
HOUSE BILL 2893

State of Washington 52nd Legislature 1992 Regular Session

By Representatives Prentice and Wood

Read first time 01/31/92. Referred to Committee on Health Care.

1 AN ACT Relating to dispensing drug outlets; amending RCW 18.64.011,
2 18.64.246, and 18.64.255; reenacting and amending RCW 18.64.165 and
3 18.64.245; adding a new section to chapter 18.64 RCW; and prescribing
4 penalties.

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

6 **Sec. 1.** RCW 18.64.011 and 1989 1st ex.s. c 9 s 412 are each
7 amended to read as follows:

8 Unless the context clearly requires otherwise, definitions of terms
9 shall be as indicated when used in this chapter.

10 (1) "Person" means an individual, corporation, government,
11 governmental subdivision or agency, business trust, estate, trust,
12 partnership or association, or any other legal entity.

13 (2) "Board" means the Washington state board of pharmacy.

14 (3) "Drugs" means:

1 (a) Articles recognized in the official United States pharmacopoeia
2 or the official homeopathic pharmacopoeia of the United States;

3 (b) Substances intended for use in the diagnosis, cure, mitigation,
4 treatment, or prevention of disease in man or other animals;

5 (c) Substances (other than food) intended to affect the structure
6 or any function of the body of man or other animals; or

7 (d) Substances intended for use as a component of any substances
8 specified in (a), (b), or (c) of this subsection, but not including
9 devices or their component parts or accessories.

10 (4) "Device" means instruments, apparatus, and contrivances,
11 including their components, parts, and accessories, intended (a) for
12 use in the diagnosis, cure, mitigation, treatment, or prevention of
13 disease in man or other animals, or (b) to affect the structure or any
14 function of the body of man or other animals.

15 (5) "Nonlegend" or "nonprescription" drugs means any drugs which
16 may be lawfully sold without a prescription.

17 (6) "Legend drugs" means any drugs which are required by any
18 applicable federal or state law or regulation to be dispensed on
19 prescription only or are restricted to use by practitioners only.

20 (7) "Controlled substance" means a drug or substance, or an
21 immediate precursor of such drug or substance, so designated under or
22 pursuant to the provisions of chapter 69.50 RCW.

23 (8) "Prescription" means an order for drugs or devices issued by a
24 practitioner duly authorized by law or rule in the state of Washington
25 to prescribe drugs or devices in the course of his or her professional
26 practice for a legitimate medical purpose.

27 (9) "Practitioner" means a physician, dentist, veterinarian, nurse,
28 or other person duly authorized by law or rule in the state of
29 Washington to prescribe drugs.

1 (10) "Pharmacist" means a person duly licensed by the Washington
2 state board of pharmacy to engage in the practice of pharmacy.

3 (11) "Practice of pharmacy" includes the practice of and
4 responsibility for: Interpreting prescription orders; the compounding,
5 dispensing, labeling, administering, and distributing of drugs and
6 devices; the monitoring of drug therapy and use; the initiating or
7 modifying of drug therapy in accordance with written guidelines or
8 protocols previously established and approved for his or her practice
9 by a practitioner authorized to prescribe drugs; the participating in
10 drug utilization reviews and drug product selection; the proper and
11 safe storing and distributing of drugs and devices and maintenance of
12 proper records thereof; the providing of information on legend drugs
13 which may include, but is not limited to, the advising of therapeutic
14 values, hazards, and the uses of drugs and devices.

15 (12) "Pharmacy" means every place properly licensed by the board of
16 pharmacy where the practice of pharmacy is conducted.

17 (13) The words "drug" and "devices" shall not include surgical or
18 dental instruments or laboratory materials, gas and oxygen, therapy
19 equipment, x-ray apparatus or therapeutic equipment, their component
20 parts or accessories, or equipment, instruments, apparatus, or
21 contrivances used to render such articles effective in medical,
22 surgical, or dental treatment, or for use or consumption in or for
23 mechanical, industrial, manufacturing, or scientific applications or
24 purposes, nor shall the word "drug" include any article or mixture
25 covered by the Washington pesticide control act (chapter 15.58 RCW), as
26 enacted or hereafter amended, nor medicated feed intended for and used
27 exclusively as a feed for animals other than man.

