State of Arizona
House of Representatives
Fifty-fifth Legislature
Second Regular Session
2022

CHAPTER 98

HOUSE BILL 2490

AN ACT

AMENDING SECTION 32-1901, ARIZONA REVISED STATUTES; REPEALING SECTION 32-1970, ARIZONA REVISED STATUTES; AMENDING TITLE 32, CHAPTER 18, ARTICLE 3, ARIZONA REVISED STATUTES, BY ADDING A NEW SECTION 32-1970; RELATING TO THE ARIZONA STATE BOARD OF PHARMACY.

(TEXT OF BILL BEGINS ON NEXT PAGE)

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

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

32-1901. Definitions

In this chapter, unless the context otherwise requires:

1. "Administer" means directly applying a controlled substance, prescription-only drug, dangerous drug or narcotic drug, whether by injection, inhalation, ingestion or any other means, to the body of a patient or research subject by a practitioner or by the practitioner's authorized agent or the patient or research subject at the direction of the practitioner.

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

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

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

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

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

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

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

6. "Automated prescription-dispensing kiosk" means a mechanical system that is operated as an extension of a pharmacy, that maintains all transaction information within the pharmacy operating system, that is separately permitted from the pharmacy and that performs operations that either:

(a) Accept a prescription or refill order, store prepackaged or repackaged medications, label and dispense patient-specific prescriptions and provide counseling on new or refilled prescriptions.

(b) Dispense or deliver a prescription or refill that has been prepared by or on behalf of the pharmacy that oversees the automated prescription-dispensing kiosk.

7. "Board" or "board of pharmacy" means the Arizona state board of pharmacy.

8. "Certificate of composition" means a list of a product's ingredients.

9. "Certificate of free sale" means a document that authenticates a product that is generally and freely sold in domestic or international channels of trade.

10. "Color additive" means a material that either:

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

(b) If added or applied to a drug, or to the human body or any part of the human body, is capable of imparting color, except that color additive does not include any material that has been or may be exempted under the federal act. Color includes black, white and intermediate grays.

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

12. "Compressed medical gas distributor" means a person that holds a current permit issued by the board to distribute compressed medical gases to compressed medical gas suppliers and other entities that are registered, licensed or permitted to use, administer or distribute compressed medical gases.

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

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

15. "Compressed medical gas supplier" means a person that holds a current permit issued by the board to supply compressed medical gases pursuant to a compressed medical gas order and only to the consumer or the patient.

16. "Controlled substance" means a drug, substance or immediate precursor that is identified, defined or listed in title 36, chapter 27, article 2 or the rules adopted pursuant to title 36, chapter 27, article 2.

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

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

19. "Dangerous drug" has the same meaning prescribed in section 13-3401.

20. "Day" means a business day.

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

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

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

24. "Device", except as used in paragraph 18 of this section, section 32-1965, paragraph 4 and section 32-1967, subsection A, paragraph 15 and subsection C, means an instrument, apparatus or contrivance, including its components, parts and accessories, including all such items under the federal act, that is intended either:

(a) For use in diagnosing, curing, mitigating, treating or preventing disease in the human body or other animals.

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

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

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

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

28. "Dispenser" means a practitioner who dispenses.

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

30. "Distributor" means a person who distributes.

31. "Drug" means:

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

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

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

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

32. "Drug enforcement administration" means the drug enforcement administration of the United States department of justice or its successor agency.

33. "Drug or device manufacturing" means producing, preparing, propagating or processing a drug or device, either directly or indirectly, by extraction from substances of natural origin or independently by means of chemical synthesis and includes any packaging or repackaging of substances or labeling or relabeling of its container and promoting and marketing the same. Drug or device manufacturing does not include compounding.

34. "Durable medical equipment" means technologically sophisticated medical equipment as prescribed by the board in rule that a patient or consumer may use in a home or residence and that may be a prescription-only device.

35. "Durable medical equipment distributor":

(a) Means a person that stores or distributes durable medical equipment other than to the patient or consumer.

(b) Includes a virtual durable medical equipment distributor as prescribed in rule by the board.

36. "Durable medical equipment supplier":

(a) Means a person that sells, leases or supplies durable medical equipment to the patient or consumer.

(b) Includes a virtual durable medical equipment supplier as prescribed in rule by the board.

37. "Economic poison" means any substance that alone, in chemical combination with or in formulation with one or more other substances is a pesticide within the meaning of the laws of this state or the federal insecticide, fungicide and rodenticide act and that is used in producing, storing or transporting raw agricultural commodities.

