ENGROSSED SUBSTITUTE SENATE BILL 5017

State of Washington 57th Legislature 2001 Regular Session

By Senate Committee on Judiciary (originally sponsored by Senators Franklin, Winsley and Regala)

READ FIRST TIME 02/07/01.

AN ACT Relating to precursor drugs; amending RCW 69.43.010, 69.43.020, 69.43.040, and 69.43.090; adding new sections to chapter 69.43 RCW; creating a new section; and prescribing penalties.

4 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF WASHINGTON:

NEW SECTION. Sec. 1. Communities all over the state of Washington 5 experienced an increase in the illegal manufacture of 6 have 7 Illegal methamphetamine labs create a significant methamphetamine. threat to the health and safety of the people of the state. 8 Some of the chemicals and compounds used to make methamphetamine, and the toxic 9 10 wastes the process generates, are hazards to the public health. Increases in crime, violence, and the abuse and neglect of children 11 12 present at laboratory sites are also associated with the increasing 13 number of illeqal laboratory sites. The druqs ephedrine, pseudoephedrine, and phenylpropanolamine, which are used in the illegal 14 15 manufacture of methamphetamine, have been identified as factors in the increase in the number of illegal methamphetamine labs. Therefore, it 16 17 is the intent of the legislature to place restrictions on the sale and possession of those three drugs in order to reduce the proliferation of 18

illegal methamphetamine laboratories and the associated threats to
 public health and safety.

3 Sec. 2. RCW 69.43.010 and 1998 c 245 s 107 are each amended to 4 read as follows: (1) ((Beginning July 1, 1988,)) A report to the state board of 5 pharmacy shall be submitted in accordance with this chapter by a б 7 manufacturer, <u>wholesaler</u>, retailer, or other person who sells, transfers, or otherwise furnishes to any person ((in this state)) any 8 9 of the following substances or their salts or isomers: (a) Anthranilic acid; 10 (b) Barbituric acid; 11 12 (c) Chlorephedrine; 13 (d) Diethyl malonate; 14 (e) D-lysergic acid; 15 (f) Ephedrine; 16 (g) Ergotamine tartrate; (h) Ethylamine; 17 18 (i) Ethyl malonate; 19 (j) Ethylephedrine; (k) Lead acetate; 20 (1) Malonic acid; 21 22 (m) Methylamine; 23 (n) ((Methylformanide)) Methylformamide; 24 (o) Methylephedrine; 25 (p) Methylpseudoephedrine; 26 (q) N-acetylanthranilic acid; 27 (r) Norpseudoephedrine; (s) Phenylacetic acid; 28 (t) Phenylpropanolamine; 29 (u) Piperidine; 30 (v) Pseudoephedrine; and 31 (w) Pyrrolidine. 32 33 (2) The state board of pharmacy shall administer this chapter and 34 may, by rule adopted pursuant to chapter 34.05 RCW, add a substance to

35 or remove a substance from the list in subsection (1) of this section.
36 In determining whether to add or remove a substance, the board shall
37 consider the following:

(a) The likelihood that the substance is useable as a precursor in 1 2 the illegal production of a controlled substance as defined in chapter 3 69.50 RCW;

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(b) The availability of the substance;

5 (c) The relative appropriateness of including the substance in this chapter or in chapter 69.50 RCW; and 6

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(d) The extent and nature of legitimate uses for the substance.

8 (3)(a) ((Beginning on July 1, 1988,)) Any manufacturer, wholesaler, 9 retailer, or other person shall, before selling, transferring, or 10 otherwise furnishing any substance specified in subsection (1) of this section to ((a)) any person ((in this state)), require proper 11 12 identification from the purchaser.

13 (b) For the purposes of this subsection, "proper identification" means((, in the case of a face-to-face purchase,)): 14

(i) A motor vehicle operator's license or other official state-15 16 issued identification of the purchaser containing a photograph of the 17 purchaser, and includes the residential or mailing address of the purchaser, other than a post office box number((τ)); 18

19 (ii) The motor vehicle license number of any motor vehicle owned or 20 operated by the purchaser $((-))_{i}$

(iii) A letter of authorization from any business for which any 21 substance specified in subsection (1) of this section is being 22 23 furnished, which includes the business license number and address of 24 the business((-));

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(iv) A description of how the substance is to be used((τ)); and 26 (v) The signature of the purchaser.

The person selling, transferring, or otherwise furnishing any 27 substance specified in subsection (1) of this section shall affix his 28 or her signature as a witness to the signature and identification of 29 30 the purchaser. ((The state board of pharmacy shall provide by rule for the proper identification of purchasers in other than face-to-face 31 32 purchases.))

(c) A violation of or a failure to comply with this subsection is 33 34 a misdemeanor.

