Agency # 070.00

### REGULATIONRULE 9 — PHARMACEUTICAL CARE/PATIENT

COUNSELING

09-00: PATIENT COUNSELING

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### 09-00-0001--PATIENT INFORMATION, DRUG USE EVALUATION, AND PATIENT RESEARCH **COUNSELING**

The intent of this regulation is to improve pharmaceutical care by defining basic standards of care. Pharmacy care/pharmaceutical care is defined as the responsible provision of drug therapy for the purpose of achieving definite outcomes that improve a patient's quality of life. These outcomes are: (1) cure of disease, (2) elimination or reduction of a patient's symptomatology, (3) arresting or slowing a disease process, or (4) preventing a disease or symptomatology.

Pharmaceutical care (clinical pharmacy) involves four major functions on behalf of the patient: (1) identifying potential and actual drug-related problems, (2) resolving actual drug related problems, (3) preventing potential drug-related problems, and (4) optimizing patient therapy outcomes. It is recognized that the patient might be best served if medication is not provided.

#### (a) Patient information (profile)

In order to effectively counsel patients, the pharmacist must, through communication with the patient or caregiver, make a reasonable effort to obtain, record, and maintain the following information for each patient. It is recognized that most of this can be obtained using pharmacy technicians and designed forms, etc.

- (1) Name, address, telephone number;
- (2) Date of birth (age);
- (3) Gender;
- (4) Medical history
  - (A) Significant patient health problems known to the pharmacist;
  - (B) Prescription drug reactions/prescription drug allergies;
  - (C) List of prescription medications and legend drug administration devices known to the pharmacist.
- (5) Transitory patients or situations where the pharmacy will only provide medication one

In obtaining patient information, if the pharmacist knows or is informed by the patient that this is a one-time situation, the pharmacist may forego the above requirement to record and maintain the information.

- (6) Pharmacist comments
- (b) Drug use evaluation for new and refill prescriptions

Drug use evaluation or drug utilization review includes the following activities:

- (1) The pharmacist shall evaluate the prescription or medication order for:
  - (A) Reasonable dose and route of administration;
  - (B) Reasonable directions for use.
- (2) The pharmacist shall evaluate medication orders and patient information for:

- (A) Duplication of therapy is the patient taking the same or similar medication(s)?;
- (B) Prescription drug-prescription drug interactions;
- (C) Proper utilization (over or underutilization);
- (D) Known drug allergies.
- (3) Drug-drug contraindications as defined by the Board. (Is this medication contraindicated with another medication the patient is taking?)
- (4) It is recognized that the ultimate decision to use the medication or not use the medication rests with the physician who has more complete patient information. It is the pharmacist's responsibility to monitor the patient's medication therapy in the areas addressed in this regulation and inform the physician of the suspected problem.
- (5) If a problem is suspected and the physician is informed, the pharmacist shall document the process.

#### (c) Patient counseling.

- (1) A pharmacist shall counsel the patient or caregiver "face to face" if the patient or caregiver is in the pharmacy. If not, a pharmacist shall make a reasonable effort to counsel the patient or caregiver;
- (2) Alternative forms of patient information may be used to supplement, but not replace face-to-face patient counseling;
- (3) Patient counseling, as described herein, shall also be required for outpatients of hospitals and institutions when medications are dispensed on discharge from the hospital or institution.
- (4) Patient counseling as described in this regulation shall not be required for inpatients of a hospital or institution where a nurse or other licensed health care professional is authorized to administer the medication. However, the pharmacist shall provide drug therapy counseling it is when professionally deemed to be appropriate and when medications are provided by the pharmacy, and when a pharmacist is on duty and a patient is discharged from the hospital or institution.
- (5) The pharmacist shall maintain and make available to all patients appropriate patientoriented reference materials USP-DI or *Facts and Comparisons Patient Drug Facts* or an equivalent or better publication as determined by the Board.
- (6) It is recognized that the ultimate decision to not provide patient counseling rests with the physician. If the physician in specific instances (blanket requests not accepted) requests that information NOT be provided to the patient and gives reason, the pharmacist should honor that request in almost all instances.
- (d) "Patient counseling" shall mean the effective communication by the pharmacist of information, as defined in this act to the patient or caregiver, in order to improve therapeutic outcome by encouraging proper use of prescription medications and drug delivery devices.
  - (1) For original prescription medication orders, (excluding renewed or updated prescriptions the patient has been recently taking) and orders for legend devices, specific areas of counseling shall include:
    - (A) Name and general description of the medication dispensed, i.e. antibiotic, antihistamine, blood pressure medicine, etc.

