

State of Arizona
Senate
Fifty-second Legislature
Second Regular Session
2016

CHAPTER 274
SENATE BILL 1327

AN ACT

AMENDING SECTION 32-1901, ARIZONA REVISED STATUTES; AMENDING TITLE 36,
CHAPTER 4, ARTICLE 1, ARIZONA REVISED STATUTES, BY ADDING SECTION 36-416;
RELATING TO HOSPITALS.

(TEXT OF BILL BEGINS ON NEXT PAGE)

1 Be it enacted by the Legislature of the State of Arizona:

2 Section 1. Section 32-1901, Arizona Revised Statutes, is amended to
3 read:

4 32-1901. Definitions

5 In this chapter, unless the context otherwise requires:

6 1. "Administer" means the direct application of a controlled
7 substance, prescription-only drug, dangerous drug or narcotic drug, whether
8 by injection, inhalation, ingestion or any other means, to the body of a
9 patient or research subject by a practitioner or by the practitioner's
10 authorized agent or the patient or research subject at the direction of the
11 practitioner.

12 2. "Advertisement" means all representations disseminated in any
13 manner or by any means, other than by labeling, for the purpose of inducing,
14 or that are likely to induce, directly or indirectly, the purchase of drugs,
15 devices, poisons or hazardous substances.

16 3. "Advisory letter" means a nondisciplinary letter to notify a
17 licensee or permittee that either:

18 (a) While there is insufficient evidence to support disciplinary
19 action, the board believes that continuation of the activities that led to
20 the investigation may result in further board action against the licensee or
21 permittee.

22 (b) The violation is a minor or technical violation that is not of
23 sufficient merit to warrant disciplinary action.

24 (c) While the licensee or permittee has demonstrated substantial
25 compliance through rehabilitation, remediation or reeducation that has
26 mitigated the need for disciplinary action, the board believes that
27 repetition of the activities that led to the investigation may result in
28 further board action against the licensee or permittee.

29 4. "Antiseptic", if a drug is represented as such on its label, means
30 a representation that it is a germicide, except in the case of a drug
31 purporting to be, or represented as, an antiseptic for inhibitory use as a
32 wet dressing, ointment or dusting powder or other use that involves prolonged
33 contact with the body.

34 5. "Authorized officers of the law" means legally empowered peace
35 officers, compliance officers of the ~~state~~ board of pharmacy and agents of
36 the division of narcotics enforcement and criminal intelligence of the
37 department of public safety.

38 6. "Board" or "board of pharmacy" means the Arizona state board of
39 pharmacy.

40 7. "Color additive" means a material that either:

41 (a) Is any dye, pigment or other substance made by a process of
42 synthesis or similar artifice, or extracted, isolated or otherwise derived,
43 with or without intermediate or final change of identity, from any vegetable,
44 animal, mineral or other source.

45 (b) If added or applied to a drug, or to the human body or any part of
46 the human body, is capable of imparting color, except that color additive

1 does not include any material that has been or may be exempted under the
2 federal act. Color includes black, white and intermediate grays.

3 8. "Compounding" means the preparation, mixing, assembling, packaging
4 or labeling of a drug by a pharmacist or an intern or pharmacy technician
5 under the pharmacist's supervision, for the purpose of dispensing to a
6 patient based on a valid prescription order. Compounding includes the
7 preparation of drugs in anticipation of prescription orders prepared on
8 routine, regularly observed prescribing patterns and the preparation of drugs
9 as an incident to research, teaching or chemical analysis or for
10 administration by a medical practitioner to the medical practitioner's
11 patient and not for sale or dispensing. Compounding does not include the
12 preparation of commercially available products from bulk compounds or the
13 preparation of drugs for sale to pharmacies, practitioners or entities for
14 the purpose of dispensing or distribution.

15 9. "Compressed medical gas distributor" means a person who holds a
16 current permit issued by the board to distribute compressed medical gases
17 pursuant to a compressed medical gas order to compressed medical gas
18 suppliers and other entities that are registered, licensed or permitted to
19 use, administer or distribute compressed medical gases.

20 10. "Compressed medical gas order" means an order for compressed
21 medical gases that is issued by a medical practitioner.