38. "Enteral feeding" means nourishment that is provided by means of a tube inserted into the stomach or intestine.

39. "Established name", with respect to a drug or ingredient of a drug, means any of the following:

(a) The applicable official name.

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

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

40. "Executive director" means the executive director of the board of pharmacy.

41. "Federal act" means the federal laws and regulations that pertain to drugs, devices, poisons and hazardous substances and that are official at the time any drug, device, poison or hazardous substance is affected by this chapter.

42. "Full-service wholesale permittee":

(a) Means a permittee who may distribute prescription-only drugs and devices, controlled substances and over-the-counter drugs and devices to pharmacies or other legal outlets from a place devoted in whole or in part to wholesaling these items.

(b) Includes a virtual wholesaler as defined in rule by the board.

43. "Good manufacturing practice" means a system for ensuring that products are consistently produced and controlled according to quality standards and covering all aspects of design, monitoring and control of manufacturing processes and facilities to ensure that products do not pose any risk to the consumer or public.

44. "Highly toxic" means any substance that falls within any of the following categories:

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

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

(c) Produces death within fourteen days in half or more than half of a group of ten or more rabbits tested in a dosage of two hundred milligrams or less per kilogram of body weight, if administered by continuous contact with the bare skin for twenty-four hours or less. If the board finds that available data on human experience with any substance indicate results different from those obtained on animals in the dosages

or concentrations prescribed in this paragraph, the human data shall take precedence.

45. "Hospital" means any institution for the care and treatment of the sick and injured that is approved and licensed as a hospital by the department of health services.

46. "Intern" means a pharmacy intern.

47. "Internship" means the practical, experiential, hands-on training of a pharmacy intern under the supervision of a preceptor.

48. "Irritant" means any substance, other than a corrosive, that on immediate, prolonged or repeated contact with normal living tissue will induce a local inflammatory reaction.

49. "Jurisprudence examination" means a board-approved pharmacy law examination that is written and administered in cooperation with the national association of boards of pharmacy or another board-approved pharmacy law examination.

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

51. "Labeling" means all labels and other written, printed or graphic matter that either:

(a) Is on any article or any of its containers or wrappers.

(b) Accompanies that article.

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

53. "Limited service pharmacy" means a pharmacy that is approved by the board to practice a limited segment of pharmacy as indicated by the permit issued by the board.

54. "Manufacture" or "manufacturer":

(a) Means every person who prepares, derives, produces, compounds, processes, packages or repackages or labels any drug in a place, other than a pharmacy, that is devoted to manufacturing the drug.

(b) Includes a virtual manufacturer as defined in rule by the board.

55. "Marijuana" has the same meaning prescribed in section 13-3401.

56. "Medical practitioner" means any medical doctor, doctor of osteopathic medicine, dentist, podiatrist, veterinarian or other person who is licensed and authorized by law to use and prescribe drugs and devices to treat sick and injured human beings or animals or to diagnose or prevent sickness in human beings or animals in this state or any state, territory or district of the United States.

57. "Medication order" means a written or verbal order from a medical practitioner or that person's authorized agent to administer a drug or device.

58. "Narcotic drug" has the same meaning prescribed in section 13-3401.

59. "New drug" means either:

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

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

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

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

(b) A controlled substance.

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

(d) A drug that is intended for human use by hypodermic injection.

61. "Nonprescription drug wholesale permittee":

(a) Means a permittee who may distribute only over-the-counter drugs and devices to pharmacies or other lawful outlets from a place devoted in whole or in part to wholesaling these items.

(b) Includes a virtual wholesaler as defined in rule by the board.

62. "Notice" means personal service or the mailing of a copy of the notice by certified mail and email addressed either to the person at the person's latest address of record in the board office or to the person and the person's attorney using the most recent information provided to the board in the board's licensing database.

63. "Nutritional supplementation" means vitamins, minerals and caloric supplementation. Nutritional supplementation does not include medication or drugs.

64. "Official compendium" means the latest revision of the United States pharmacopeia and the national formulary or any current supplement.

65. "Other jurisdiction" means one of the other forty-nine states, the District of Columbia, the Commonwealth of Puerto Rico or a territory of the United States of America.

66. "Package" means a receptacle that is defined or described in the United States pharmacopeia and the national formulary as adopted by the board.