- (B) Name, general description and directions for use of drug delivery devices, i.e., insulin syringes, morphine pump, etc.
- (C) Explanation of route of administration, dosage, times of administration, and continuity of therapy;
- (D) Special directions for storage as deemed necessary by the pharmacist;
- (E) If the drug has been determined to have a significant side effect by the Board of Pharmacy, the patient shall be properly counseled to the extent deemed necessary by the pharmacist.
- (F) When the prescription drug dispensed has a significant side effect, if taken with over-the-counter drugs, the pharmacist should counsel the patient about that interaction. (Example: coumadin with aspirin)
- (G) If the prescription medication is significantly affected by food or diet, the pharmacist should so advise the patient. (Example: tetracycline with milk or food)
- (H) The pharmacist shall inform the patient or caregiver that he/she is available to answer questions about medications or general health information.
- (2) Refills--On refills the pharmacist shall present the opportunity for the patient or caregiver to ask questions. However, counseling on refills is not required except when needed in the professional judgment of the pharmacist.
- (d) Drug interactions significant side effects

Recognizing that a pharmacist cannot be expected to recognize all possible drug interactions and also recognizing that the pharmacist and the patient do not have time to explain the numerous side effects of drugs, the pharmacy shall maintain a computer program which will identify significant drug interactions. (These are drugs with side effects which may be managed most effectively if the patient is aware of the specific side effect and what to do if it occurs.) The pharmacist in charge will be responsible for assuring that the computer system adequately flags and warns the pharmacist of any occurrence of significant drug interactions or significant side effects. (If a pharmacy was in business before September 1, 1997, and at that time, did not have a computer system, said pharmacy may substitute *Patient Drug Facts* or other drug interaction manuals to reference drug interactions and side effects for effective patient counseling. This method should only be used until such time as the pharmacy acquires an adequate computer program as described in this section.) The pharmacist will be responsible for counseling the patient on these interactions with verbal and, where appropriate, written information. (2/12/91, 2/10/98, 07/15/2004)

### 09-00-0002—PRESCRIPTION ORDERS TO ADMINISTER MEDICATION AND/OR IMMUNIZATIONS

Except as limited by these rules or Arkansas statutes §17-92-101, an Arkansas licensed pharmacist, intern or pharmacy technician has the ability to administer medications they have been trained to administer.

Authority to administer medications/immunizations:

(1) An Authority to Administer is a written protocol, as defined in ACA § 17-92-101, from a practitioner for administration by a pharmacist of an approved medication or immunization.

- (2) Pharmacists may provide pharmaceutical care to patients seven (7) years of age and older by administering medications or immunizations to an eligible patient upon receiving an Authority to Administer or a valid prescription order by a practitioner so authorized to prescribe such medications or immunizations as provided in ACA § 17-92-101(16)(A)(i). After completing the course of study described in (b)(5)(B)—(E) of this section, licensed interns, as defined by Regulation 02-01-0003 (a), may provide pharmaceutical care to patients seven (7) years of age and older by administering medications or immunizations to an eligible patient, under the supervision of an appropriately licensed pharmacist with an Authority to Administer and in accordance with Regulations 02-01-004 and 02-01-0005(h).
- (2) An Authority to Administer, once granted, is valid for a time period not to exceed one (1) year—unless such an order is invalidated by the practitioner granting the authority.
- (3) An Authority to Administer is valid only for the pharmacist meeting the requirements set forth by the Arkansas State Board of Pharmacy and is not transferable.
- (4) Unless otherwise specifically authorized by the Board, a person must possess a
  Certification for the Authority to Administer Medications/Immunizations issued by
  the Board to be qualified to accept an Authority to Administer. Certification for the
  credential (Authority to Administer Medications/Immunizations) will be issued to
  pharmacists who:
  - (A) obtain and maintain a license to practice pharmacy issued by the Arkansas State Board of Pharmacy;
  - (B) successfully complete a Board approved course of study, examination, and certification consisting of a training program that includes the current guidelines and recommendations of the Centers of Disease Control and Prevention. The course of study should include, at a minimum:
    - (i) basic immunology, including the human immune response;
    - (ii) the mechanism of immunity, adverse effects, dose, and administration schedule of available vaccines and approved medication/immunization;
    - (iii)how to handle an emergency situation in the event one should arise as a result of the administration of the medication /immunization;
    - (iv)how to persuade patients to be immunized and options for record keeping for patients that do get immunized;
    - (v) how to administer subcutaneous, intradermal, and intramuscular injection; and (vi)record keeping requirements for these medications as required by law or regulation.
  - (C) obtain supervised instructions on the physical administration of vaccines during such course of study and certification;
  - (D) obtain and maintain current certification in Cardiopulmonary Resuscitation (CPR) or Basic Cardiac Life Support (BCLS), these certification courses must contain a live component where proficiency is tested; and
  - (E) successful completion of the above described course of study may be accomplished by:
    - (i) successfully completing the Board-approved course of study in a College of Pharmacy curriculum; or