22 11. "Compressed medical gas supplier" means a person who holds a
23 current permit issued by the board to supply compressed medical gases
24 pursuant to a compressed medical gas order and only to the consumer or the
25 patient.

26 12. "Compressed medical gases" means gases and liquid oxygen that a
27 compressed medical gas distributor or manufacturer has labeled in compliance
28 with federal law.

29 13. "Controlled substance" means a drug, substance or immediate
30 precursor THAT IS identified, defined or listed in title 36, chapter 27,
31 article 2.

32 14. "Corrosive" means any substance that when it comes in contact with
33 living tissue will cause destruction of tissue by chemical action.

34 15. "Counterfeit drug" means a drug that, or the container or labeling
35 of which, without authorization, bears the trademark, trade name or other
36 identifying mark, imprint, number or device, or any likeness of these, of a
37 manufacturer, distributor or dispenser other than the person who in fact
38 manufactured, distributed or dispensed that drug.

39 16. "Dangerous drug" has the same meaning prescribed in section
40 13-3401.

41 17. "Decree of censure" means an official action that is taken by the
42 board and that may include a requirement for restitution of fees to a patient
43 or consumer.

44 18. "Deliver" or "delivery" means the actual, constructive or attempted
45 transfer from one person to another whether or not there is an agency
46 relationship.

1 19. "Deputy director" means a pharmacist who is employed by the board
2 and selected by the executive director to perform duties as prescribed by the
3 executive director.

4 20. "Device", except as used in paragraph 15 of this section, section
5 32-1965, paragraph 4 and section 32-1967, subsection A, paragraph 15 and
6 subsection C, means instruments, apparatus and contrivances, including their
7 components, parts and accessories, including all such items under the federal
8 act, intended either:

9 (a) For use in the diagnosis, cure, mitigation, treatment or
10 prevention of disease in the human body or other animals.

11 (b) To affect the structure or any function of the human body or other
12 animals.

13 21. "Direct supervision of a pharmacist" means the pharmacist is
14 present. If relating to the sale of certain items, direct supervision of a
15 pharmacist means that a pharmacist determines the legitimacy or advisability
16 of a proposed purchase of those items.

17 22. "Director" means the director of the division of narcotics
18 enforcement and criminal investigation of the department of public safety.

19 23. "Dispense" means to deliver to an ultimate user or research subject
20 by or pursuant to the lawful order of a practitioner, including the
21 prescribing, administering, packaging, labeling or compounding necessary to
22 prepare for that delivery.

23 24. "Dispenser" means a practitioner who dispenses.

24 25. "Distribute" means to deliver, other than by administering or
25 dispensing.

26 26. "Distributor" means a person who distributes.

27 27. "Drug" means:

28 (a) Articles recognized, or for which standards or specifications are
29 prescribed, in the official compendium.

30 (b) Articles intended for use in the diagnosis, cure, mitigation,
31 treatment or prevention of disease in the human body or other animals.

32 (c) Articles other than food intended to affect the structure or any
33 function of the human body or other animals.

34 (d) Articles intended for use as a component of any articles specified
35 in subdivision (a), (b) or (c) of this paragraph but does not include devices
36 or their components, parts or accessories.

37 28. "Drug enforcement administration" means the drug enforcement
38 administration of the United States department of justice or its successor
39 agency.

40 29. "Drug or device manufacturing" means the production, preparation,
41 propagation or processing of a drug or device, either directly or indirectly,
42 by extraction from substances of natural origin or independently by means of
43 chemical synthesis and includes any packaging or repackaging of substances or
44 labeling or relabeling of its container and the promotion and marketing of
45 the same. Drug or device manufacturing does not include compounding.

1 30. "Economic poison" means any substance that alone, in chemical
2 combination or in formulation with one or more other substances is a
3 pesticide within the meaning of the laws of this state or the federal
4 insecticide, fungicide and rodenticide act and that is used in the
5 production, storage or transportation of raw agricultural commodities.

6 31. "ENTERAL FEEDING" MEANS NOURISHMENT PROVIDED BY MEANS OF A TUBE
7 INSERTED INTO THE STOMACH OR INTESTINE.

