H-1128.1			

HOUSE BILL 1424

55th Legislature

1997 Regular Session

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By Representatives Skinner and Murray

State of Washington

Read first time 01/27/97. Referred to Committee on Health Care.

- 1 AN ACT Relating to kidney dialysis centers; and amending RCW
- 2 18.64.011.
- 3 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF WASHINGTON:
- 4 Sec. 1. RCW 18.64.011 and 1995 c 319 s 2 are each amended to read 5 as follows:
- Unless the context clearly requires otherwise, definitions of terms shall be as indicated when used in this chapter.
- 8 (1) "Person" means an individual, corporation, government,
- 9 governmental subdivision or agency, business trust, estate, trust,
- 10 partnership or association, or any other legal entity.
- 11 (2) "Board" means the Washington state board of pharmacy.
- 12 (3) "Drugs" means:
- 13 (a) Articles recognized in the official United States pharmacopoeia
- 14 or the official homeopathic pharmacopoeia of the United States;
- 15 (b) Substances intended for use in the diagnosis, cure, mitigation,
- 16 treatment, or prevention of disease in man or other animals;
- 17 (c) Substances (other than food) intended to affect the structure
- 18 or any function of the body of man or other animals; or

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- 1 (d) Substances intended for use as a component of any substances 2 specified in (a), (b), or (c) of this subsection, but not including 3 devices or their component parts or accessories.
- 4 (4) "Device" means instruments, apparatus, and contrivances, 5 including their components, parts, and accessories, intended (a) for 6 use in the diagnosis, cure, mitigation, treatment, or prevention of 7 disease in man or other animals, or (b) to affect the structure or any 8 function of the body of man or other animals.
- 9 (5) "Nonlegend" or "nonprescription" drugs means any drugs which 10 may be lawfully sold without a prescription.
- 11 (6) "Legend drugs" means any drugs which are required by any 12 applicable federal or state law or regulation to be dispensed on 13 prescription only or are restricted to use by practitioners only.
- 14 (7) "Controlled substance" means a drug or substance, or an 15 immediate precursor of such drug or substance, so designated under or 16 pursuant to the provisions of chapter 69.50 RCW.
- 17 (8) "Prescription" means an order for drugs or devices issued by a 18 practitioner duly authorized by law or rule in the state of Washington 19 to prescribe drugs or devices in the course of his or her professional 20 practice for a legitimate medical purpose.
- (9) "Practitioner" means a physician, dentist, veterinarian, nurse, or other person duly authorized by law or rule in the state of Washington to prescribe drugs.
- 24 (10) "Pharmacist" means a person duly licensed by the Washington 25 state board of pharmacy to engage in the practice of pharmacy.
 - (11) "Practice of pharmacy" includes the practice of and responsibility for: Interpreting prescription orders; the compounding, dispensing, labeling, administering, and distributing of drugs and devices; the monitoring of drug therapy and use; the initiating or modifying of drug therapy in accordance with written guidelines or protocols previously established and approved for his or her practice by a practitioner authorized to prescribe drugs; the participating in drug utilization reviews and drug product selection; the proper and safe storing and distributing of drugs and devices and maintenance of proper records thereof; the providing of information on legend drugs which may include, but is not limited to, the advising of therapeutic values, hazards, and the uses of drugs and devices.
- 38 (12) "Pharmacy" means every place properly licensed by the board of 39 pharmacy where the practice of pharmacy is conducted.

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- (13) The words "drug" and "devices" shall not include surgical or 1 dental instruments or laboratory materials, gas and oxygen, therapy 2 3 equipment, X-ray apparatus or therapeutic equipment, their component 4 parts or accessories, or equipment, instruments, apparatus, or contrivances used to render such articles effective in medical, 5 surgical, or dental treatment, or for use or consumption in or for 6 7 mechanical, industrial, manufacturing, or scientific applications or 8 purposes, nor shall the word "drug" include any article or mixture 9 covered by the Washington pesticide control act (chapter 15.58 RCW), as 10 enacted or hereafter amended, nor medicated feed intended for and used exclusively as a feed for animals other than man. 11
- 12 (14) The word "poison" shall not include any article or mixture 13 covered by the Washington pesticide control act (chapter 15.58 RCW), as 14 enacted or hereafter amended.
- 15 (15) "Deliver" or "delivery" means the actual, constructive, or 16 attempted transfer from one person to another of a drug or device, 17 whether or not there is an agency relationship.
- (16) "Dispense" means the interpretation of a prescription or order for a drug, biological, or device and, pursuant to that prescription or order, the proper selection, measuring, compounding, labeling, or packaging necessary to prepare that prescription or order for delivery.
- 22 (17) "Distribute" means the delivery of a drug or device other than 23 by administering or dispensing.
- 24 (18) "Compounding" shall be the act of combining two or more 25 ingredients in the preparation of a prescription.
- (19) "Wholesaler" shall mean a corporation, individual, or other entity which buys drugs or devices for resale and distribution to corporations, individuals, or entities other than consumers.
- 29 (20) "Manufacture" means the production, preparation, propagation, 30 compounding, or processing of a drug or other substance or device or 31 the packaging or repackaging of such substance or device, or the labeling or relabeling of the commercial container of such substance or 32 device, but does not include the activities of a practitioner who, as 33 34 an incident to his or her administration or dispensing such substance 35 or device in the course of his or her professional practice, prepares, compounds, packages, or labels such substance or device. 36
- 37 (21) "Manufacturer" shall mean a person, corporation, or other 38 entity engaged in the manufacture of drugs or devices.

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- 1 (22) "Labeling" shall mean the process of preparing and affixing a 2 label to any drug or device container. The label must include all 3 information required by current federal and state law and pharmacy 4 rules.
- 5 (23) "Administer" means the direct application of a drug or device, 6 whether by injection, inhalation, ingestion, or any other means, to the 7 body of a patient or research subject.
- 8 (24) "Master license system" means the mechanism established by 9 chapter 19.02 RCW by which master licenses, endorsed for individual 10 state-issued licenses, are issued and renewed utilizing a master 11 application and a master license expiration date common to each 12 renewable license endorsement.
- 13 (25) "Department" means the department of health.
- 14 (26) "Secretary" means the secretary of health or the secretary's designee.
- 16 (27) "Health care entity" means an organization that provides 17 health care services in a setting that is not otherwise licensed by the 18 state. Health care entity includes a free-standing outpatient surgery 19 center((-,)) or a free-standing cardiac care center((-, or a kidney 20 dialysis center)). It does not include an individual practitioner's 21 office or a multipractitioner clinic.

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