As Passed by the House

136th General Assembly

Regular Session

Sub. H. B. No. 324

2025-2026

Representatives Mathews, A., Craig

Cosponsors: Representatives Schmidt, Gross, King, Miller, M., Stewart, Barhorst, Bird, Click, Dovilla, Fowler Arthur, Hall, T., John, Mathews, T., McClain, Newman, Odioso, Richardson, Salvo, Thomas, D., Williams, Young

То	enact section 3715.39 of the Revised Code to	1
	establish conditions on the prescribing of	2
	prescription drugs causing severe adverse	3
	effects and to name this act the Patient	4
	Protection Act.	5

BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF OHIO:

Section 1. That section 3715.39 of the Revised Code be	6
enacted to read as follows:	7
Sec. 3715.39. (A) As used in this section:	8
(1) "Dangerous drug" and "prescriber" have the same	9
meanings as in section 4729.01 of the Revised Code, except that	10
a prescriber does not include a veterinarian licensed under	11
Chapter 4741. of the Revised Code.	12
(2) "Severe adverse effect" means any of the following:	13
(a) Death;	14
(b) Infection requiring hospitalization;	15
(c) Hemorrhaging requiring hospitalization;	16

(d) Organ failure;	17
(e) Sepsis.	18
(B) Before a prescriber may issue for a patient a	19
prescription for a dangerous drug that causes one or more severe	20
adverse effects in greater than five per cent of the drug's	21
users, the prescriber shall do all of the following:	22
(1) Conduct an in-person examination of the patient;	23
(2) Inform the patient that the drug causes one or more	24
severe adverse effects in greater than five per cent of the	25
<pre>drug's users;</pre>	26
(3) Schedule the patient for a follow-up appointment.	27
(C)(1) For purposes of this section, the director of	28
health is responsible for determining if a dangerous drug causes	
one or more severe adverse effects in greater than five per cent	30
of the drug's users. In making such a determination, both of the	31
<pre>following apply:</pre>	32
(a) The director shall consult with the superintendent of	33
insurance and executive directors of the state board of pharmacy	34
and state medical board.	35
(b) The director shall base the determination on the	36
greater of insurance claims, patient reports of severe adverse	37
effects to health care professionals, and any applicable data	38
available from the United States food and drug administration.	39
(2) The director of health shall prepare and update as	40
needed a list containing each dangerous drug that the director	41
determines causes one or more severe adverse effects in greater	42
than five per cent of the drug's users. The director shall make	43
the list, and each of its updates, available to the public on	44

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the internet web site maintained by the department of health.	45
Section 2. This act shall be known as the Patient	46
Protection Act.	47