SENATE MOTION

MR. PRESIDENT:

I move that Senate Bill 282 be amended to read as follows:

Page 1, delete lines 1 through 17.

Page 2, delete lines 1 through 3.

Page 2, delete lines 7 through 42, begin a new paragraph and insert:

"**Chapter 22.5. Drugs: Compounding**

Sec. 1. As used in this chapter, "compounding pharmacy" means a pharmacy licensed under IC 25-26 that is subject to compounding drug regulation under 21 U.S.C. 353a.

Sec. 2. As used in this chapter, "monograph" means quality standards for prescription drug medicines and dietary supplements that specify the quality expectations for the medicine or dietary supplement, including the identity, strength, purity, and performance of the medicine or dietary supplement.

Sec. 3. A patient has a right to determine, with assistance and guidance of health care providers, individual courses of treatment through the use of medications and treatments obtained from a compounding pharmacy.

Sec. 4. (a) Except as provided in subsection (b), a compounding pharmacy shall have access to active pharmaceutical ingredients that meet federal Food and Drug Administration pharmacopeia monographs for use in compounding if the active pharmaceutical ingredient is:

(1) prepared for use by a manufacturer or repackager that is registered with the federal Food and Drug Administration; and

(2) shipped into Indiana in compliance with state law and accompanied with a certificate of analysis detailing quality specifications.

This subsection includes medications, dietary supplements, and

amino acids that are already in use by compounding pharmacies in Indiana and are provided to patients with a prescribed individual course of treatment by a prescribing health care provider.

(b) Subsection (a) does not apply if the active pharmaceutical ingredient is deemed unsafe for compounding by the federal Food and Drug Administration.

(c) Subject to the enforcement discretion of the federal Food and Drug Administration, a compounding pharmacy may use substances placed on the federal Food and Drug Administration 503A Category 1 bulk drug substance list or active pharmaceutical ingredients.

(d) Subsection (a) does not restrict the use of an active pharmaceutical ingredient that is a component of a drug approved by the federal Food and Drug Administration.

Sec. 5. This chapter may not be deemed to allow any treatment or use of medication that is intended to cause the death of a patient.

Sec. 6. Nothing in this chapter may be construed to preempt, limit, or supersede any Indiana law or administrative rule governing abortion or gender transition treatment.".

Delete pages 3 through 4.

Page 5, delete lines 1 through 38.

Renumber all SECTIONS consecutively.

(Reference is to SB 282 as printed January 23, 2026.)

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Senator GARTEN