

HOUSE BILL NO. 4352

February 24, 2021, Introduced by Reps. Allor, Brann, Rendon, Whiteford, Borton, Glenn, Farrington, Bellino, Yaroach, Wozniak, O'Malley and Calley and referred to the Committee on Health Policy.

A bill to amend 1978 PA 368, entitled
"Public health code,"
by amending section 17757 (MCL 333.17757), as amended by 2016 PA
383, and by adding section 17757b.

THE PEOPLE OF THE STATE OF MICHIGAN ENACT:

1 telephone.

2 Every pharmacy has the current selling prices of both generic
3 and brand name drugs dispensed by the pharmacy.

4 Ask your pharmacist if a lower-cost generic drug is available
5 to fill your prescription. A generic drug contains the same
6 medicine as a brand name drug and is a suitable substitute in most
7 instances.

8 A generic drug may not be dispensed by your pharmacist if your
9 doctor has written "dispense as written" or the initials "d.a.w."
10 on the prescription.

11 If you have questions about the drugs that have been
12 prescribed for you, ask your doctor or pharmacist for more
13 information.

14 To avoid dangerous drug interactions, let your doctor and
15 pharmacist know about any other medications you are taking. This is
16 especially important if you have more than 1 doctor or have
17 prescriptions filled at more than 1 pharmacy.

18 (4) The notice required under subsection (2) must also contain
19 the address and phone number of the board and the department. The
20 text of the notice must be in at least 32-point bold type and ~~must~~
21 be printed on paper at least 11 inches by 17 inches in size. The
22 notice may be printed on multiple pages.

23 (5) The department shall provide a copy of the notice required
24 under subsection (2) to each licensee. The department shall provide
25 additional copies if needed. A person may duplicate or reproduce
26 the notice if the duplication or reproduction is a true copy of the
27 notice as produced by the department, without any additions or
28 deletions.

29 (6) The pharmacist shall furnish to the purchaser of a

1 prescription drug at the time the drug is delivered to the
2 purchaser a receipt evidencing the transactions that contains all
3 of the following:

4 (a) The brand name of the drug, if applicable.

5 (b) The name of the manufacturer or the supplier of the drug,
6 if the drug does not have a brand name.

7 (c) The strength of the drug, if significant.

8 (d) The quantity dispensed, if applicable.

9 (e) The name and address of the pharmacy.

10 (f) The serial number of the prescription or a reference to
11 the standing order issued under section 17744e.

12 (g) The date the prescription was originally dispensed.

13 (h) The name of the prescriber or, if prescribed under the
14 prescriber's delegatory authority, the name of the delegatee.

15 (i) Except as otherwise authorized under section 5110, 17744a,
16 17744b, or 17744e, the name of the patient for whom the drug was
17 prescribed.

18 (j) The price for which the drug was sold to the purchaser.

19 (7) The items required under subsection (6) (a), (b), and (c)
20 may be omitted from a receipt by a pharmacist only if the omission
21 is expressly required by the prescriber. The pharmacist shall
22 retain a copy of each receipt furnished under subsection (6) for 90
23 days. The inclusion of the items required under subsection (6) on
24 the prescription container label is a valid receipt to the
25 purchaser. Including the items required under subsection (6) on the
26 written prescription form and retaining the form constitutes
27 retention of a copy of the receipt.

28 (8) The department, in consultation with the board, may
29 promulgate rules to implement this section.

1 Sec. 17757b. (1) A pharmacy or pharmacist engaged in the
2 business of selling drugs shall not enter into a contract with a
3 pharmacy benefit manager that violates section 26 of the third
4 party administrator act, 1984 PA 218, MCL 550.926, or that prevents
5 or interferes with in any manner a patient's choice to receive an
6 eligible prescription drug from a 340b entity or a pharmacy when
7 dispensing a 340b drug.

8 (2) As used in this section:

9 (a) "340b drug" means a covered drug as that term is defined
10 in 42 USC 256b.

11 (b) "340b entity" means a covered entity as that term is
12 defined in 42 USC 256b.

13 (c) "Pharmacy benefit manager" means that term as defined in
14 section 2 of the third party administrator act, 1984 PA 218, MCL
15 550.902.

16 Enacting section 1. This amendatory act does not take effect
17 unless Senate Bill No. ____ or House Bill No. 4351 (request no.
18 00166'21) of the 101st Legislature is enacted into law.