67. "Packaging" means the act or process of placing a drug item or device in a container for the purpose or intent of dispensing or distributing the item or device to another.

68. "Parenteral nutrition" means intravenous feeding that provides an individual with fluids and essential nutrients the individual needs while the individual is unable to receive adequate fluids or feedings by mouth or by enteral feeding.

69. "Person" means an individual, partnership, corporation and association, and their duly authorized agents.

70. "Pharmaceutical care" means the provision of drug therapy and other pharmaceutical patient care services.

71. "Pharmacist" means an individual who is currently licensed by the board to practice the profession of pharmacy in this state.

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

73. "Pharmacist licensure examination" means a board-approved examination that is written and administered in cooperation with the national association of boards of pharmacy or any other board-approved pharmacist licensure examination.

74. "Pharmacy":

(a) Means:

(i) Any place where drugs, devices, poisons or related hazardous substances are offered for sale at retail or where prescription orders are dispensed by a licensed pharmacist.

(ii) Any place that has displayed on it or in it the words "pharmaceutical chemist", "apothecary", "druggist", "pharmacy", "drugstore", "drugs" or "drug sundries" or any of these words or combinations of these words, or words of similar import either in English or any other language, or that is advertised by any sign containing any of these words.

(iii) Any place where the characteristic symbols of pharmacy or the characteristic prescription sign "Rx" is exhibited and where drugs are stored or dispensed.

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

(v) A remote dispensing site pharmacy.

(vi) A remote hospital site pharmacy, as defined by the board in rule, that operates under direct or remote supervision by a pharmacist pursuant to rules adopted by the board.

(b) Includes a satellite pharmacy.

75. "Pharmacy intern" means a person who has all of the qualifications and experience prescribed in section 32-1923.

76. "Pharmacy technician" means a person who is licensed pursuant to this chapter.

77. "Pharmacy technician trainee" means a person who is licensed pursuant to this chapter.

78. "Poison" or "hazardous substance" includes any of the following if intended and suitable for household use or use by children:

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

(b) A toxic substance.

(c) A highly toxic substance.

(d) A corrosive substance.

(e) An irritant.

(f) A strong sensitizer.

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

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

79. "Practice of pharmacy":

(a) Means furnishing the following health care services as a medical professional:

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

(ii) Compounding drugs pursuant to or in anticipation of a prescription order.

(iii) Labeling drugs and devices in compliance with state and federal requirements.

(iv) Participating in drug selection and drug utilization reviews, drug administration, drug or drug-related research and drug therapy monitoring or management.

(v) Providing patient counseling necessary to provide pharmaceutical care.

(vi) Properly and safely storing drugs and devices in anticipation of dispensing.

(vii) Maintaining required records of drugs and devices.

(viii) Offering or performing acts, services, operations or transactions that are necessary to conduct, operate, manage and control a pharmacy.

(ix) ~~Initiating, monitoring and modifying drug therapy~~ PROVIDING PATIENT CARE SERVICES pursuant to a ~~protocol-based drug therapy~~ COLLABORATIVE PRACTICE agreement with a provider as outlined in section 32-1970.

(x) Initiating and administering immunizations or vaccines pursuant to section 32-1974.

(b) Does not include initiating a prescription order for any medication, drug or other substance used to induce or cause a medication abortion as defined in section 36-2151.

80. "Practitioner" means any physician, dentist, veterinarian, scientific investigator or other person who is licensed, registered or otherwise permitted to distribute, dispense, conduct research with respect to or administer a controlled substance in the course of professional practice or research in this state, or any pharmacy, hospital or other institution that is licensed, registered or otherwise permitted to distribute, dispense, conduct research with respect to or administer a controlled substance in the course of professional practice or research in this state.

81. "Preceptor" means a pharmacist who is serving as the practical instructor of an intern and who complies with section 32-1923.

82. "Precursor chemical" means a substance that is:

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

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

83. "Prescription" means either a prescription order or a prescription medication.

84. "Prescription medication" means any drug, including label and container according to context, that is dispensed pursuant to a prescription order.

85. "Prescription-only device" includes:

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

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

86. "Prescription-only drug" does not include a controlled substance but does include:

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

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

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

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

87. "Prescription order" means any of the following:

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

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

(c) An order that is initiated by a pharmacist pursuant to a ~~protocol-based drug therapy~~ COLLABORATIVE PRACTICE agreement with a provider as outlined in section 32-1970, or immunizations or vaccines administered by a pharmacist pursuant to section 32-1974.