35 (4) ((Beginning on July 1, 1988,)) Any manufacturer, wholesaler, retailer, or other person who sells, transfers, or otherwise furnishes 36 37 the substance specified in subsection (1) of this section to ((a)) any person ((in this state)) shall, not less than twenty-one days before 38 39 delivery of the substance, submit a report of the transaction, which 1 includes the identification information specified in subsection (3) of 2 this section to the state board of pharmacy. However, the state board 3 of pharmacy may authorize the submission of the reports on a monthly 4 basis with respect to repeated, regular transactions between the 5 furnisher and the recipient involving the same substance if the state 6 board of pharmacy determines that either of the following exist:

7 (a) A pattern of regular supply of the substance exists between the
8 manufacturer, wholesaler, retailer, or other person who sells,
9 transfers, or otherwise furnishes such substance and the recipient of
10 the substance; or

11 (b) The recipient has established a record of using the substance 12 for lawful purposes.

(5) Any person specified in subsection (4) of this section who does
not submit a report as required by ((that)) subsection (4) of this
section is guilty of a gross misdemeanor.

16 **Sec. 3.** RCW 69.43.020 and 1988 c 147 s 2 are each amended to read 17 as follows:

18 (1) ((Beginning on July 1, 1988,)) Any manufacturer, wholesaler, 19 retailer, or other person ((subject to any other reporting requirements 20 in this chapter,)) who receives from a source outside of this state any 21 substance specified in RCW 69.43.010(1)((τ)) shall submit a report of 22 such transaction to the state board of pharmacy under rules adopted by 23 the board.

(2) Any person specified in subsection (1) of this section who does
not submit a report as required by subsection (1) of this section is
guilty of a gross misdemeanor.

27 <u>NEW SECTION.</u> **Sec. 4.** A new section is added to chapter 69.43 RCW 28 to read as follows:

(1) Any manufacturer or wholesaler who sells, transfers, or otherwise furnishes any substance specified in RCW 69.43.010(1) to any person in a suspicious transaction shall report the transaction in writing to the state board of pharmacy.

(2) Any person specified in subsection (1) of this section who does
not submit a report as required by subsection (1) of this section is
guilty of a gross misdemeanor.

36 (3) For the purposes of this section, "suspicious transaction"37 means a sale or transfer to which any of the following applies:

(a) The circumstances of the sale or transfer would lead a 1 reasonable person to believe that the substance is likely to be used 2 for the purpose of unlawfully manufacturing a controlled substance 3 4 under chapter 69.50 RCW, based on such factors as the amount involved, the method of payment, the method of delivery, and any past dealings 5 with any participant in the transaction. The state board of pharmacy 6 7 shall adopt by rule criteria for determining whether a transaction is 8 suspicious, taking into consideration the recommendations in appendix 9 A of the report to the United States attorney general by the suspicious 10 orders task force under the federal comprehensive methamphetamine control act of 1996. 11

(b) The transaction involves payment for any substance specified in
RCW 69.43.010(1) in cash or money orders in a total amount of more than
two hundred dollars.

15 <u>NEW SECTION.</u> Sec. 5. A new section is added to chapter 69.43 RCW 16 to read as follows:

(1) Any manufacturer or wholesaler who sells, transfers, or otherwise furnishes any substance specified in RCW 69.43.010(1) to any person shall maintain a record of each such sale or transfer. The records must contain:

21 (a) The name of the substance;

22 (b) The quantity of the substance sold, transferred, or furnished;

23 (c) The date the substance was sold, transferred, or furnished;

(d) The name and address of the person buying or receiving thesubstance; and

26 (e) The method of and amount of payment for the substance.

(2) The records of sales and transfers required by this section
shall be available for inspection by the state board of pharmacy and
its authorized representatives and shall be maintained for two years.
(3) A violation of this section is a gross misdemeanor.

31 <u>NEW SECTION.</u> Sec. 6. A new section is added to chapter 69.43 RCW 32 to read as follows:

A manufacturer, wholesaler, retailer, or other person who sells, transfers, or otherwise furnishes any substance specified in RCW 69.43.010(1) and who is subject to the reporting or recordkeeping requirements of this chapter may satisfy the requirements by submitting to the state board of pharmacy, and its authorized representatives:

1 (1) Computer readable data from which all of the required 2 information may be readily derived; or

3 (2) Copies of reports that are filed under federal law that contain 4 all of the information required by the particular reporting or 5 recordkeeping requirement of this chapter which it is submitted to 6 satisfy.