8 ~~31-~~ 32. "Established name", with respect to a drug or ingredient of a
9 drug, means any of the following:

10 (a) The applicable official name.

11 (b) If there is no such name and the drug or ingredient is an article
12 recognized in an official compendium, the official title in an official
13 compendium.

14 (c) If neither subdivision (a) nor (b) of this paragraph applies, the
15 common or usual name of such drug.

16 ~~32-~~ 33. "Executive director" means the executive director of the board
17 of pharmacy.

18 ~~33-~~ 34. "Federal act" means the federal laws and regulations that
19 pertain to drugs, devices, poisons and hazardous substances and that are
20 official at the time any drug, device, poison or hazardous substance is
21 affected by this chapter.

22 ~~34-~~ 35. "Full service wholesale permittee" means a permittee who may
23 distribute prescription-only drugs and devices, controlled substances and
24 over-the-counter drugs and devices to pharmacies or other legal outlets from
25 a place devoted in whole or in part to wholesaling these items.

26 ~~35-~~ 36. "Graduate intern" means a person who has graduated from a
27 college, school or program of pharmacy approved by the board and who meets
28 the qualifications and experience for a pharmacy intern as provided in
29 section 32-1923.

30 ~~36-~~ 37. "Highly toxic" means any substance that falls within any of
31 the following categories:

32 (a) Produces death within fourteen days in half or more than half of a
33 group of ten or more laboratory white rats each weighing between two hundred
34 and three hundred grams, at a single dose of fifty milligrams or less per
35 kilogram of body weight, when orally administered.

36 (b) Produces death within fourteen days in half or more than half of a
37 group of ten or more laboratory white rats each weighing between two hundred
38 and three hundred grams, if inhaled continuously for a period of one hour or
39 less at an atmospheric concentration of two hundred parts per million by
40 volume or less of gas or vapor or two milligrams per liter by volume or less
41 of mist or dust, provided the concentration is likely to be encountered by
42 humans if the substance is used in any reasonably foreseeable manner.

43 (c) Produces death within fourteen days in half or more than half of a
44 group of ten or more rabbits tested in a dosage of two hundred milligrams or
45 less per kilogram of body weight, if administered by continuous contact with
46 the bare skin for twenty-four hours or less.

1 If the board finds that available data on human experience with any substance
2 indicate results different from those obtained on animals in the dosages or
3 concentrations prescribed in this paragraph, the human data shall take
4 precedence.

5 ~~37.~~ 38. "Hospital" means any institution for the care and treatment of
6 the sick and injured that is approved and licensed as a hospital by the
7 department of health services.

8 ~~38.~~ 39. "Intern" means a pharmacy intern and a graduate intern.

9 ~~39.~~ 40. "Internship" means the practical, experiential, hands-on
10 training of a pharmacy intern under the supervision of a preceptor.

11 ~~40.~~ 41. "Irritant" means any substance, other than a corrosive, that
12 on immediate, prolonged or repeated contact with normal living tissue will
13 induce a local inflammatory reaction.

14 ~~41.~~ 42. "Jurisprudence examination" means a ~~board-approved~~
15 BOARD-APPROVED pharmacy law examination that is written and administered in
16 cooperation with the national association of boards of pharmacy or another
17 ~~board-approved~~ BOARD-APPROVED pharmacy law examination.

18 ~~42.~~ 43. "Label" means a display of written, printed or graphic matter
19 on the immediate container of any article that, unless easily legible through
20 the outside wrapper or container, also appears on the outside wrapper or
21 container of the article's retail package. For the purposes of this
22 paragraph, the immediate container does not include package liners.

23 ~~43.~~ 44. "Labeling" means all labels and other written, printed or
24 graphic matter either:

25 (a) On any article or any of its containers or wrappers.

26 (b) Accompanying that article.

27 ~~44.~~ 45. "Letter of reprimand" means a disciplinary letter that is a
28 public document issued by the board and that informs a licensee or permittee
29 that the licensee's or permittee's conduct violates state or federal law and
30 may require the board to monitor the licensee or permittee.

