- 9 electronic transmission of a prescription shall be communicated in
- 10 a retrievable, recognizable form acceptable to the intended
- 11 recipient. The electronic form utilized in the transmission of a
- 12 prescription shall not include "dispense as written" or "d.a.w." as
- 13 the default setting.
- 14 (3) Prior to BEFORE dispensing a prescription that is
- 15 electronically transmitted, the pharmacist shall exercise
- 16 professional judgment regarding the accuracy, validity, and
- 17 authenticity of the transmitted prescription.
- 18 (4) An electronically transmitted prescription that meets the
- 19 requirements of this section is the original prescription.
- 20 Sec. 17757. (1) Upon a request made in person or by telephone,
- 21 a pharmacist engaged in the business of selling drugs at retail
- 22 shall provide the current selling price of a drug dispensed by that
- 23 pharmacy or comparative current selling prices of generic and brand
- 24 name drugs dispensed by that pharmacy. The information shall be
- 25 provided to the person making the request before a drug is
- 26 dispensed to the person. A person who makes a request for price
- 27 information under this subsection shall—IS not be—obligated to

- 1 purchase the drug for which the price or comparative prices are
- 2 requested.
- 3 (2) A pharmacist engaged in the business of selling drugs at
- 4 retail shall conspicuously display the notice described in
- 5 subsection (3) at each counter over which prescription drugs are
- 6 dispensed.
- 7 (3) The notice required under subsection (2) shall be in
- 8 substantially the following form:
- 9 NOTICE TO CONSUMERS
- 10 ABOUT PRESCRIPTION DRUGS
- 11 Under Michigan law, you have the right to find out the price
- 12 of a prescription drug before the pharmacist fills the
- 13 prescription. You are under no obligation to have the prescription
- 14 filled here and may use this price information to shop around at
- 15 other pharmacies. You may request price information in person or by
- 16 telephone.
- 17 Every pharmacy has the current selling prices of both generic
- 18 and brand name drugs dispensed by the pharmacy.
- 19 Ask your pharmacist if a lower-cost generic drug is available
- 20 to fill your prescription. A generic drug contains the same
- 21 medicine as a brand name drug and is a suitable substitute in most
- 22 instances.
- 23 A generic drug may not be dispensed by your pharmacist if your
- 24 doctor has written "dispense as written" or the initials "d.a.w."
- 25 on the prescription.
- 26 If you have questions about the drugs which THAT have been
- 27 prescribed for you, ask your doctor or pharmacist for more

- 1 information.
- 2 To avoid dangerous drug interactions, let your doctor and
- 3 pharmacist know about any other medications you are taking. This is
- 4 especially important if you have more than 1 doctor or have
- 5 prescriptions filled at more than 1 pharmacy.
- 6 (4) The notice required under subsection (2) shall also
- 7 contain the address and phone number of the board and the
- 8 department. The text of the notice shall be in at least 32-point
- 9 bold type and shall be printed on paper at least 11 inches by 17
- 10 inches in size. The notice may be printed on multiple pages.
- 11 (5) A copy of the notice required under subsection (2) shall
- 12 be provided to each licensee by the department. Additional THE
- 13 DEPARTMENT SHALL PROVIDE ADDITIONAL copies shall be available if
- 14 needed. from the department. A person may duplicate or reproduce
- 15 the notice if the duplication or reproduction is a true copy of the
- 16 notice as produced by the department, without any additions or
- 17 deletions. whatsoever.
- 18 (6) The pharmacist shall furnish to the purchaser of a
- 19 prescription drug at the time the drug is delivered to the
- 20 purchaser a receipt evidencing the transactions, which contains ALL
- 21 OF the following:
- (a) The brand name of the drug, if applicable.
- 23 (b) The name of the manufacturer or the supplier of the drug,
- 24 if the drug does not have a brand name.
- 25 (c) The strength of the drug, if significant.
- 26 (d) The quantity dispensed, if applicable.
- (e) The name and address of the pharmacy.

- 1 (f) The serial number of the prescription.
- 2 (g) The date the prescription was originally dispensed.
- 3 (h) The name of the prescriber or, if prescribed under the
- 4 prescriber's delegatory authority, shall list the name of the
- 5 delegatee.
- 6 (i) The EXCEPT AS OTHERWISE AUTHORIZED UNDER SECTION 17744A,
- 7 THE name of THE patient for whom the drug was prescribed.
- 8 (j) The price for which the drug was sold to the purchaser.
- 9 (7) Subsection THE ITEMS REQUIRED UNDER SUBSECTION (6)(a),
- 10 (b), and (c) may be omitted by a pharmacist only if the omission is
- 11 expressly required by the prescriber. The pharmacist shall retain a
- 12 copy of each receipt for 90 days. The inclusion of THE ITEMS
- 13 REQUIRED UNDER subsection (6) on the prescription container label
- 14 is a valid receipt to the purchaser. Including THE ITEMS REQUIRED
- 15 UNDER subsection (6) on the written prescription form and retaining
- 16 the form constitutes retention of a copy of the receipt.
- 17 (8) The board may promulgate rules to implement this section.
- 18 Enacting section 1. This amendatory act does not take effect
- 19 unless Senate Bill No. or House Bill No. 4353 (request no.
- 20 01326'13) of the 97th Legislature is enacted into law.

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