7 Sec. 7. RCW 69.43.040 and 1989 1st ex.s. c 9 s 441 are each 8 amended to read as follows:

9 (1) The department of health, in accordance with rules developed by 10 the state board of pharmacy shall provide a common reporting form for 11 the substances in RCW 69.43.010 that contains at least the following 12 information:

13 (a) Name of the substance;

14 (b) Quantity of the substance sold, transferred, or furnished;

15 (c) The date the substance was sold, transferred, or furnished;

16 (d) The name and address of the person buying or receiving the 17 substance; and

(e) The name and address of the manufacturer, wholesaler, retailer,or other person selling, transferring, or furnishing the substance.

(2) Monthly reports authorized under ((subsection (1)(e) of this
 section)) <u>RCW 69.43.010(4)</u> may be computer-generated in accordance with
 rules adopted by the department.

23 **Sec. 8.** RCW 69.43.090 and 1989 1st ex.s. c 9 s 443 are each 24 amended to read as follows:

(1) Any manufacturer, wholesaler, retailer, or other person who 25 sells, transfers, or otherwise furnishes any substance specified in RCW 26 27 69.43.010 to ((a)) any person ((in this state)) or who receives from a source outside of the state any substance specified in RCW 69.43.010 28 29 shall obtain a permit for the conduct of that business from the state board of pharmacy. However, a permit shall not be required of any 30 manufacturer, wholesaler, retailer, or other person for the sale, 31 32 transfer, furnishing, or receipt of any drug that contains ephedrine, 33 phenylpropanolamine, or pseudoephedrine, or of any cosmetic that contains a substance specified in RCW 69.43.010(1), if such drug or 34 35 cosmetic is lawfully sold, transferred, or furnished over the counter without a prescription or by a prescription under chapter 69.04 or 36 37 69.41 RCW.

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1 (2) Applications for permits shall be filed with the department in 2 writing and signed by the applicant, and shall set forth the name of 3 the applicant, the business in which the applicant is engaged, the 4 business address of the applicant, and a full description of any 5 substance sold, transferred, or otherwise furnished, or received.

6 (3) The board may grant permits on forms prescribed by it. The 7 permits shall be effective for not more than one year from the date of 8 issuance.

9 (4) Each applicant shall pay at the time of filing an application 10 for a permit a fee determined by the department.

(5) A permit granted under this chapter may be renewed on a date to be determined by the board, and annually thereafter, upon the filing of a renewal application and the payment of a permit renewal fee determined by the department.

(6) Permit fees charged by the department shall not exceed thecosts incurred by the department in administering this chapter.

(7) Selling, transferring, or otherwise furnishing, or receiving
any substance specified in RCW 69.43.010 without a required permit, is
a gross misdemeanor.

20 <u>NEW SECTION.</u> **Sec. 9.** A new section is added to chapter 69.43 RCW 21 to read as follows:

(1) It is unlawful for a pharmacy licensed by, or shopkeeper or
itinerant vendor registered with, the department of health under
chapter 18.64 RCW, or an employee thereof, knowingly to sell, transfer,
or to otherwise furnish, in a single transaction:

(a) More than three packages of one or more products that he or she
knows to contain ephedrine, pseudoephedrine, or phenylpropanolamine,
their salts, isomers, or salts of isomers; or

(b) A single package of any product that he or she knows to contain more than three grams of ephedrine, pseudoephedrine, or phenylpropanolamine, their salts, isomers, or salts of isomers, or a combination of any of these substances.

(2) It is unlawful for a person who is not a manufacturer, wholesaler, pharmacy, practitioner, shopkeeper, or itinerant vendor licensed by or registered with the department of health under chapter 18.64 RCW to purchase or acquire, in any twenty-four hour period, more than the quantities of the substances specified in subsection (1) of this section.

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(3) A violation of this section is a gross misdemeanor.

2 <u>NEW SECTION.</u> Sec. 10. A new section is added to chapter 69.43 RCW
3 to read as follows:

4 (1) Any person who possesses more than fifteen grams of ephedrine,
5 pseudoephedrine, or phenylpropanolamine, their salts, isomers, or salts
6 of isomers, or a combination of any of those substances, is guilty of
7 a gross misdemeanor.

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(2) This section does not apply to any of the following:

9 (a) A pharmacist or other authorized person who sells or furnishes 10 ephedrine, pseudoephedrine, or phenylpropanolamine, their salts, 11 isomers, or salts of isomers upon the prescription of a practitioner, 12 as defined in RCW 69.41.010;

(b) A practitioner who administers or furnishes ephedrine,
pseudoephedrine, or phenylpropanolamine, their salts, isomers, or salts
of isomers to his or her patients;

16 (c) A pharmacy, manufacturer, or wholesaler licensed by, or 17 shopkeeper or itinerant vendor registered with, the department of 18 health under chapter 18.64 RCW;

(d) A person in the course of his or her business of selling,
transporting, or storing ephedrine, pseudoephedrine, or
phenylpropanolamine, their salts, isomers, or salts of isomers, for a
person described in (a), (b), or (c) of this subsection; or

(e) A person in possession of more than fifteen grams of ephedrine, pseudoephedrine, or phenylpropanolamine, their salts, isomers, or salts of isomers in their home or residence under circumstances consistent with typical medicinal or household use as indicated by, but not limited to, storage location and possession of products in a variety of strengths, brands, types, purposes, and expiration dates.

