

CHAPTER 5 ARTICLE IX PRESCRIBING CONTROLLED SUBSTANCES

BUREAU (15)
LEGISLATIVE RESEARCH

Section 1- Arkansas optometrist licensed as optometric physician who applies for and possess a DEA number shall:

- A. Prescribe schedules III, IV, and V controlled substances only.
- B. Administer and prescribe controlled substances for the diagnosis and treatment of diseases and conditions of the eye, lids, and adnexa.
- C. Not sell any prescription medication including controlled substances.
- D. Be responsible for knowing and abiding by all state and federal regulations pertaining to controlled substances with emphasis on the "Mid-Level Practitioner's Manual", published by the DEA, and all State Board rules and regulations pertaining to controlled substances. Record the names and directions of prescribed controlled substances in the patient's record.
- E. A prescriber who has been found by the Arkansas State Board of Optometry to be in violation of a rule or law involving prescription drugs shall be required by the board to register with the Prescription Drug Monitoring Program and access patient information before writing a prescription for an opioid. The board, in its discretion, may remove this requirement after a period of time if the board deems removal of the requirement appropriate.
- F. Within the first two (2) years of being granted a license in the state, a prescriber shall obtain a minimum of three (3) hours of prescribing education approved by the Arkansas State Board of Optometry. The education approved by the board under this section shall include:
 - 1. Options for online and in-person programs; and
 - 2. Information on prescribing rules, regulations, and laws that apply to individuals who are licensed in the state.
 - 3. Information and instructions on prescribing controlled substances, record keeping and maintaining safe and professional boundaries.

This section-shall apply to all-prescribers licensed after December 31, 2015.

- G. A prescriber shall check the information in the Prescription Drug Monitoring Program when prescribing:
- 1. An opioid from Schedule II through Schedule III for every time prescribing the medication to a patient; and
 - 2. A benzodiazepine medication for the first time prescribing the medication to a patient.
- H. A practitioner who fails to access the Prescription Drug Monitoring Program as required is subject to disciplinary action by the Board.

Section 2-

- A. Only optometrists certified as optometric physicians, and/or approved by the Board, shall apply for or possess a DEA number to prescribe controlled substances.
- B. Optometrists not specifically approved by the Board to prescribe controlled substances:
 - 1. Cannot apply for, obtain or possess a DEA number
 - 2. Cannot prescribe controlled substances without being in violation of State and Federal laws.

Stricken language would be deleted from and underlined language would be added to present law. Act 820 of the Regular Session

1	State of Arkansas As Engrossed: S2/20/17 S3/9/17 S3/13/17 S3/14/17 S3/15/17
2	H3/17/17
3	91st General Assembly A Bill
4	Regular Session, 2017 SENATE BILL 339
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6	By: Senator J. Hutchinson
7	By: Representative Hammer
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9	For An Act To Be Entitled
10	AN ACT TO AMEND THE PRESCRIPTION DRUG MONITORING
11	PROGRAM TO MANDATE PRESCRIBERS CHECK THE PRESCRIPTION
12	DRUG MONITORING PROGRAM WHEN PRESCRIBING CERTAIN
13	MEDICATIONS; AND FOR OTHER PURPOSES.
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16	Subtitle
17	TO AMEND THE PRESCRIPTION DRUG MONITORING
18	PROGRAM TO MANDATE PRESCRIBERS CHECK THE
19	PRESCRIPTION DRUG MONITORING PROGRAM WHEN
20	PRESCRIBING CERTAIN MEDICATIONS.
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22	DE THE DIVISION DIVINING CONTROL ACCOUNTY OF THE CONTROL OF THE CO
23	BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF ARKANSAS:
24 25	CECTION 1 Adheren Cala C 20 7 (0//1) and and the manifestation
26	SECTION 1. Arkansas Code § 20-7-604(d), concerning the requirements for the Prescription Drug Monitoring Program, is amended to read as follows:
27	(d)(1) Practitioners Except as required in subdivision (d)(2) of this
28	section, practitioners are encouraged to access or check the information in
29	the controlled substance database created under this subchapter before
30	prescribing, dispensing, or administering medications.
31	(2)(A) A prescriber shall check the information in the
32	Prescription Drug Monitoring Program when prescribing:
33	(i) An opioid from Schedule II or Schedule III for
34	every time prescribing the medication to a patient; and
35	(ii) A benzodiazepine medication for the first time
36	prescribing the medication to a patient.