31 ~~45.~~ 46. "Limited service pharmacy" means a pharmacy that is approved
32 by the board to practice a limited segment of pharmacy as indicated by the
33 permit issued by the board.

34 ~~46.~~ 47. "Manufacture" or "manufacturer" means every person who
35 prepares, derives, produces, compounds, processes, packages or repackages or
36 labels any drug in a place, other than a pharmacy, devoted to manufacturing
37 the drug.

38 ~~47.~~ 48. "Marijuana" has the same meaning prescribed in section
39 13-3401.

40 ~~48.~~ 49. "Medical practitioner" means any medical doctor, doctor of
41 osteopathy, dentist, podiatrist, veterinarian or other person WHO IS licensed
42 and authorized by law to use and prescribe drugs and devices for the
43 treatment of sick and injured human beings or animals or for the diagnosis or
44 prevention of sickness in human beings or animals in this state or any state,
45 territory or district of the United States.

1 ~~49.~~ 50. "Medication order" means a written or verbal order from a
2 medical practitioner or that person's authorized agent to administer a drug
3 or device.

4 ~~50.~~ 51. "Narcotic drug" has the same meaning prescribed in section
5 13-3401.

6 ~~51.~~ 52. "New drug" means either:

7 (a) Any drug the composition of which is such that the drug is not
8 generally recognized among experts qualified by scientific training and
9 experience to evaluate the safety and effectiveness of drugs as safe and
10 effective for use under the conditions prescribed, recommended or suggested
11 in the labeling.

12 (b) Any drug the composition of which is such that the drug, as a
13 result of investigations to determine its safety and effectiveness for use
14 under such conditions, has become so recognized, but that has not, other than
15 in the investigations, been used to a material extent or for a material time
16 under those conditions.

17 ~~52.~~ 53. "Nonprescription drug" or "over-the-counter drug" means any
18 nonnarcotic medicine or drug that may be sold without a prescription and is
19 prepackaged and labeled for use by the consumer in accordance with the
20 requirements of the laws of this state and federal law. Nonprescription drug
21 does not include:

22 (a) A drug that is primarily advertised and promoted professionally to
23 medical practitioners and pharmacists by manufacturers or primary
24 distributors.

25 (b) A controlled substance.

26 (c) A drug that is required to bear a label that states "Rx only". ~~"~~

27 (d) A drug THAT IS intended for human use by hypodermic injection.

28 ~~53.~~ 54. "Nonprescription drug wholesaler permittee" means a permittee
29 who may distribute only over-the-counter drugs and devices to pharmacies or
30 other lawful outlets from a place devoted in whole or in part to wholesaling
31 these items.

32 ~~54.~~ 55. "Notice" means personal service or the mailing of a copy of
33 the notice by certified mail addressed either to the person at the person's
34 latest address of record in the board office or to the person's attorney.

35 56. "NUTRITIONAL SUPPLEMENTATION" MEANS VITAMINS, MINERALS AND CALORIC
36 SUPPLEMENTATION. NUTRITIONAL SUPPLEMENTATION DOES NOT INCLUDE MEDICATION OR
37 DRUGS.

38 ~~55.~~ 57. "Official compendium" means the latest revision of the United
39 States pharmacopeia and the national formulary or any current supplement.

40 ~~56.~~ 58. "Other jurisdiction" means one of the other forty-nine states,
41 the District of Columbia, the Commonwealth of Puerto Rico or a territory of
42 the United States of America.

43 ~~57.~~ 59. "Package" means a receptacle defined or described in the
44 United States pharmacopeia and the national formulary as adopted by the
45 board.

1 ~~58.~~ 60. "Packaging" means the act or process of placing a drug item or
2 device in a container for the purpose or intent of dispensing or distributing
3 the item or device to another.

4 61. "PARENTERAL NUTRITION" MEANS INTRAVENOUS FEEDING THAT PROVIDES A
5 PERSON WITH FLUIDS AND ESSENTIAL NUTRIENTS THE PERSON NEEDS WHILE THE PERSON
6 IS UNABLE TO RECEIVE ADEQUATE FLUIDS OR FEEDINGS BY MOUTH OR BY ENTERAL
7 FEEDING.