- (ii) successfully completing an American Council of Pharmaceutical
  Education (ACPE) Certificate Program of not less than twelve (12) hours
  on the course of study described in paragraph (b)(5)(B) above.
- (F) The College of Pharmacy or the provider of said course of study shall provide participants a certificate of completion. A copy of said certificate shall be mailed to the Board of Pharmacy offices and placed in the pharmacist's permanent file.
- (5) Pharmacists who complete items (A) through (E) of section (5) above may apply to the Board for a Certification for the Authority to Administer

  Medications/Immunizations. The certificate is valid until the pharmacist's license expires. The certificate shall be displayed in the pharmacy at which the pharmacist is working, and may be renewed when the pharmacist renews his or her license biennially after demonstrating continuing competency for certification.
- (6) Continuing competency for certification for Authority to Administer must be maintained. A minimum of two (2) of the thirty (30) hour requirement for continuing education, each biennium, must be dedicated to this area of practice. In addition, the pharmacist must maintain a current certificate in cardiopulmonary resuscitation (CPR) or basic cardiac life support (BCLS).
- (7) An Authority to Administer order shall meet the following requirements:
  - (A) must properly identify the practitioner issuing the order;
  - (B) must identify the medication or vaccine covered in any such order;
  - (C) must identify the medication or vaccine administered, site of the administration, dose administered, identity of pharmacist administering the dose; and
  - (D) must bear the date of the original order.
- (c) Record keeping: Pharmacists shall maintain the following information for a minimum of two years:
  - (1) Authority to Administer
  - (2) Signed Patient Consent Form containing at least the following information
    - (a) Name of Patient
    - (b) Description of the medication or vaccine
    - (c) Description of the risks and possible side effects of the medication or vaccine
    - (d) Lot number of the medication or vaccine
    - (e) Expiration date of the medication or vaccine
    - (f) Date of administration

(Revised 07/15/2004, 03/14/2006, 7/5/2007, 7/27/2011 and 12/1/2017)

#### 09-01: DISEASE STATE MANAGEMENT

#### 09-01-0001 DISEASE STATE MANAGEMENT

The purpose of this regulation is to provide standards for the maintenance of records of a pharmacist engaged in the provision of disease state management as authorized in §17-92-101 (16) and §17-92-205 (a).

- (a) Definitions. The following words and terms, when used in this regulation, shall have the following meanings, unless the context clearly indicates otherwise:
  - (1) "Act" means the Arkansas Pharmacy Practice Act
  - (2) "Board" means the Arkansas State Board of Pharmacy

