

EHB 1538 - S COMM AMD
By Committee on Health Care

1 Strike everything after the enacting clause and insert the
2 following:

3 "NEW SECTION. **Sec. 1.** A new section is added to chapter 70.05 RCW
4 to read as follows:

5 (1) A registered nurse who is licensed under chapter 18.79 RCW, and
6 who is an employee of a local health department or district or a clinic
7 or facility under contract with a local health department or district,
8 may dispense a drug or device for purposes of prevention or treatment
9 of a communicable disease or family planning.

10 (2) Such dispensing must be pursuant to the order of a person
11 authorized to prescribe a drug or device in the state of Washington.

12 (3) The local health officer must establish policies and procedures
13 that include the following:

14 (a) Procedures for drug dispensing, documentation storage,
15 security, and accountability;

16 (b) Maintenance of all drug records required by federal and state
17 law.

18 (4) A drug may only be dispensed by a practitioner authorized by
19 law to prescribe the drug or by a registered nurse licensed under
20 chapter 18.79 RCW, and must be dispensed in a container complying with
21 the federal poison prevention packaging act unless the patient requests
22 a noncomplying container. A registered nurse may only dispense a drug
23 or device for purposes of prevention or treatment of a communicable
24 disease or family planning. Each drug that is dispensed must be
25 labeled with the following:

26 (a) Name of patient;

27 (b) Name of prescriber;

28 (c) Name, address, and phone number of the clinic;

29 (d) Date of dispensing;

1 (e) Name and strength of the drug. If the drug does not have a
2 brand name, then the generic name of the drug and the drug manufacturer
3 must be stated;

4 (f) Directions for use;

5 (g) Initials of the person dispensing;

6 (h) Cautionary statements, if any, as required by law;

7 (i) Manufacturer's expiration date, or an earlier date if
8 preferable, after which the patient should not use the drug.

9 A drug information fact sheet must accompany each drug dispensed from
10 a local health department or district or clinic or facility under
11 contract with a local health department or district.

12 (5) A drug repackaged for dispensing must be in a container meeting
13 United States pharmacopeia standards and labeled to identify at a
14 minimum:

15 (a) Brand name, or generic name and manufacturer;

16 (b) Strength;

17 (c) Lot number;

18 (d) Manufacturer's expiration date or an earlier date if
19 preferable. An internal control number that references manufacturer
20 and lot number may be used.

21 (6) In the absence of a dispensing practitioner or a registered
22 nurse, drugs must be kept in a locked drug cabinet or drug room which
23 is sufficiently secure to deny access to unauthorized persons. Only
24 dispensing practitioners and registered nurses may have access to the
25 drug cabinet or drug room. In their absence, the drug cabinet or drug
26 room must remain locked. All drugs must be stored in areas that will
27 assure proper sanitation, temperature, light, ventilation and moisture
28 control as recommended by the manufacturer. Drugs that are outdated,
29 damaged, deteriorated, misbranded, or adulterated must be quarantined
30 and physically separated from other drugs until they are destroyed or
31 returned to their supplier.

32 (7) A dispensing record must be maintained separately from the
33 patient chart and kept for a minimum of three years. The record must
34 show, at a minimum, the following:

35 (a) Name of patient;

36 (b) Brand name of drug, or generic name and name of manufacturer or
37 distributor;

38 (c) Amount dispensed;

1 (d) Date;

2 (e) Initials of person dispensing the prescription.

3 All records of receipt and disposal of drugs must be kept for a minimum
4 of three years. All records required by these rules or by federal and
5 state law must be readily retrievable for inspection by the board of
6 pharmacy.

7 (8) Notwithstanding any other requirements in this section, when a
8 drug is dispensed in the practice of the expedited partner therapy
9 treatment protocol, the name of the patient may be omitted from the
10 label, the patient's name may be omitted from the records and a drug
11 may be dispensed to the patient to be given to the patient's partner
12 even if the partner has not been examined by a licensed health care
13 provider acting within his or her scope of practice.

14 (9) The pharmacy quality assurance commission may adopt rules
15 necessary to implement this section."

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16 On page 1, line 2 of the title, after "medications;" strike the
17 remainder of the title and insert "and adding a new section to chapter
18 70.05 RCW."

EFFECT: Corrects a reference to the pharmacy quality assurance
commission.

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