28 (14) The word "poison" shall not include any article or mixture
29 covered by the Washington pesticide control act (chapter 15.58 RCW), as
30 enacted or hereafter amended.

1 (15) "Deliver" or "delivery" means the actual, constructive, or
2 attempted transfer from one person to another of a drug or device,
3 whether or not there is an agency relationship.

4 (16) "Dispense" means the interpretation of a prescription or order
5 for a drug, biological, or device and, pursuant to that prescription or
6 order, the proper selection, measuring, compounding, labeling, or
7 packaging necessary to prepare that prescription or order for delivery.

8 (17) "Dispensing drug outlet" means a drug outlet where drugs are
9 dispensed by a health care practitioner, other than a pharmacist, who
10 is authorized to dispense to his or her patients only.

11 (18) "Distribute" means the delivery of a drug or device other than
12 by administering or dispensing.

13 (~~(18)~~) (19) "Compounding" shall be the act of combining two or
14 more ingredients in the preparation of a prescription.

15 (~~(19)~~) (20) "Wholesaler" shall mean a corporation, individual, or
16 other entity which buys drugs or devices for resale and distribution to
17 corporations, individuals, or entities other than consumers.

18 (~~(20)~~) (21) "Manufacture" means the production, preparation,
19 propagation, compounding, or processing of a drug or other substance or
20 device or the packaging or repackaging of such substance or device, or
21 the labeling or relabeling of the commercial container of such
22 substance or device, but does not include the activities of a
23 practitioner who, as an incident to his or her administration or
24 dispensing such substance or device in the course of his or her
25 professional practice, prepares, compounds, packages, or labels such
26 substance or device.

27 (~~(21)~~) (22) "Manufacturer" shall mean a person, corporation, or
28 other entity engaged in the manufacture of drugs or devices.

29 (~~(22)~~) (23) "Labeling" shall mean the process of preparing and
30 affixing a label to any drug or device container. The label must

1 include all information required by current federal and state law and
2 pharmacy rules.

3 ~~((+23+))~~ (24) "Administer" means the direct application of a drug
4 or device, whether by injection, inhalation, ingestion, or any other
5 means, to the body of a patient or research subject.

6 ~~((+24+))~~ (25) "Master license system" means the mechanism
7 established by chapter 19.02 RCW by which master licenses, endorsed for
8 individual state-issued licenses, are issued and renewed utilizing a
9 master application and a master license expiration date common to each
10 renewable license endorsement.

11 ~~((+25+))~~ (26) "Department" means the department of health.

12 ~~((+26+))~~ (27) "Secretary" means the secretary of health or the
13 secretary's designee.

14 **Sec. 2.** RCW 18.64.165 and 1989 1st ex.s. c 9 s 404 and 1989 c 352
15 s 4 are each reenacted and amended to read as follows:

16 The board shall have the power to refuse, suspend, or revoke the
17 license of any manufacturer, wholesaler, pharmacy or dispensing drug
18 outlet, shopkeeper, itinerant vendor, peddler, poison distributor, or
19 precursor chemical distributor upon proof that:

20 (1) The license was procured through fraud, misrepresentation, or
21 deceit;

22 (2) The licensee has violated or has permitted any employee to
23 violate any of the laws of this state or the United States relating to
24 drugs, controlled substances, cosmetics, or nonprescription drugs, or
25 has violated any of the rules ~~((and regulations))~~ of the board of
26 pharmacy or has been convicted of a felony.

27 **Sec. 3.** RCW 18.64.245 and 1989 1st ex.s. c 9 s 402 and 1989 c 352
28 s 2 are each reenacted and amended to read as follows:

1 Every proprietor or manager of a pharmacy or dispensing drug outlet
2 shall keep readily available a suitable record of prescriptions which
3 shall preserve for a period of not less than two years the record of
4 every prescription dispensed at such pharmacy or dispensing drug outlet
5 which shall be numbered, dated, and filed, and shall produce the same
6 in court or before any grand jury whenever lawfully required to do so.
7 The record shall be maintained either separately from all other records
8 of the pharmacy or dispensing drug outlet or in such form that the
9 information required is readily retrievable from ordinary business
10 records of the pharmacy or dispensing drug outlet. All record-keeping
11 requirements for controlled substances must be complied with. Such
12 record of prescriptions shall be for confidential use in the pharmacy
13 or dispensing drug outlet, only. The record of prescriptions shall be
14 open for inspection by the board of pharmacy or any officer of the law,
15 who is authorized to enforce chapter 18.64, 69.41, or 69.50 RCW.