8 ~~59.~~ 62. "Person" means an individual, partnership, corporation and
9 association, and their duly authorized agents.

10 ~~60.~~ 63. "Pharmaceutical care" means the provision of drug therapy and
11 other pharmaceutical patient care services.

12 ~~61.~~ 64. "Pharmacist" means an individual WHO IS currently licensed by
13 the board to practice the profession of pharmacy in this state.

14 ~~62.~~ 65. "Pharmacist in charge" means the pharmacist who is responsible
15 to the board for a licensed establishment's compliance with the laws and
16 administrative rules of this state and of the federal government pertaining
17 to the practice of pharmacy, the manufacturing of drugs and the distribution
18 of drugs and devices.

19 ~~63.~~ 66. "Pharmacist licensure examination" means a ~~board-approved~~
20 BOARD-APPROVED examination that is written and administered in cooperation
21 with the national association of boards of pharmacy or any other ~~board~~
22 ~~approved~~ BOARD-APPROVED pharmacist licensure examination.

23 ~~64.~~ 67. "Pharmacy" means any place:

24 (a) Where drugs, devices, poisons or related hazardous substances are
25 offered for sale at retail.

26 (b) In which the profession of pharmacy is practiced or where
27 prescription orders are compounded and dispensed.

28 (c) That has displayed on it or in it the words "pharmacist,"
29 "pharmaceutical chemist," "apothecary," "druggist," "pharmacy," "drugstore,"
30 "drugs" or "drug sundries" or any of these words or combinations of these
31 words, or words of similar import either in English or any other language, or
32 that is advertised by any sign containing any of these words.

33 (d) Where the characteristic symbols of pharmacy or the characteristic
34 prescription sign "Rx" is exhibited.

35 (e) Or a portion of any building or structure that is leased, used or
36 controlled by the permittee to conduct the business authorized by the board
37 at the address for which the permit was issued and that is enclosed and
38 secured when a pharmacist is not in attendance.

39 ~~65.~~ 68. "Pharmacy intern" means a person who has all of the
40 qualifications and experience prescribed in section 32-1923.

41 ~~66.~~ 69. "Pharmacy technician" means a person who is licensed pursuant
42 to this chapter.

43 ~~67.~~ 70. "Pharmacy technician trainee" means a person who is licensed
44 pursuant to this chapter.

1 ~~68-~~ 71. "Poison" or "hazardous substance" includes, but is not limited
2 to, any of the following if intended and suitable for household use or use by
3 children:

4 (a) Any substance that, according to standard works on medicine,
5 pharmacology, pharmacognosy or toxicology, if applied to, introduced into or
6 developed within the body in relatively small quantities by its inherent
7 action uniformly produces serious bodily injury, disease or death.

8 (b) A toxic substance.

9 (c) A highly toxic substance.

10 (d) A corrosive substance.

11 (e) An irritant.

12 (f) A strong sensitizer.

13 (g) A mixture of any of the substances described in this paragraph, if
14 the substance or mixture of substances may cause substantial personal injury
15 or substantial illness during or as a proximate result of any customary or
16 reasonably foreseeable handling or use, including reasonably foreseeable
17 ingestion by children.

18 (h) A substance **THAT IS** designated by the board to be a poison or
19 hazardous substance. This subdivision does not apply to radioactive
20 substances, economic poisons subject to the federal insecticide, fungicide
21 and rodenticide act or the state pesticide act, foods, drugs and cosmetics
22 subject to state laws or the federal act or substances intended for use as
23 fuels when stored in containers and used in the heating, cooking or
24 refrigeration system of a house. This subdivision applies to any substance
25 or article that is not itself an economic poison within the meaning of the
26 federal insecticide, fungicide and rodenticide act or the state pesticide
27 act, but that is a poison or hazardous substance within the meaning of this
28 paragraph by reason of bearing or containing an economic poison or hazardous
29 substance.

30 ~~69-~~ 72. "Practice of pharmacy" means furnishing the following health
31 care services as a medical professional:

32 (a) Interpreting, evaluating and dispensing prescription orders in the
33 patient's best interests.

34 (b) Compounding drugs pursuant to or in anticipation of a prescription
35 order.