29 <u>NEW SECTION.</u> Sec. 11. A new section is added to chapter 69.43 RCW 30 to read as follows:

31 Sections 9 and 10 of this act do not apply to:

(1) Pediatric products primarily intended for administration to 32 33 children under twelve years of age, according to label instructions, either: (a) In solid dosage form whose individual dosage units do not 34 35 exceed fifteen milligrams of ephedrine, pseudoephedrine, or phenylpropanolamine; or (b) in liquid form whose recommended dosage, 36 according to label instructions, does not exceed fifteen milligrams of 37

ephedrine, pseudoephedrine, or phenylpropanolamine per five milliliters
 of liquid product;

3 (2) Pediatric liquid products primarily intended for administration 4 to children under two years of age for which the recommended dosage 5 does not exceed two milliliters and the total package content does not 6 exceed one fluid ounce; or

7 (3) Products that the state board of pharmacy, upon application of 8 a manufacturer, exempts by rule from sections 9 and 10 of this act 9 because the product has been formulated in such a way as to effectively 10 prevent the conversion of the active ingredient into methamphetamine, 11 or its salts or precursors.

12 <u>NEW SECTION.</u> Sec. 12. A new section is added to chapter 69.43 RCW 13 to read as follows:

(1) In addition to the other penalties provided for in this chapter or in chapter 18.64 RCW, the state board of pharmacy may impose a civil penalty, not to exceed ten thousand dollars for each violation, on any licensee or registrant who has failed to comply with this chapter or the rules adopted under this chapter. In the case of a continuing violation, every day the violation continues shall be considered a separate violation.

21 (2) The state board of pharmacy may waive the suspension or 22 revocation of a license or registration issued under chapter 18.64 RCW, 23 or waive any civil penalty under this chapter, if the licensee or 24 registrant establishes that he or she acted in good faith to prevent 25 violations of this chapter, and the violation occurred despite the licensee's or registrant's exercise of due diligence. In making such 26 a determination, the state board of pharmacy may consider evidence that 27 28 an employer trained employees on how to sell, transfer, or otherwise 29 furnish substances specified in RCW 69.43.010(1) in accordance with applicable laws. 30

31 <u>NEW SECTION.</u> Sec. 13. A new section is added to chapter 69.43 RCW 32 to read as follows:

This chapter is applicable and uniform throughout this state and in all counties, cities, code cities, and towns therein. A county, city, code city, or town may not adopt or enforce any ordinance, pertaining to this chapter, which prohibits conduct that is not prohibited under this chapter, or defining violations or penalties different from those

1 provided under this chapter. However, this section does not preclude 2 a county, city, code city, or town from revoking, canceling, 3 suspending, or otherwise limiting a business or professional license it 4 has issued for conduct that violates any provision of this chapter.

5 <u>NEW SECTION.</u> **Sec. 14.** A new section is added to chapter 69.43 RCW 6 to read as follows:

7 (1) To prevent violations of section 9 of this act, every licensee 8 and registrant under chapter 18.64 RCW, who sells at retail any 9 products containing ephedrine, pseudoephedrine, or phenylpropanolamine, 10 or their salts, isomers, or salts of isomers, shall do either or may do 11 both of the following:

(a) Program scanners, cash registers, or other electronic devices used to record sales in a manner that will alert persons handling transactions to potential violations of section 9(1) of this act and/or prevent such violations; or

(b) Place one or more signs on the premises to notify customers of the prohibitions of section 9 of this act. Any such sign may, but is not required to, conform to the language and format prepared by the department of health under subsection (2) of this section.

(2) The department of health shall prepare language and format for 20 a sign summarizing the prohibitions in sections 9 and 10 of this act 21 and make the language and format available to licensees and registrants 22 23 under chapter 18.64 RCW, for voluntary use in their places of business 24 to inform customers and employees of the prohibitions. Nothing in this 25 section requires the department of health to provide licensees or registrants with copies of signs, or any licensee or registrant to use 26 the specific language or format prepared by the department under this 27 28 subsection.

29 <u>NEW SECTION.</u> Sec. 15. If any provision of this act or its 30 application to any person or circumstance is held invalid, the 31 remainder of the act or the application of the provision to other 32 persons or circumstances is not affected.