SENATE MOTION

MR. PRESIDENT:

I move that Senate Bill 262 be amended to read as follows:

Page 1, delete lines 1 through 14, begin a new paragraph and insert:

"SECTION 1. IC 25-26-24-3, AS ADDED BY P.L.51-2019, SECTION 8, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2026]: Sec. 3. ~~(a)~~ As used in this chapter, "dispense" means to deliver a controlled substance to an ultimate user or research subject by or pursuant to the lawful order of a practitioner and includes the prescribing, administering, packaging, labeling, or compounding necessary to prepare the substance for that delivery.

~~(b) The term does not apply to the following:~~

~~(1) A drug administered directly to a patient.~~

~~(2) A drug dispensed by a practitioner, if the quantity dispensed is not more than a seventy-two (72) hour supply of a controlled substance listed in schedule II, III, IV, or V as set forth in IC 35-48-3-9.~~".

Page 2, between lines 10 and 11, begin a new paragraph and insert:

"SECTION 3. IC 25-26-24-17, AS AMENDED BY P.L.17-2021, SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2026]: Sec. 17. (a) **Except as provided in section 17.5 of this chapter,** the board shall provide for an ephedrine, pseudoephedrine, and controlled substance prescription monitoring program that includes the following components:

(1) Each time ephedrine, pseudoephedrine, or a controlled substance designated by the board under IC 35-48-2-5 through IC 35-48-2-10 is dispensed, the dispenser shall transmit to the INSPECT program the following information:

(A) The ephedrine, pseudoephedrine, or controlled substance recipient's name.

(B) The ephedrine, pseudoephedrine, or controlled substance recipient's or the recipient representative's identification number or the identification number or phrase designated by the INSPECT program.

(C) The ephedrine, pseudoephedrine, or controlled substance recipient's date of birth.

(D) The national drug code number of the ephedrine, pseudoephedrine, or controlled substance dispensed.

(E) The date the ephedrine, pseudoephedrine, or controlled substance is dispensed.

(F) The quantity of the ephedrine, pseudoephedrine, or controlled substance dispensed.

(G) The number of days of supply dispensed.

(H) The dispenser's United States Drug Enforcement Agency registration number.

(I) The prescriber's United States Drug Enforcement Agency registration number.

(J) An indication as to whether the prescription was transmitted to the pharmacist orally or in writing.

(K) Other data required by the board.

(2) The information required to be transmitted under this section must be transmitted not more than twenty-four (24) hours after the date on which ephedrine, pseudoephedrine, or a controlled substance is dispensed. However, if the dispenser's pharmacy is closed the day following the dispensing, the information must be transmitted by the end of the next business day. A dispenser who is also authorized to prescribe is required only to report actual dispensations within twenty-four (24) hours of the dispensation.

(3) A dispenser shall transmit the information required under this section by:

(A) uploading to the INSPECT ~~Internet web site;~~ **website;** or

(B) another electronic method that meets specifications prescribed by the board.

(4) The board may require that prescriptions for ephedrine, pseudoephedrine, or controlled substances be written on a one (1) part form that cannot be duplicated. However, the board may not apply such a requirement to prescriptions filled at a pharmacy with a Category II permit (as described in IC 25-26-13-17) and operated by a hospital licensed under IC 16-21, or prescriptions ordered for and dispensed to bona fide enrolled patients in facilities licensed under IC 16-28. The board may not require multiple copy prescription forms for any prescriptions written. The board may not require different prescription forms for any individual drug or group of drugs. Prescription forms required under this subdivision must be approved by the Indiana board of pharmacy created by IC 25-26-13-3.

(5) The costs of the program.

(6) As part of the information to be completed in the data base and, if available, an entry where a dispenser indicates that a patient is participating in a pain management contract with a designated practitioner.

(b) The board shall consider the recommendations of the committee concerning the INSPECT program.

(c) This subsection applies only to a retail pharmacy. A pharmacist, pharmacy technician, or person authorized by a pharmacist to dispense ephedrine, pseudoephedrine, or a controlled substance may not dispense ephedrine, pseudoephedrine, or a controlled substance to a person who is not personally known to the pharmacist, pharmacy technician, or person authorized by a pharmacist to dispense a controlled substance unless the person taking possession of the ephedrine, pseudoephedrine, or controlled substance provides documented proof of the person's identification to the pharmacist, pharmacy technician, or person authorized by a pharmacist to dispense ephedrine, pseudoephedrine, or a controlled substance.

SECTION 4. IC 25-26-24-17.5 IS ADDED TO THE INDIANA CODE AS A **NEW** SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2026]: **Sec. 17.5. A dispenser is not required to transmit information required under section 17 of this chapter to the INSPECT program in the following circumstances:**

(1) A drug is administered directly to a patient.

(2) A drug is dispensed by a veterinary practitioner, if the quantity dispensed is not more than a seventy-two (72) hour supply of a controlled substance listed in schedule II, III, IV, or V as set forth in IC 35-48-3-9.".

Renumber all SECTIONS consecutively.

(Reference is to SB 262 as printed January 16, 2026.)

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Senator CRIDER