

# SENATE BILL NO. 317

May 20, 2025, Introduced by Senators JOHNSON, DAMOOSE, VICTORY, DALEY, LINDSEY, WEBBER and CHANG and referred to Committee on Health Policy.

A bill to allow for the establishment of a wholesale prescription drug importation program; to provide for the powers and duties of certain state and local governmental officers and entities; and to allow for the promulgation of rules.

## **THE PEOPLE OF THE STATE OF MICHIGAN ENACT:**

- 1       Sec. 1. This act may be cited as the "prescription drug
- 2       importation act".
- 3       Sec. 3. As used in this act:
- 4       (a) "Canadian supplier" means a Canadian prescription drug

1 supplier that is regulated under the laws of Canada or a province  
2 of Canada.

3 (b) "Department" means the department of health and human  
4 services.

5 (c) "Eligible importer" means a licensed drug wholesaler that  
6 contracts with the department under section 5.

7 (d) "Licensed drug wholesaler" means a wholesale distributor  
8 as that term is defined in section 17709 of the public health code,  
9 1978 PA 368, MCL 333.17709.

10 (e) "Pharmacy" means that term as defined in section 17707 of  
11 the public health code, 1978 PA 368, MCL 333.17707.

12 (f) "Prescription drug" means that term as defined in section  
13 17708 of the public health code, 1978 PA 368, MCL 333.17708.

14 (g) "Program" means the wholesale prescription drug  
15 importation program developed in section 5.

16 Sec. 5. The department, in consultation with interested  
17 stakeholders and appropriate federal officials, may develop a  
18 wholesale prescription drug importation program. The program must  
19 meet all of the following requirements:

20 (a) Comply with the applicable requirements of 21 USC 384,  
21 including requirements on safety and cost savings.

22 (b) Require the department to contract with a licensed drug  
23 wholesaler for the purposes of seeking federal certification and  
24 approval to import prescription drugs into this state.

25 (c) Require the use of a Canadian supplier.

26 (d) Ensure that only prescription drugs that meet safety,  
27 effectiveness, and other standards of the United States Food and  
28 Drug Administration are imported by or on behalf of this state.

29 (e) Ensure that the program does not import generic

1 prescription drugs that would violate federal patent laws on brand  
2 name prescription drugs in the United States.

3 (f) Ensure that only prescription drugs that are expected to  
4 result in a significant reduction in the cost of those prescription  
5 drugs to consumers in this state are imported.

6 (g) Ensure that the tracking and tracing requirements of 21  
7 USC 360eee and 360eee-1 are complied with to the extent possible  
8 before imported prescription drugs come into the possession of an  
9 eligible importer and that the tracking and tracing requirements of  
10 21 USC 360eee and 360eee-1 are fully complied with after imported  
11 prescription drugs come into the possession of an eligible  
12 importer.

13 (h) Ensure that any prescription drug imported under the  
14 program is not distributed, dispensed, or sold outside of this  
15 state.

16 (i) Establish policies and procedures for record keeping and  
17 that allow for a periodic audit of the records to ensure the  
18 department's compliance with this section.

19 Sec. 7. (1) Before implementing the program, the department  
20 shall seek certification of the program from the secretary of the  
21 United States Department of Health and Human Services and, on  
22 receiving the certification, shall do all of the following in  
23 implementing the program:

24 (a) Comply with the program requirements described in section  
25 5.

26 (b) Develop a registration process for health insurers,  
27 pharmacies, and health care providers who administer prescription  
28 drugs, that are willing to participate in the program.

29 (c) Create and maintain a list of the wholesale acquisition

1 cost of each prescription drug that is imported under the program.  
2 The department shall make the list available to the public and the  
3 persons described in subdivision (b).

4 (d) Develop and implement an outreach and marketing plan to  
5 generate awareness about the program.

6 (e) Conduct any other activity that the department considers  
7 important for the successful implementation of the program.

8 (2) The department shall also seek the appropriate federal  
9 approval, waiver, exemption, or agreement necessary to allow a  
10 covered entity enrolled in or eligible for the federal 340B program  
11 to participate in this state's program to the fullest extent  
12 possible without jeopardizing eligibility for the federal 340B  
13 program. As used in this subsection, "federal 340B program" means  
14 the 340B drug pricing program established under section 602 of the  
15 veterans health care act of 1992, Public Law 102-585.

16 Sec. 9. (1) Beginning on the first January 15 after the date  
17 of the program's implementation, and annually after that, the  
18 department shall do all of the following:

19 (a) Prepare a report on the operation of the program during  
20 the previous calendar year.

21 (b) Publish the report described in subdivision (a) on the  
22 department's website in a location that is available to the public.

23 (c) Submit a copy of the report described in subdivision (a)  
24 to the senate and house of representatives standing committees on  
25 health policy.

26 (2) The report described in subsection (1) must include all of  
27 the following information for the calendar year covered by the  
28 report:

29 (a) A list of each prescription drug that was included in the

1 program.

2 (b) The number of pharmacies, health care providers, and  
3 health insurance plans that participated in the program.

4 (c) The number of prescriptions dispensed under the program.

5 (d) The estimated cost savings to consumers, health insurers,  
6 employers, and this state. The report must also include the total  
7 estimated cost savings to consumers, health insurance plans,  
8 employers, and this state since the date of the program's  
9 implementation.

10 (e) Information on the implementation of the auditing  
11 procedure developed by the department and any audit findings.

12 (f) Any other information that the department considers  
13 relevant.

14 Sec. 11. The department shall consult with the department of  
15 attorney general to identify the potential for and to monitor  
16 anticompetitive behavior in industries that are affected by a  
17 wholesale prescription drug importation program.

18 Sec. 13. The department may promulgate rules under the  
19 administrative procedures act of 1969, 1969 PA 306, MCL 24.201 to  
20 24.328, to implement this act.