- (3) "Confidential record" means any health related record maintained by a pharmacy or pharmacist—such as a patient medication record, prescription drug order, or medication order.
- (4) "Disease state management" means the performance of specific acts of disease state management delegated to a pharmacist for an individual patient by an authorized practitioner through written protocol. (Disease state management shall not include the selection of drug products not prescribed by the practitioner, unless the drug product is named in the practitioner initiated protocol.)
- (5) "Written protocol" means a practitioner's order, standing medical order, standing delegation order, or other order or protocol as defined by rule of the Arkansas State Medical Board under the Medical Practice Act.
  - (A) A written protocol must contain at a minimum the following:
    - (i) A statement identifying the individual practitioner(s) authorized to prescribe drugs and responsible for the delegation of disease state management;
    - (ii) A statement identifying the individual pharmacist(s) authorized to dispense drugs and to engage in disease state management delegated by the practitioner;
    - (iii) A statement identifying the types of disease state management decisions that the pharmacist is authorized to make which shall include:
      - (a) A statement of the ailments or diseases involved, drugs, and types of drug therapy management authorized; and
      - (b) A specific statement of the procedures, decision criteria, or plan the pharmacist shall follow when exercising disease state management authority
    - (iv) A statement of the activities the pharmacist shall follow in the course of exercising disease state management authority, including the method for documenting decisions made and a plan for communication or feedback to the authorizing physician concerning specific decisions made. Documentation shall be recorded within a reasonable time of each intervention and may be performed on the patient medication record, patient medical chart, or in a separate log book; and
    - (v) A statement that describes appropriate mechanisms and time schedule for the pharmacist to report to the physician monitoring the pharmacist's exercise of delegated disease state management and the results of the disease state management.
  - (B) A standard protocol may be used, or the attending practitioner may develop a disease state management protocol for the individual patient. If a standard protocol is used, the practitioner shall record, what deviations if any, from the standard protocol are ordered for that patient;
  - (C) Maintenance of records:
    - (i) Every patient record required to be kept under this regulation shall be kept by the pharmacist and be available, for at least two (2) years from the date of such record, for inspecting and copying by the Board or its representative and to other authorized local, state, or federal law enforcement or regulatory agencies.

- (ii) Patient records may be maintained in an alternative data retention system, such as a data processing system or direct imaging system provided:
  - (a) The records maintained in the alternative system contain all of the information required on a manual record; and
  - (b) The data processing system is capable of producing a hard copy of the record upon the request of the Board, its representative, or other authorized local, state, or federal law enforcement or regulatory agencies.

#### (D) Written protocol:

- (i) A copy of the written protocol and any patient-specific deviations from the protocol shall be maintained by the pharmacist and available for inspection by a Board Inspector upon request.
- (ii) Written protocols, including standard protocols, any patient specific deviations from a standard protocol, and any individual patient protocol, shall be reviewed by the practitioner and pharmacist at least annually and revised, if necessary. Such review shall be documented in the pharmacist's records. Documentation of all services provided to the patient, by the pharmacist, shall be reviewed by the physician on the schedule established in the protocol.
- (iii) Any protocol from a practitioner shall be maintained in the pharmacy and available for inspection by a Board Inspector upon request.

#### (E) Confidentiality:

- (i) A pharmacist shall provide adequate security to prevent indiscriminate or unauthorized access to confidential records. If confidential health information is not transmitted directly between a pharmacy and a practitioner, but is transmitted through a data communication device, the confidential health information may not be accessed or maintained by the operator of the data communication device unless specifically authorized to obtain the confidential information by this regulation.
- (ii) Confidential records are privileged and may be released only to:
  - (a) the patient or the patient's agent;
  - (b) practitioners and other pharmacists when, in the pharmacist's professional judgment, such release is necessary to protect the patient's health and wellbeing;
  - (c) other persons, the Board, or other state or federal agencies authorized by law to receive such information:
  - (d) a law enforcement agency engaged in investigation of suspected violations of the Controlled Substances Act; or
  - (e) a person employed by any state agency which licenses a practitioner as defined in the Act if such person is engaged in the performance of the person's official duties.
- (iii) This regulation shall not affect or alter the provisions relating to the confidentiality of the physician patient communication as specified in the Medical Practice Act.

  (Adopted 8/19/99)

# 09-01-0003 — QUALIFICATIONS, RESOURCES, AND RECORD KEEPING REQUIRED FOR PRACTICING DISEASE STATE MANAGEMENT IN ARKANSAS.