1 (B) A licensing board the	at licenses practitioners who have
2 the authority to prescribe shall adopt rule	s requiring the practitioners to
3 check the information in the Prescription D	rug Monitoring Program as
described in subdivision (d)(2) of this sec	tion.
(C) This subdivision (d)	(2) does not apply to:
(i) A practitioner	administering a controlled
substance:	
(a) Immediate	ely before or during surgery;
(b) During re	ecovery from a surrery while in a
healthcare facility;	
(c) In a hea.	lthcare facility; or
(d) Necessar	y to treat a patient in an
emergency situation at the scene of an emer	gency, in a licensed ground
ambulance or air ambulance, or in the inten-	sive care unit of a licensed
hospital;	
(ii) A practitione.	r prescribing or administering a
controlled substance to:	
(a) A pallia	tive care or hospice patient; or
(b) A residen	nt in a licensed nursing home
facility; or	
(iii) Situations is	n which the Prescription Drug
Monitoring Program is not accessible due to	technological or electrical
failure.	
(D) The State Board of H	ealth may amend, by rule, the
exemptions listed in subdivision $(d)(2)(C)$	of this section upon a
recommendation from the Director of the Dep	artment of Health and a showing
that the exemption or lack of exemption is	unnecessarily burdensome or has
created a hardship.	
(3) A licensed oncologist shall	1 check the Prescription Drug
Monitoring Program when prescribing to a pa	tient on an initial malignate
episodic diagnosis and every three (3) mont.	hs following the diagnosis while
continuing treatment.	
SECTION 2. Arkansas Code § 20-7-607(a)(1), concerning providing
prescription monitoring information to the	Prescription Drug Monitoring
Program, is amended to read as follows:	27

1	(a)(l)(A) $\underline{(i)}$ The Department of Health may shall review the
2	Prescription Drug Monitoring Program information, including without
3	limitation a review to identify information that appears to indicate whether
4	a person may be is obtaining prescriptions in a manner that may represent
5	misuse or abuse of controlled substances based on prescribing criteria
6	determined by the Director of the Department of Health upon consultation with
7	the Prescription Drug Monitoring Program Advisory Committee.
8	(ii) The prescribing criteria shall be posted on the
9	website of the department and be available in print upon request.
10	(B) If the information appears to indicate misuse or abuse
11	may have occurred, the department shall notify the practitioners and
12	dispensers who have prescribed or dispensed in the following manner:
13	(i) The department shall provide quarterly reports
14	to the individual practitioners and dispensers; and
15	(ii) If after twelve (12) months of providing
16	quarterly reports to the practitioners and dispensers, the information
17	appears to indicate misuse or abuse may be continuing, the department shall
18	send a report to the licensing boards of the practitioner or dispenser who
19	prescribed or dispensed the prescription.
20	(C) If information of misuse or abuse is identified, the
21	department shall notify the practitioners and dispensers who prescribed or
22	dispensed the prescriptions and the Office of Diversion Control of the United
23	States Drug Enforcement Administration.
24	(D) On or before January 1, 2019, the department shall
25	contract with a vendor to make the Prescription Drug Monitoring Program
26	interactive and to provide same-day reporting in real-time, if funding and
27	technology are available.
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29	SECTION 3. Arkansas Code § 20-7-611, concerning unlawful acts and
30	penalties regarding the Prescription Drug Monitoring Program, is amended to
31	add an additional subsection to read as follows:
32	(i) A practitioner who purposely fails to access the Prescription Drug
33	Monitoring Program as required by \$ 20-7-604(d) is subject to disciplinary
34	action by the licensing board of the practitioner.
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SECTION 4. Arkansas Code § 20-7-605(c), concerning the membership of

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1	the Prescription Drug Monitoring Program Advisory Committee, is amended to
2	read as follows:
3	(c) The committee shall consist of:
4	(1) One (1) representative designated by each of the following
5	organizations:
6	(A) The Arkansas Academy of Physician Assistants;
7	(B) The Arkansas Association of Chiefs of Police;
8	(C) The Arkansas Drug Director;
9	(D) The Arkansas Medical Society;
10	(E) The Arkansas Nurses Association;
11	(F) The Arkansas Optometric Association;
12	(G) The Arkansas Osteopathic Medical Association;
13	(H) The Arkansas Pharmacists Association;
14	(I) The Arkansas Podiatric Medical Association;
15	(J) The Arkansas Prosecuting Attorneys Association;
16	(K) The Arkansas Sheriffs' Association;
17	(L) The Arkansas State Dental Association;
18	(M) The Arkansas Veterinary Medical Association;
19	(N) The State Board of Health; and
20	(0) The Arkansas Public Defender Commission;
21	(2) One (1) mental health provider or certified drug and alcohol
22	counselor; and
23	(3) One (1) consumer appointed by the Governor;
24	(4) The chair of the Arkansas State Medical Board or his or her
25	designee who is also a member of the Arkansas State Medical Board; and
26	(5) The chair of the Arkansas State Board of Dental Examiners of
27	his or her designee who is also a member of the Arkansas State Board of
28 29	Dental Examiners.
30	SECTION 5. Arkansas Code § 17-95-303, concerning the powers and duties
31	of the Arkansas State Medical Board, is amended to add an additional
32	subdivision to read as follows:
33	(11) Promulgate rules limiting the amount of Schedule II
34	narcotics that may be prescribed and dispensed by licensees of the board.
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36	SECTION 6. Arkansas Code § 10-3-309(c), concerning the review and