(d) A diet order or an order for enteral feeding, nutritional supplementation or parenteral nutrition that is initiated by a registered dietitian or other qualified nutrition professional in a hospital pursuant to section 36-416.

88. "Professionally incompetent" means:

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

(b) When considered with other indications of professional incompetence, a pharmacist or pharmacy intern who fails to obtain a passing score on a board-approved pharmacist licensure examination or a pharmacy technician or pharmacy technician trainee who fails to obtain a passing score on a board-approved pharmacy technician licensure examination.

89. "Radioactive substance" means a substance that emits ionizing radiation.

90. "Remote dispensing site pharmacy" means a pharmacy where a pharmacy technician or pharmacy intern prepares, compounds or dispenses prescription medications under remote supervision by a pharmacist.

91. "Remote supervision by a pharmacist" means that a pharmacist directs and controls the actions of pharmacy technicians and pharmacy interns through the use of audio and visual technology.

92. "Revocation" or "revoke" means the official cancellation of a license, permit, registration or other approval authorized by the board for a period of two years unless otherwise specified by the board. A request or new application for reinstatement may be presented to the board for review before the conclusion of the specified revocation period upon review of the executive director.

93. "Safely engage in employment duties" means that a permittee or the permittee's employee is able to safely engage in employment duties related to the manufacture, sale, distribution or dispensing of drugs, devices, poisons, hazardous substances, controlled substances or precursor chemicals.

94. "Satellite pharmacy" means a work area located within a hospital or on a hospital campus that is not separated by other commercial property or residential property, that is under the direction of a pharmacist, that is a remote extension of a centrally licensed hospital pharmacy, that is owned by and dependent on the centrally licensed hospital pharmacy for administrative control, staffing and drug procurement and that is not required to be separately permitted.

95. "Symbol" means the characteristic symbols that have historically identified pharmacy, including show globes and mortar and pestle, and the sign "Rx".

96. "Third-party logistics provider" means an entity that provides or coordinates warehousing or other logistics services for the following items, but that does not take ownership of the items, and that distributes those items as directed by a manufacturer, wholesaler, dispenser or durable medical equipment supplier that is permitted by the board:

(a) Narcotic drugs or other controlled substances.

(b) Dangerous drugs as defined in section 13-3401.
(c) Prescription-only drugs and devices.
(d) Nonprescription drugs and devices.
(e) Precursor chemicals.
(f) Regulated chemicals as defined in section 13-3401.

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

98. "Ultimate user" means a person who lawfully possesses a drug or controlled substance for that person's own use, for the use of a member of that person's household or for administering to an animal owned by that person or by a member of that person's household.

Sec. 2. Repeal

Section 32-1970, Arizona Revised Statutes, is repealed.

Sec. 3. Title 32, chapter 18, article 3, Arizona Revised Statutes, is amended by adding a new section 32-1970, to read:

32-1970. Collaborative practice agreements; requirements; rules; definitions

A. A PHARMACIST WHO IS LICENSED PURSUANT TO THIS CHAPTER MAY ENTER INTO A COLLABORATIVE PRACTICE AGREEMENT WITH A PROVIDER PURSUANT TO THIS SECTION TO INITIATE, MONITOR AND MODIFY DRUG THERAPY OR PROVIDE DISEASE MANAGEMENT ASSISTANCE. THE COLLABORATIVE PRACTICE AGREEMENT MAY BE BETWEEN ONE OR MORE PHARMACISTS AND ONE OR MORE PROVIDERS. THE COLLABORATIVE PRACTICE AGREEMENT SHALL:

1. OUTLINE THE DUTIES RELATED TO DRUG THERAPY AND DISEASE MANAGEMENT THAT THE PROVIDER IS DELEGATING TO THE PHARMACIST TO PERFORM, INCLUDING DRUG THERAPY THAT THE PHARMACIST MAY INITIATE, MONITOR AND MODIFY AND LABORATORY TESTS THAT THE PHARMACIST MAY ORDER, AND THE ELIGIBLE GROUP OF PATIENTS THAT MAY BE TREATED UNDER THE COLLABORATIVE PRACTICE AGREEMENT.

2. SPECIFY, AT A MINIMUM, THE CONDITIONS TO BE MANAGED BY THE PHARMACIST THROUGH DISEASE MANAGEMENT AND DRUG THERAPY MANAGEMENT, THE CIRCUMSTANCES FOR WHICH THE PHARMACIST MUST NOTIFY THE PROVIDER AND ANY DOCUMENTATION OR RECORDKEEPING REQUIREMENTS.