- (a) To practice disease state management a pharmacist must:
  - (1) be a licensed pharmacist in the State of Arkansas
  - (2) complete requirements for a credential as established by a Board of Pharmacy approved organization.
- (b) Resource requirements for the provision of disease state management services shall include but are not limited to the following:
  - (1) Maintain a distinct area that provides privacy for the provision of disease state management services;
  - (2) Maintain references that include a current copy/edition of applicable national practice guidelines and such other resources as may be necessary for the provision of optimal care;
  - (3) Maintain devices, supplies, furniture, and equipment as may be needed for the provision of optimal care.
- (c) Record keeping requirements for disease state management.

  The pharmacist shall record, maintain, and transfer data essential to the continuity of care and consistent with all applicable state and federal laws and regulations; and these records and all related files shall be available to the Arkansas State Board of Pharmacy inspectors and professional staff upon request. Additionally, a transferable pharmaceutical care record is to be maintained and is to include:
  - (1) The written request for consultation from the patient or physician;
  - (2) The physician approved protocol and/or patient care plan, which is to recognize all concomitant diseases and the patient's complete medication history/profile;
  - (3) Pharmacy progress notes; and,
  - (4) Laboratory data. (Adopted 8/19/99, Revised 11/12/2009)

#### 09-01-0004 MINIMUM COMPETENCIES AND STANDARDS

- (a) Minimum competencies for pharmaceutical care in all disease state management areas:
  - (1) The pharmacist shall be capable of identifying and accessing the patient's current health status, health related needs and problems, and desired therapeutic outcomes.
  - (2) The pharmacist shall be capable of implementing, and evaluating a pharmaceutical care plan that assures the appropriateness of the patient's medication(s), dosing regimens, dosage forms, routes of administration, and delivery systems.
  - (3) The pharmacist shall be capable of communicating appropriate information to the patient and/or caregiver and other health care professionals regarding prescription or non-prescription medications and/or medical devices, disease states, or medical conditions, and the maintenance of health and wellness.
  - (4) The pharmacist shall be capable of monitoring and documenting the patient's progress toward identified endpoints and outcomes of the pharmaceutical care plan and shall intervene when appropriate.

    (Adopted 8/19/99, Revised 11/12/2009)

### 09-01-0005 NOTIFICATION OF CREDENTIAL IN DISEASE STATE MANAGEMENT REQUIRED:

Every pharmacist who receives a credential in disease state management from a Board approved organization must provide a copy of the credential to the Board of Pharmacy office. The Board of Pharmacy will notify any party requesting notification that the pharmacist is so qualified.

(Adopted 8/19/99, Revised 8/2001, 07/2004, 11/12/2009)

# Stricken language would be deleted from and underlined language would be added to present law. Act 63 of the Regular Session

1	State of Arkansas	A D:11	
2	93rd General Assembly	A Bill	
3	Regular Session, 2021		HOUSE BILL 1174
4			
5	By: Representative Boyd		
6	By: Senator Bledsoe		
7			
8		For An Act To Be Entitled	
9	AN ACT TO E	ELIMINATE THE NURSING HOME CONSULTA	NT
10	PHARMACIST	PERMIT AND THE DISEASE STATE MANAG	EMENT
11	CREDENTIAL;	AND FOR OTHER PURPOSES.	
12			
13			
14		Subtitle	
15	TO ELI	IMINATE THE NURSING HOME CONSULTANT	נ
16	PHARMA	ACIST PERMIT AND THE DISEASE STATE	
17	MANAGE	EMENT CREDENTIAL.	
18			
19			
20	BE IT ENACTED BY THE GE	NERAL ASSEMBLY OF THE STATE OF ARK	ANSAS:
21			
22		sas Code § 17-92-101(2), concerning	
23		g to pharmacists and pharmacies, i	s amended to read as
24	follows:		
25		ntialing" means the issuance of or	•
26		Pharmacy of a credential issued to	- ·
27		Arkansas State Board of Pharmacy co	
28		standards of competency established	-
29		for disease state management or of	ther pharmacy
30	services necessitating	a credential;	
31	CECHTON O A.l.	0 1 . 8 . 17 . 00 . 100 ( ) ( ) ( )	
32		sas Code § 17-92-108(a)(9), concern	ning fees charged by
33 34		d of Pharmacy, is repealed.	
35		rhormogist normit shall not expect	
36	dollars (\$35.00).	pharmacist permit shall not exceed	: tnirty-five
J 0	GOTTALD (933+UU).		