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     approval of proposed state agency rules by the Legislative Council, is
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     amended to read as follows:
           (c)(l) A state agency shall file a proposed rule with the Legislative
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     Council at least thirty (30) days before the expiration of the period for
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    public comment on the rule under the Arkansas Administrative Procedure Act, §
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     25-15-201 et seq., or other laws or policies pertaining to the rulemaking
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     authority of that state agency.
                 (2) The Legislative Council shall assign proposed rules to the
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    Administrative Rules and Regulations Subcommittee of the Legislative Council.
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                 (3)(A)(i) The proposed rule shall be reviewed by the
    Administrative Rules and Regulations Subcommittee of the Legislative Council.
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                                    When reviewing a rule under subdivision
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                             (ii)
     (c)(3)(A)(i) of this section, the Administrative Rules and Regulations
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     Subcommittee of the Legislative Council shall allow members of the public a
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     reasonable opportunity to comment on the proposed rule.
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                       (B)(i)(a) Except as set forth in subdivision (c)(3)(B)(ii)
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     of this subsection. Upon upon conclusion of the review of the proposed rule
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     by the Administrative Rules and Regulations Subcommittee of the Legislative
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     Council, the proposed rule shall be considered approved unless a majority of
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     a quorum present request that the Administrative Rules and Regulations
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     Subcommittee of the Legislative Council vote on the issue of approving the
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     proposed rule.
                             (ii) (b) If the Administrative Rules and Regulations
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     Subcommittee of the Legislative Council votes on the issue of approving the
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     proposed rule, the proposed rule shall be approved unless a majority of a
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     quorum present vote for the proposed rule to not be approved.
                             (ii) A proposed rule submitted by the State Board of
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     Health under Arkansas Code § 20-7-604(d)(2)(D), concerning exemptions from
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     the requirements of the Prescription Drug Monitoring Program, shall be
     considered reviewed and approved by the subcommittee upon an affirmative vote
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     of three-fourths (3/4) of the members present when a quorum is present.
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                 (4)(A)(i) Except as set forth in subdivision (c)(4)(B) of this
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     subsection. A a proposed rule approved by the Administrative Rules and
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     Regulations Subcommittee of the Legislative Council shall be considered
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     approved by the Legislative Council unless a majority of a quorum present
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     request that the Legislative Council vote on the issue of approving the
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1	proposed rule.
2	(B)(ii) If the Legislative Council votes on the issue of
3	approving the proposed rule, the proposed rule shall be approved unless a
4	majority of a quorum present vote for the proposed rule to not be approved.
5	(B) A proposed rule submitted by the State Board of Health
6	under Arkansas Code § 20-7-604(d)(2)(D), concerning exemptions from the
7	requirements of the Prescription Drug Monitoring Program, shall be considered
8	reviewed and approved by the Legislative Council upon an affirmative vote of
9	three-fourths (3/4) of the members present when a quorum is present.
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12	SECTION 7. Arkansas Code § 10-3-309(f), concerning a vote not to
13	approve a state agency rule, is amended to read as follows:
14	(f)(l) A committee or subcommittee under this section may vote to not
15	approve a rule under this section only if the rule is inconsistent with:
16	(A) State or federal law; or
17	(B) Legislative intent.
18	(2) A committee or subcommittee under this section voting not to
19	approve a rule under this section shall state the grounds under subdivision
20	(f)(l) of this section when not approving a rule.
21	(3) A committee or subcommittee under this section considering a
22	rule submitted in accordance with Arkansas Code § 20-7-604(d)(2)(D),
23	concerning exemptions from the Prescription Drug Monitoring Program, is not
24	required to state the grounds required under subdivision (f)(1) when not
25	approving a rule.
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27	SECTION 8. Arkansas Code § 17-82-208, concerning the rules and
28	regulations of the Arkansas State Board of Dental Examiners, is amended to
29	add an additional subsection to read as follows:
30	(e) The board shall promulgate rules limiting the amount of Schedule
31	II narcotics that may be prescribed and dispensed by licensees of the board.
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33	SECTION 9. Arkansas Code § 17-87-203, concerning the powers and duties
34	of the Arkansas State Board of Nursing, is amended to add an additional
35	subdivision to read as follows:
36	(21) Promulgate rules limiting the amount of Schedule II

1	narcotics that may be prescribed and dispensed by licensees of the board.
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3	SECTION 10. Arkansas Code § 17-90-204, concerning the powers and
4	duties of the State Board of Optometry, is amended to add an additional
5	subdivision to read as follows:
6	(8) Promulgate rules limiting the amount of Schedule II
7	narcotics that may be prescribed and dispensed by licensees of the board.
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9	SECTION 11. Arkansas Code § 17-92-205, concerning the rules and
10	regulations of the Arkansas State Board of Pharmacy, is amended to add an
11	additional subsection to read as follows:
12	(d) The board shall promulgate rules limiting the amount of Schedule
13	II narcotics that may be dispensed by licensees of the board.
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15	SECTION 12. Arkansas Code § 17-101-203, concerning the powers and
16	duties of the Veterinary Medical Examining Board, is amended to add an
17	additional subdivision to read as follows:
18	(12) Promulgate rules limiting the amount of Schedule II
19	narcotics that may be prescribed and dispensed by licensees of the board.
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21	/s/J. Hutchinson
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24	APPROVED: 04/03/2017
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