36 (c) Labeling of drugs and devices in compliance with state and federal
37 requirements.

38 (d) Participating in drug selection and drug utilization reviews, drug
39 administration, drug or drug-related research and drug therapy monitoring or
40 management.

41 (e) Providing patient counseling necessary to provide pharmaceutical
42 care.

43 (f) Properly and safely storing drugs and devices in anticipation of
44 dispensing.

45 (g) Maintaining required records of drugs and devices.

1 (h) Offering or performing of acts, services, operations or
2 transactions necessary in the conduct, operation, management and control of a
3 pharmacy.

4 (i) Implementing, monitoring and modifying drug therapy pursuant to a
5 protocol-based drug therapy agreement with a provider as outlined in section
6 32-1970.

7 (j) Initiating and administering immunizations or vaccines pursuant to
8 section 32-1974.

9 ~~70-~~ 73. "Practitioner" means any physician, dentist, veterinarian,
10 scientific investigator or other person who is licensed, registered or
11 otherwise permitted to distribute, dispense, conduct research with respect to
12 or administer a controlled substance in the course of professional practice
13 or research in this state, or any pharmacy, hospital or other institution
14 that is licensed, registered or otherwise permitted to distribute, dispense,
15 conduct research with respect to or administer a controlled substance in the
16 course of professional practice or research in this state.

17 ~~71-~~ 74. "Preceptor" means a pharmacist who is serving as the practical
18 instructor of an intern and complies with section 32-1923.

19 ~~72-~~ 75. "Precursor chemical" means a substance that is:

20 (a) The principal compound that is commonly used or that is produced
21 primarily for use and that is an immediate chemical intermediary used or
22 likely to be used in the manufacture of a controlled substance, the control
23 of which is necessary to prevent, curtail or limit manufacture.

24 (b) Listed in section 13-3401, paragraph 26 or 27.

25 ~~73-~~ 76. "Prescription" means either a prescription order or a
26 prescription medication.

27 ~~74-~~ 77. "Prescription medication" means any drug, including label and
28 container according to context, that is dispensed pursuant to a prescription
29 order.

30 ~~75-~~ 78. "Prescription-only device" includes:

31 (a) Any device that is limited by the federal act to use under the
32 supervision of a medical practitioner.

33 (b) Any device required by the federal act to bear on its label
34 essentially the legend "Rx only".

35 ~~76-~~ 79. "Prescription-only drug" does not include a controlled
36 substance but does include:

37 (a) Any drug that because of its toxicity or other potentiality for
38 harmful effect, the method of its use, or the collateral measures necessary
39 to its use is not generally recognized among experts, qualified by scientific
40 training and experience to evaluate its safety and efficacy, as safe for use
41 except by or under the supervision of a medical practitioner.

42 (b) Any drug that is limited by an approved new drug application under
43 the federal act or section 32-1962 to use under the supervision of a medical
44 practitioner.

45 (c) Every potentially harmful drug, the labeling of which does not
46 bear or contain full and adequate directions for use by the consumer.

1 (d) Any drug, other than a controlled substance, required by the
2 federal act to bear on its label the legend "Rx only".

3 ~~77.~~ 80. "Prescription order" means any of the following:

4 (a) An order to a pharmacist for drugs or devices issued and signed by
5 a duly licensed medical practitioner in the authorized course of the
6 practitioner's professional practice.

7 (b) An order transmitted to a pharmacist through word of mouth,
8 telephone or other means of communication directed by that medical
9 practitioner. Prescription orders received by word of mouth, telephone or
10 other means of communication shall be maintained by the pharmacist pursuant
11 to section 32-1964, and the record so made by the pharmacist constitutes the
12 original prescription order to be dispensed by the pharmacist. This
13 paragraph does not alter or affect laws of this state or any federal act
14 requiring a written prescription order.

15 (c) An order initiated by a pharmacist pursuant to a protocol-based
16 drug therapy agreement with a provider as outlined in section 32-1970, or
17 immunizations or vaccines administered by a pharmacist pursuant to section
18 32-1974.