1	(B) The fee for the renewal of a nursing home consultant
2	pharmacist permit shall not exceed thirty-five dollars (\$35.00) per year;
3	
4	SECTION 3. Arkansas Code § 17-92-310(a)(1)(A), concerning failure to
5	renew a license, permit, or credential with the Arkansas State Board of
6	Pharmacy, is amended to read as follows:
7	(a)(l)(A) All retail pharmacy permits, out-of-state pharmacy permits,
8	specialty pharmacy permits, nursing home consultant pharmacist permits, and
9	pharmacist licenses shall expire on December 31 of the first odd-numbered
10	year following the date of issuance.
11	
12	SECTION 4. Arkansas Code § 17-92-310(b), concerning failure to renew a
13	license, permit, or credential with the Arkansas State Board of Pharmacy, is
14	amended to read as follows:
15	(b) The penalty for late payment of renewal for pharmacists,
16	pharmacies, wholesaler/manufacturer of legend drugs and controlled
17	substances, hospital, and institutional, and nursing home consultant permits
18	shall be as listed in § 17-92-108, and if renewal remains unpaid $\Theta + \Phi$ April
19	l of any the year, the license shall be void.
20	
21	SECTION 5. Arkansas Code § 17-92-316(a) and (b), concerning
22	credentials required for professional pharmacy services, are amended to read
23	as follows:
24	(a)(1) The Arkansas State Board of Pharmacy may provide by rule for
25	credentialing and approval of pharmacists to practice disease state
26	management and any other pharmacy services determined by the board to require
27	a credential.
28	(2)(A) The credentials may be issued by agencies approved by the
29	board to pharmacists who qualify pursuant to as a result of meeting the
30	minimum competencies, standards, objectives, and qualifications determined by
31	the board.
32	(B) However, a credential shall not authorize the
33	pharmacist to practice credentialed pharmacy service services in Arkansas
34	until after the board has determined that the credentialed pharmacist meets
35	the minimum competencies, standards, objectives, and qualifications
36	determined by the board.

(b) The board shall adopt rules necessary and appropriate to implement
the credentialing and the board's approval of pharmacists to practice disease
state management and other credentialed pharmacy services, including:

- (1) Identification of areas of credentialed pharmacy services;
- (2) Identification of the minimum competencies, standards, objectives, and qualifications necessary for a credential and the board's approval to practice in each area of credentialed pharmacy service;
- (3) Identification of the standards for qualifying an agency to issue credentials for areas of pharmacy services;
- (4) The procedure and standards, which may include a practical examination, for the board's review and approval of a credential and determination of a pharmacist's qualifications to practice disease state management or other credentialed pharmacy service services;
- (5) The conversion of a credential previously issued by the board for the practice of disease state management or other pharmacy service services to a credential issued by an approved credentialing agency; and
- (6) Continuing professional education and other measures to maintain pharmacists' continuing competency in disease state management and other credentialed pharmacy services.

- SECTION 6. Arkansas Code § 17-92-412 is amended to read as follows: 17-92-412. Nursing home consultant permit pharmacist.
- (a)(1) The Arkansas State Board of Pharmacy shall provide for the issuance of nursing home consultant permits by rule.
- (2) The A nursing home consultant pharmacist—in-charge and the nursing home administrator shall be jointly responsible to ensure that a valid permit pharmacist license is posted at the facility at all times.
- (b) The board Arkansas State Board of Pharmacy shall set by rule the standards by which the controlled and legend drugs and devices will be maintained in the nursing home or long-term care facility.
- (c) The consultant pharmacist-in-charge pharmacist, in conjunction with the nursing home administrator and director of nurses, shall ensure the proper control and accountability, storage, and proper utilization of drugs and other legend devices dispensed to patients residing in the facility according to board standards as well as those standards established by state and federal guidelines

  APPROVED: 2/4/21

### Stricken language would be deleted from and underlined language would be added to present law. Act 406 of the Regular Session

