



Sen. David Koehler

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1 AMENDMENT TO SENATE BILL 2385

2 AMENDMENT NO. _____. Amend Senate Bill 2385 by replacing
3 everything after the enacting clause with the following:

4 "Section 1. Short title. This Act may be cited as the
5 Patient Access to Pharmacy Protection Act.

6 Section 5. Findings. The General Assembly finds that:

7 (1) It is within the traditional authority of the State to
8 regulate the acquisition and delivery of drugs to pharmacies
9 and providers.

10 (2) The federal 340B statute is silent on distribution of
11 340B-acquired drugs to 340B covered entities and their
12 contract pharmacy partners.

13 (3) The State's compelling interest in preserving and
14 improving access to health care services requires it to ensure
15 that 340B covered entities continue to be allowed to contract
16 with pharmacies to receive 340B drugs and dispense them to the

1 patients of 340B covered entities in accordance with federal
2 law.

3 (4) Addressing accessibility of these life-saving
4 medications is a matter of health, safety, and welfare for the
5 people of the State of Illinois.

6 Section 10. Definitions. As used in this Act:

7 "340B drug discount program" means the program established
8 under Section 340B of the federal Public Health Service Act,
9 42 U.S.C. 256b.

10 "340B contract pharmacy" means any pharmacy that is under
11 contract with a 340B covered entity to dispense 340B drugs on
12 behalf of the 340B covered entity and is either (i) located in
13 Illinois and qualifies as a pharmacy under Section 3 of the
14 Pharmacy Practice Act; or (ii) is located in a state,
15 commonwealth, or territory of the United States, other than
16 Illinois, and dispenses 340B drugs on behalf of the 340B
17 covered entity.

18 "340B covered entity" means an entity in Illinois that
19 qualifies as a covered entity under Section 340B of the
20 federal Public Health Service Act, 42 U.S.C. 256b(a)(4).

21 "340B drug" means a drug that has been subject to any offer
22 for reduced prices by a manufacturer pursuant to 42 U.S.C.
23 256b and is purchased by a 340B covered entity.

24 "340B grantee" means an entity in Illinois that qualifies
25 as a covered entity under subparagraphs (A)-(K) of paragraph

1 (4) of subsection (a) of Section 340B of the federal Public
2 Health Service Act, 42 U.S.C. 256b(a)(4)(A)-(K).

3 "Charity care" has the meaning given to that term in line
4 23 of worksheet 5-10 to the Medicare cost report or in any
5 successor form.

6 "Children's hospital" has the meaning given to that term
7 in Section 5-5.02 of the Illinois Public Aid Code.

8 "Critical Access Hospital" has the meaning given to that
9 term in paragraph (4) of subsection (b) of Section 5-5e of the
10 Illinois Public Aid Code.

11 "Hospital" means a hospital licensed under the Hospital
12 Licensing Act or University of Illinois Hospital Act.

13 "Manufacturer" or "Pharmaceutical Manufacturer" has the
14 meaning given to the term "manufacturer" in the Wholesale Drug
15 Distribution Licensing Act.

16 "Person" includes a natural person, partnership,
17 association, corporation, or any other legal business entity.
18 "Person" does not include any federal or State government
19 entity or body.

20 "Safety-Net Hospital" has the meaning given to that term
21 in Section 5-5e.1 of the Illinois Public Aid Code.

22 Section 15. Protection of patient access to pharmacy.

23 (a) No person, including a pharmaceutical manufacturer,
24 may deny, restrict, prohibit, condition, or otherwise
25 interfere with, either directly or indirectly, the acquisition

1 of a 340B drug by, or delivery of a 340B drug to, a 340B
2 covered entity or a 340B contract pharmacy authorized to
3 receive 340B drugs on behalf of the 340B covered entity unless
4 the receipt is prohibited by federal law.

5 (b) No person, including a pharmaceutical manufacturer,
6 may impose any restriction on the ability of a 340B covered
7 entity to contract with or designate a 340B contract pharmacy,
8 including restrictions relating to the number, location,
9 ownership, or type of 340B contract pharmacy.

10 (c) Each individual transaction, as defined in 21 U.S.C.
11 360eee-24, of 340B drugs that is subject to a prohibited act in
12 subsections (a) and (b) shall constitute a separate violation
13 of this Act.

