

1-1 By: Perry S.B. No. 241
 1-2 (In the Senate - Filed November 16, 2022; February 15, 2023,
 1-3 read first time and referred to Committee on Health & Human
 1-4 Services; March 20, 2023, reported adversely, with favorable
 1-5 Committee Substitute by the following vote: Yeas 9, Nays 0;
 1-6 March 20, 2023, sent to printer.)

1-7 COMMITTEE VOTE

	Yea	Nay	Absent	PNV
1-8				
1-9	X			
1-10	X			
1-11	X			
1-12	X			
1-13	X			
1-14	X			
1-15	X			
1-16	X			
1-17	X			

1-18 COMMITTEE SUBSTITUTE FOR S.B. No. 241 By: Hancock

1-19 A BILL TO BE ENTITLED
 1-20 AN ACT

1-21 relating to written notification provided by drug manufacturers
 1-22 regarding the cause of generic or biosimilar insulin prescription
 1-23 drug unavailability.

1-24 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF TEXAS:

1-25 SECTION 1. Chapter 439, Health and Safety Code, is amended
 1-26 by adding Subchapter D to read as follows:

1-27 SUBCHAPTER D. INSULIN

1-28 Sec. 439.101. DEFINITION. In this subchapter,
 1-29 "manufacturer" has the meaning assigned by Section 531.070,
 1-30 Government Code.

1-31 Sec. 439.102. WRITTEN VERIFICATION REQUIRED FOR BRAND NAME
 1-32 INSULIN DRUG MANUFACTURER. (a) The manufacturer of a brand name
 1-33 insulin prescription drug for which a generic or biosimilar
 1-34 prescription drug is not available and that is included in the
 1-35 Medicaid vendor drug program formulary must submit to the Health
 1-36 and Human Services Commission a written verification stating
 1-37 whether or not the unavailability of the generic or biosimilar
 1-38 prescription drug is the result, wholly or partly, of:

1-39 (1) a scheme by the manufacturer to pay a generic or
 1-40 biosimilar prescription drug manufacturer to delay marketing the
 1-41 generic or biosimilar drug;

1-42 (2) a legal or business strategy to extend the life of
 1-43 a patent on the brand name prescription drug;

1-44 (3) the manufacturer directly manipulating a patent on
 1-45 the brand name prescription drug; or

1-46 (4) the manufacturer facilitating an action described
 1-47 by Subdivisions (1)-(3) on behalf of another entity.

1-48 (b) The executive commissioner shall adopt rules
 1-49 prescribing the form and manner for submission of the written
 1-50 verification required under Subsection (a).

1-51 SECTION 2. This Act takes effect September 1, 2024.

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