19 (d) A DIET ORDER OR AN ORDER FOR ENTERAL FEEDING, NUTRITIONAL
20 SUPPLEMENTATION OR PARENTERAL NUTRITION THAT IS INITIATED BY A REGISTERED
21 DIETITIAN OR OTHER QUALIFIED NUTRITION PROFESSIONAL IN A HOSPITAL PURSUANT TO
22 SECTION 36-416.

23 ~~78.~~ 81. "Professionally incompetent" means:

24 (a) Incompetence based on a variety of factors, including a lack of
25 sufficient pharmaceutical knowledge or skills or experience to a degree
26 likely to endanger the health of patients.

27 (b) When considered with other indications of professional
28 incompetence, a pharmacist, pharmacy intern or graduate intern who fails to
29 obtain a passing score on a ~~board-approved~~ BOARD-APPROVED pharmacist
30 licensure examination or a pharmacy technician or pharmacy technician trainee
31 who fails to obtain a passing score on a ~~board-approved~~ BOARD-APPROVED
32 pharmacy technician licensure examination.

33 ~~79.~~ 82. "Radioactive substance" means a substance that emits ionizing
34 radiation.

35 ~~80.~~ 83. "Safely engage in employment duties" means that a permittee or
36 the permittee's employee is able to safely engage in employment duties
37 related to the manufacture, sale, distribution or dispensing of drugs,
38 devices, poisons, hazardous substances, controlled substances or precursor
39 chemicals.

40 ~~81.~~ 84. "Symbol" means the characteristic symbols that have
41 historically identified pharmacy, including ~~"show globes"~~, AND ~~"mortar and~~
42 ~~pestle, "~~ and the sign "Rx".

43 ~~82.~~ 85. "Toxic substance" means a substance, other than a radioactive
44 substance, that has the capacity to produce injury or illness in humans
45 through ingestion, inhalation or absorption through any body surface.

1 ~~83.~~ 86. "Ultimate user" means a person who lawfully possesses a drug
2 or controlled substance for that person's own use, for the use of a member of
3 that person's household or for administering to an animal owned by that
4 person or by a member of that person's household.

5 Sec. 2. Title 36, chapter 4, article 1, Arizona Revised Statutes, is
6 amended by adding section 36-416, to read:

7 36-416. Registered dietitians; hospital orders; definition

8 A. A HOSPITAL THAT IS LICENSED PURSUANT TO SECTION 36-422 MAY ALLOW A
9 REGISTERED DIETITIAN OR OTHER QUALIFIED NUTRITION PROFESSIONAL TO ORDER
10 DIETS, ENTERAL FEEDING, NUTRITIONAL SUPPLEMENTATION OR PARENTERAL NUTRITION
11 IF AUTHORIZED BY MEDICAL STAFF PURSUANT TO 42 CODE OF FEDERAL REGULATIONS
12 SECTION 482.28(b) AND IF BOTH:

13 1. THE HOSPITAL'S WRITTEN POLICIES AND PROCEDURES ALLOW REGISTERED
14 DIETITIANS OR OTHER QUALIFIED NUTRITION PROFESSIONALS TO ISSUE SUCH ORDERS.

15 2. THE HOSPITAL HAS WRITTEN POLICIES AND PROCEDURES THAT ADDRESS THE
16 HOSPITAL'S RESPONSE TO ADVERSE EVENTS, IF ANY, THAT ARISE AS A RESULT OF
17 ORDERS ISSUED BY A REGISTERED DIETITIAN OR OTHER QUALIFIED NUTRITION
18 PROFESSIONAL.

19 B. FOR THE PURPOSES OF THIS SECTION:

20 1. "QUALIFIED NUTRITION PROFESSIONAL" MEANS A NUTRITION PROFESSIONAL
21 WHO IS DEEMED QUALIFIED BY A HOSPITAL FOR WHICH THE PERSON WORKS.

22 2. "REGISTERED DIETITIAN" MEANS A PERSON WHO MEETS THE QUALIFICATIONS
23 OF THE CREDENTIALING AGENCY FOR THE AMERICAN ACADEMY OF NUTRITION AND
24 DIETETICS.

APPROVED BY THE GOVERNOR MAY 17, 2016.

FILED IN THE OFFICE OF THE SECRETARY OF STATE MAY 17, 2016.