14 Section 20. Reporting. On or before August 1, 2026 and
15 each August 1 thereafter, a 340B covered entity shall submit a
16 report to the General Assembly pursuant to this Section. For
17 the purposes of this Section, the following covered entities
18 are exempt until January 1, 2028 and will report on or before
19 August 1, 2028 and each August 1 thereafter: hospitals with
20 fewer than 100 licensed beds, Critical Access Hospitals,
21 Safety-Net Hospitals, children's hospitals, and 340B grantees.
22 The report must include all of the following for the 340B
23 covered entity's 340B program:

24 (1) the name of the 340B covered entity submitting the
25 report;

1 (2) a copy of the 340B covered entity's annual 340B
2 program recertification;

3 (3) whether a community benefits plan report is
4 required under Section 20 of the Community Benefits Act
5 and, if so, a copy of the 340B covered entity's community
6 benefits plan report;

7 (4) the aggregate acquisition cost for prescription
8 drugs obtained under the 340B program and dispensed or
9 administered to patients;

10 (5) the aggregate payment amount received for all
11 drugs obtained under the 340B program and dispensed or
12 administered to patients;

13 (6) the aggregate payments made to contract pharmacies
14 to dispense drugs obtained under the 340B program;

15 (7) the 340B covered entity's total costs for charity
16 care;

17 (8) the number of claims for prescription drugs
18 received under the 340B program;

19 (9) the percentage of the 340B covered entity's claims
20 that were for prescription drugs obtained under the 340B
21 program;

22 (10) a description of any adverse 340B program audits
23 within the preceding 12 months; and

24 (11) a description of the impact of the 340B program
25 on the patients and the community served by the 340B
26 covered entity.

1 Section 25. Medicaid study.

2 (a) By January 1, 2028, the Department of Healthcare and
3 Family Services shall report to the General Assembly on the
4 following for the total aggregated covered outpatient drug
5 units dispensed or administered in this State for the prior
6 calendar year in connection with the medical assistance
7 program under the Illinois Public Aid Code, categorized by (i)
8 fee-for-service and (ii) each managed care plan:

9 (1) the number of dispensed or administered covered
10 outpatient drug units;

11 (2) the number of dispensed or administered covered
12 outpatient drug units that were subject to a rebate under
13 42 U.S.C. 1396r-8; and

14 (3) a reasonable estimate of net costs or savings to
15 the State's medical assistance program due to 340B covered
16 entity purchases of covered outpatient drug units at 340B
17 pricing.

18 (b) To the extent the Department of Healthcare and Family
19 Services lacks information to provide a data element required
20 under subsection (a), it shall provide a reasonable estimate
21 based on all available information and an explanation of the
22 information that it lacks.

23 Section 30. 340B prescription drug applicability. Each
24 340B covered entity shall dispense or administer 340B drugs

1 only when in connection with an outpatient health care service
2 received by the patient within the last 18 months and
3 furnished at a registered location of the 340B covered entity.

4 Section 35. Preventing duplication of 340B discounts. Each
5 340B covered entity shall develop and maintain a policy that
6 ensures it is not placing an order for a 340B drug to replenish
7 a prior pharmacy dispense if any other 340B covered entity
8 will place an order for a 340B drug to replenish the same prior
9 pharmacy dispense. The policy shall also include a process to
10 reimburse a manufacturer for any duplicate 340B discount the
11 covered entity receives. The policy shall be filed annually
12 with the Attorney General.

13 Section 40. Preemption.

14 (a) Nothing in this Act shall be construed or applied to be
15 less restrictive than federal law for a person regulated by
16 this Act.

17 (b) Nothing in this Act shall be construed or applied in a
18 manner that would conflict with:

19 (1) applicable federal law; or

20 (2) other laws of this State if the State law is
21 compatible with applicable federal law.

22 (c) Limited distribution of a drug required under 21
23 U.S.C. 355-1 may not to be construed as a violation of this
24 Act.

1 Section 97. Severability. If any provision of this Act or
2 its application to any person or circumstance is held invalid,
3 the invalidity of that provision or application does not
4 affect other provisions or applications of this Act that can
5 be given effect without the invalid provision or application.
6 Each paragraph defining "340B contract pharmacy" in Section 10
7 is severable.

8 Section 99. Effective date. This Act takes effect upon
9 becoming law."