16 **Sec. 4.** RCW 18.64.246 and 1984 c 153 s 13 are each amended to read
17 as follows:

18 To every box, bottle, jar, tube or other container of a
19 prescription which is dispensed there shall be fixed a label bearing
20 the name and address of the pharmacy or dispensing drug outlet wherein
21 the prescription is compounded, the corresponding serial number of the
22 prescription, the name of the prescriber, his directions, the name of
23 the medicine and the strength per unit dose, name of patient, date, the
24 expiration date, and initials of the licensed pharmacist who has
25 compounded the prescription, and the security of the cover or cap on
26 every bottle or jar shall meet safety standards promulgated by the
27 state board of pharmacy: PROVIDED, That at the physician's request,
28 the name and dosage of the drug need not be shown. If the prescription
29 is for a combination drug product, the generic names of the drugs

1 combined or the trade name used by the manufacturer or distributor for
2 the product shall be noted on the label. This section shall not apply
3 to the dispensing of medicines to in-patients in hospitals.

4 **Sec. 5.** RCW 18.64.255 and 1984 c 153 s 14 are each amended to read
5 as follows:

6 Nothing in this chapter shall operate in any manner:

7 (1) To restrict the scope of authorized practice of any
8 practitioner other than a pharmacist, duly licensed as such under the
9 laws of this state. However, any health care practitioner authorized
10 to dispense drugs, and operating a dispensing drug outlet, shall comply
11 with all the state and federal laws and rules relating to the
12 dispensing of drugs and the practice of pharmacy; or

13 (2) In the absence of the pharmacist from the hospital pharmacy, to
14 prohibit a registered nurse designated by the hospital and the
15 responsible pharmacist from obtaining from the hospital pharmacy such
16 drugs as are needed in an emergency: PROVIDED, That proper record is
17 kept of such emergency, including the date, time, name of prescriber,
18 the name of the nurse obtaining the drugs, and a list of what drugs and
19 quantities of same were obtained; or

20 (3) To prevent shopkeepers, itinerant vendors, peddlers, or
21 salesmen from dealing in and selling nonprescription drugs, if such
22 drugs are sold in the original packages of the manufacturer, or in
23 packages put up by a licensed pharmacist in the manner provided by the
24 state board of pharmacy, if such shopkeeper, itinerant vendor,
25 salesman, or peddler shall have obtained a registration.

26 NEW SECTION. **Sec. 6.** A new section is added to chapter 18.64 RCW
27 to read as follows:

1 (1) The owner of each dispensing drug outlet shall pay an original
2 license fee to be determined by the secretary, and annually thereafter,
3 on or before a date to be determined by the secretary, a fee to be
4 determined by the secretary, for which he or she shall receive a
5 license of location, which shall entitle the owner to operate the
6 dispensing drug outlet at the location specified, or a temporary
7 location as the secretary may approve, for the period ending on a date
8 to be determined by the secretary, and each owner shall at the time of
9 filing proof of payment of the fee as provided in RCW 18.64.045, file
10 with the department on a blank provided by the department. A
11 declaration of ownership and location filed with the department under
12 this section shall be deemed presumptive evidence of ownership of the
13 dispensing drug outlet.

14 (2) It shall be the duty of the owner to immediately notify the
15 department of any change of location or ownership and to keep the
16 license of location or the renewal of the license properly exhibited in
17 the dispensing drug outlet.

18 (3) Failure to comply with this section is a misdemeanor, and each
19 day that the failure continues is a separate offense.

20 (4) In the event the license fee remains unpaid for sixty days from
21 the date due, no renewal or new license may be issued except upon
22 payment of the license renewal fee and a penalty equal to the original
23 license fee.