3. SPECIFY THAT THE PHARMACIST MUST FOLLOW THE WRITTEN DRUG THERAPY AND DISEASE MANAGEMENT GUIDELINES PROVIDED BY THE PROVIDER AND MAY PROVIDE DRUG THERAPY AND DISEASE MANAGEMENT SERVICES ONLY PURSUANT TO THOSE GUIDELINES. THE GUIDELINES SHALL SPECIFY, AT A MINIMUM, THE SPECIFIC DRUG, DRUGS OR DRUG CLASSES AND THE CONDITIONS TO BE MANAGED BY THE PHARMACIST, THE CONDITIONS AND EVENTS FOR WHICH THE PHARMACIST MUST NOTIFY THE PROVIDER AND THE LABORATORY TESTS THE PHARMACIST MAY ORDER.

B. A PROVIDER WHO ENTERS INTO A COLLABORATIVE PRACTICE AGREEMENT UNDER THIS SECTION MUST HAVE A PREVIOUSLY ESTABLISHED PROVIDER-PATIENT RELATIONSHIP WITH A PATIENT IN ORDER FOR THAT PATIENT TO BE A PART OF THE

ELIGIBLE GROUP OF PATIENTS WHO MAY BE INCLUDED UNDER THE COLLABORATIVE PRACTICE AGREEMENT.

C. A LICENSEE WHO VIOLATES THIS SECTION COMMITS AN ACT OF UNPROFESSIONAL CONDUCT.

D. A PHARMACIST IS RESPONSIBLE FOR THE PHARMACIST'S NEGLIGENT ACTS THAT ARE THE RESULT OF THE CLINICAL DECISIONS MADE PURSUANT TO THE COLLABORATIVE PRACTICE AGREEMENT. THIS SUBSECTION DOES NOT LIMIT A PROVIDER'S LIABILITY FOR NEGLIGENT ACTS THAT ARE NOT RELATED TO A PHARMACIST'S CHANGE OF MEDICATION PURSUANT TO THE COLLABORATIVE PRACTICE AGREEMENT.

E. THE PHARMACIST SHALL MAINTAIN A COPY OF THE COLLABORATIVE PRACTICE AGREEMENT AND MAKE THE COLLABORATIVE PRACTICE AGREEMENT AVAILABLE TO THE BOARD ON REQUEST.

F. THE ARIZONA STATE BOARD OF PHARMACY, THE ARIZONA MEDICAL BOARD, THE ARIZONA BOARD OF OSTEOPATHIC EXAMINERS IN MEDICINE AND SURGERY AND THE ARIZONA STATE BOARD OF NURSING MAY ADOPT RULES RELATING TO COLLABORATIVE PRACTICE AGREEMENTS.

G. FOR THE PURPOSES OF THIS SECTION:

1. "COLLABORATIVE PRACTICE AGREEMENT" MEANS AN AGREEMENT BETWEEN A PHARMACIST AND A PROVIDER THAT OUTLINES THE DRUG THERAPY AND DISEASE MANAGEMENT SERVICES, INCLUDING INITIATING, MONITORING AND MODIFYING PRESCRIPTION DRUG AND LABORATORY TEST ORDERS THAT ARE AUTHORIZED BY THE PROVIDER AND DELEGATED TO THE PHARMACIST FOR THE PURPOSES OF DRUG THERAPY MANAGEMENT OR DISEASE MANAGEMENT BASED ON THE PHARMACIST'S SKILLS OR TRAINING.

2. INITIATE, MONITOR AND MODIFY DOES NOT INCLUDE A PHARMACIST'S SELECTION OF DRUG PRODUCTS THAT ARE NOT PRESCRIBED BY THE PROVIDER UNLESS THE SELECTION OF SPECIFIC DRUG PRODUCTS IS AUTHORIZED BY THE COLLABORATIVE PRACTICE AGREEMENT.

3. "PROVIDER" MEANS A PHYSICIAN WHO IS LICENSED PURSUANT TO CHAPTER 13 OR 17 OF THIS TITLE OR A REGISTERED NURSE PRACTITIONER WHO IS LICENSED PURSUANT TO CHAPTER 15 OF THIS TITLE.

APPROVED BY THE GOVERNOR MARCH 30, 2022.

FILED IN THE OFFICE OF THE SECRETARY OF STATE MARCH 30, 2022.