1	State of Arkansas		H2/25/21 H3/8/21	
2	93rd General Assembly	A	Bill	
3	Regular Session, 2021			HOUSE BILL 1134
4				
5	By: Representative Boyd			
6	By: Senator Gilmore			
7				
8		For An Act	To Be Entitled	
9	AN ACT T	O AMEND THE DEFIN	ITION OF "PRACTICE OF	
10	PHARMACY	" TO ALLOW PHARMA	CISTS TO PRESCRIBE,	
11	ADMINIST	ER, DELIVER, DIST	RIBUTE, OR DISPENSE	
12	VACCINES	, IMMUNIZATIONS, A	AND MEDICATIONS TO TREAT	Г
13	ADVERSE	REACTIONS TO ADMIN	VISTERED VACCINES OR	
14	IMMUNIZA'	TIONS TO A PERSON	WHO IS THREE (3) YEARS	OF
15	AGE OR O	LDER; AND FOR OTHE	ER PURPOSES.	
16				
17				
18		Su	btitle	
19	TO	ALLOW PHARMACISTS	TO PRESCRIBE,	
20	ADM	INISTER, DELIVER,	DISTRIBUTE, OR	
21	DIS	PENSE VACCINES, I	MMUNIZATIONS, AND	
22	MED	ICATIONS TO TREAT	ADVERSE REACTIONS TO	
23	ADM	INISTERED VACCINE	5.	
24				
25				
26	BE IT ENACTED BY THE	GENERAL ASSEMBLY	OF THE STATE OF ARKANSA	۵:
27				
28	SECTION 1. Ark	kansas Code § 17-9	2-101(17)(A)(i), concer	ning the
29	definition of "practi	ice of pharmacy",	is amended to read as f	follows:
30		(i)(a) Dispen	sing, selling, distribu	ting,
31	transferring possessi	lon of, vending, b	artering, or, in accord	ance with rules
32	adopted by the Arkans	sas State Board of	Pharmacy, administerin	g drugs,
33	medicines, poisons, o	or chemicals that	under the laws of the U	nited States or
34	the State of Arkansas	may be sold or d	ispensed only on the pr	escription and
35	order of a practition	er authorized by	law to prescribe drugs,	medicines,
36	poisons, or chemicals			



1	(b) Except as limited by rules adopted by the
2	Arkansas State Board of Pharmacy, a pharmacist has the ability to administer
3	medications.
4	(c) Influenza vaccines and influenza
5	immunizations may be administered to a person seven (7) years of age or older
6	under a general written protocol.
7	(d) Vaccines and immunizations other than
8	influenza vaccines and influenza immunizations may be administered to a
9	person from seven (7) years of age to eighteen (18) years of age under a
10	general written protocol and subject to reporting required under § 20-15-1203
11	if written consent of the parent or legal guardian of the minor is obtained
12	before the administration of the vaccine or immunization.
13	(e) Vaccines and immunizations other than
14	influenza vaccines and influenza immunizations may be administered to a
15	person eighteen (18) years of age or older under a general written protocol.
16	(f) Medications other than vaccines and
17	immunizations may be administered to a person seven (7) years of age or older
18	under a patient-specific order or prescription and subject to reporting of
19	the administration to the prescribing physician.
20	(c) A pharmacist may prescribe, administer,
21	deliver, distribute, or dispense medications that treat adverse reactions
22	associated with the administration of vaccines and immunizations, vaccines,
23	and immunizations to or for a person three (3) years of age or older.
24	(d) A pharmacist who prescribes and
25	administers vaccines and immunizations other than for influenza or
26	coronavirus 2019 (COVID-19) to a person who is three (3) years of age to six
27	(6) years of age shall:
28	(1) Participate in the federal Vaccines
29	for Children Program; and
30	(2) Inform the person who is three (3)
31	years of age to six (6) years of age and adult caregivers accompanying the
32	person who is three (3) years of age to six (6) years of age of the
33	importance of a well-child visit with a pediatrician or other licensed
34	primary care provider and recommend a well-child visit at least yearly.
35	(e)(1) A pharmacist may administer medications
36	other than medications that treat adverse reactions associated with the

1	administration of vaccines and immunizations, vaccines, and immunizations
2	under a patient-specific order or prescription or general written protocol.
3	(2) The administration of the medication
4	under subdivision $(17)(A)(i)(e)(l)$ of this section is subject to reporting to
5	the prescribing physician, if applicable.
6	$\frac{(g)(f)}{f}$ A general written protocol under
7	subdivisions (17)( $\Lambda$ )(i)(c) and (e) of this section and patient-specific
8	orders or prescriptions under subdivisions (17)(A)(i)(d) and (f) subdivision
9	(17)(A)(i)(e) of this section shall be from a physician licensed by the
10	Arkansas State Medical Board and practicing in Arkansas or within fifty (50)
11	miles of the Arkansas border.
12	$\frac{h}{g}$ Under a statewide protocol, a
13	pharmacist may initiate therapy and administer or dispense, or both, drugs
14	that include Naloxone and nicotine replacement therapy products;
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16	/s/Boyd
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19	APPROVED: 3/22/21
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# Stricken language would be deleted from and underlined language would be added to present law. Act 407 of the Regular Session

1	State of Arkansas As Engrossed: H1/19/21 H3/8/21
2	93rd General Assembly A B1II
3	Regular Session, 2021 HOUSE BILL 1135
4	
5	By: Representatives Boyd, Pilkington
6	By: Senator Gilmore
7	
8	For An Act To Be Entitled
9	AN ACT TO AUTHORIZE PHARMACY TECHNICIANS TO
10	ADMINISTER VACCINES AND IMMUNIZATIONS; TO AMEND THE
11	DEFINITION OF "PRACTICE OF PHARMACY" TO ALLOW
12	PHARMACY TECHNICIANS TO ADMINISTER VACCINES OR
13	IMMUNIZATIONS TO A PERSON WHO IS THREE (3) YEARS OF
14	AGE OR OLDER; AND FOR OTHER PURPOSES.
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17	Subtitle
18	TO AUTHORIZE PHARMACY TECHNICIANS TO
19	ADMINISTER VACCINES AND IMMUNIZATIONS.
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22	BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF ARKANSAS:
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24	SECTION 1. Arkansas Code § 17-92-101(17)(C), concerning the
25	definition of "practice of pharmacy" relating to pharmacists and pharmacies,
26	is amended to read as follows:
27	(C)(i) The Arkansas State Board of Pharmacy may permit
28	pharmacy technicians other than pharmacists or interns to perform some or all
29	of those functions described in Arkansas State Board of Pharmacy rules under
30	the direct, personal supervision of a licensed pharmacist pursuant to under
31 32	rules defining the minimum qualifications of such employees, the ratio of
33	pharmacy technicians to supervising pharmacists, and the scope of the duties,
	practices, and procedures that the Arkansas State Board of Pharmacy
34 35	determines will promote the delivery of competent, professional
	pharmaceutical services and promote the public health and welfare. Nothing in
36	this chapter shall be construed as allowing pharmacy technicians to

1	administer medications.
2	(ii) A pharmacy technician may administer vaccines
3	and immunizations to a person three (3) years of age or older if delegated to
4	do so by a supervising pharmacist, but may not administer other medications.
5	(iii) The conduct of a pharmacy technician is the
6	responsibility of the pharmacist-in-charge and supervising pharmacist of the
7	pharmacy who shall not permit the employee to perform any act, task, or
8	function that involves the exercise of independent judgment by the employee.
9	(iii)(iv) Pharmacy products prepared by pharmacy
10	technicians shall be verified for accuracy by the supervising pharmacist
11	before release for patient use, and the verification shall be documented.
12	(iv)(v) The use of pharmacy technicians in a manner
13	not authorized by this chapter or rules promulgated hereunder shall be
14	unprofessional conduct by the pharmacist-in-charge and the supervising
15	pharmacist.
16	(v)(vi) It is recognized that hospital pharmacy
17	technicians as defined in § 17-92-602(4) are governed by the Hospital
18	Pharmacies Act, § 17-92-601 et seq., and related Arkansas State Board of
19	Pharmacy rules developed pursuant to that subchapter under the Hospital
20	Pharmacies Act, § 17-92-601 et seq.;
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22	/s/Boyd
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25	APPROVED: BECAME LAW ON 3/22/21 WITHOUT THE GOVERNOR'S SIGNATURE.
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