

1	INTERIM STUDY PROPOSAL 2019-	-174
2	State of Arkansas	
3	92nd General Assembly A Bill	JMB/JMB
4	Second Extraordinary Session, 2020	HOUSE BILL
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6	By: Representative Love	
7	Filed with: House Committee or	n Public Health, Welfare, and Labor
8		pursuant to A.C.A. §10-3-217.
9	For An Act To Be Entitled	l
10	AN ACT TO AUTHORIZE PHARMACISTS TO DISPE	INSE HIV
11	PREEXPOSURE AND POSTEXPOSURE PROPHYLAXIS	; AND FOR
12	OTHER PURPOSES.	
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15	Subtitle	
16	TO AUTHORIZE PHARMACISTS TO DISPENSE HIV	
17	PREEXPOSURE AND POSTEXPOSURE PROPHYLAXIS.	
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20	BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE C	OF ARKANSAS:
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22	SECTION 1. Arkansas Code § 17-92-101(17)(A)(i	)(h), concerning the
23	definition of the "practice of pharmacy", is amended	l to read as follows:
24	(h) Under a statewide	e protocol, a pharmacist
25	may initiate therapy and administer or dispense, or	both, drugs that include
26	Naloxone <u>, and</u> nicotine replacement therapy products,	HIV preexposure
27	prophylaxis, and HIV postexposure prophylaxis;	
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29	SECTION 2. Arkansas Code § 17-92-101, concern	ing definitions regarding
30	pharmacy and pharmacists, is amended to add an addit	ional subdivision to read
31	as follows:	
32	(26) "HIV" means the human immunodeficiency virus or any other	
33	identified causative agent of acquired immunodeficie	ency syndrome (AIDS);
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35	SECTION 2. Arkansas Code § 17-92-115 is amend	led to read as follows:

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1 17-92-115. Requirements for administering and dispensing under 2 statewide protocol. 3 (a) When initiating therapy and administering or dispensing, or both, 4 under a statewide protocol, a pharmacist shall: 5 (1) Notify the primary care provider of the patient of any drug 6 or device furnished to the patient or enter the appropriate information in a 7 patient record system shared with the primary care provider, as permitted by 8 the primary care provider; 9 (2) Provide the patient with a written record of the drugs or 10 devices furnished and advise the patient to consult a physician of the patient's choice, if the patient does not have a primary care provider; and 11 12 (3)(A) Make a standardized fact sheet available to the recipient 13 of the drug or device. 14 (B) The standardized fact sheet shall include without 15 limitation: 16 (i) The indications and contraindications for the 17 use of the drug or device; 18 (ii) The appropriate method for the use of the drug 19 or device; 20 (iii) The need for medical follow-up; and 21 (iv) Other appropriate information. 22 (b)(1) In addition to the requirements under subsection (a) of this 23 section, when initiating therapy and administering or dispensing, or both, for HIV preexposure prophylaxis or HIV postexposure prophylaxis, or both, 24 under a statewide protocol, a pharmacist shall: 25 26 (A) Within twelve (12) months of initiating therapy and 27 administering or dispensing, or both, complete a training program approved by the Arkansas State Board of Pharmacy on the use of HIV preexposure 28 29 prophylaxis and HIV postexposure prophylaxis, which shall include information 30 about: 31 (i) Financial assistance programs for HIV 32 preexposure prophylaxis and HIV postexposure prophylaxis; and 33 (ii) Relevant federal guidelines regarding HIV preexposure prophylaxis and HIV postexposure prophylaxis; and 34 35 (B) Not permit a patient to waive consultation for HIV 36 preexposure prophylaxis or HIV postexposure prophylaxis.

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1	(2) Under a statewide protocol, a pharmacist shall dispense at	
2	least a thirty-day supply and up to a sixty-day supply of HIV preexposure	
3	prophylaxis if:	
4	(A)(i) The patient is HIV negative as documented by a	
5	negative HIV test result obtained within the previous seven (7) days from:	
6	(a) An HIV antigen/antibody test;	
7	(b) An HIV antibody-only test; or	
8	(c) A rapid, point-of-care fingerstick blood	
9	test approved by the United States Food and Drug Administration.	
10	(ii) If the test results are not transmitted	
11	directly to the pharmacist, the pharmacist shall verify the test results.	
12	(iii) If the patient tests positive for HIV	
13	infection, the pharmacist shall direct the patient to a primary care provider	
14	and provide a list of providers and clinics in the region;	
15	(B) The patient does not report:	
16	(i) Any signs or symptoms of acute HIV infection on	
17	a self-reported checklist of acute HIV infection signs and symptoms; and	
18	(ii) Usage of any contraindicated medication;	
19	(C) The pharmacist provides counseling to the patient on	
20	the ongoing use of HIV preexposure prophylaxis, which shall include education	
21	about:	
22	(i) Side effects;	
23	(ii) Safety during pregnancy and breastfeeding;	
24	(iii) Adherence to recommended dosing;	
25	(iv) The importance of timely testing and treatment	
26	for HIV, renal function, hepatitis B, hepatitis C, sexually transmitted	
27	diseases, and pregnancy for individuals of childbearing capacity; and	
28	(v) The requirement that subsequent prescriptions	
29	for HIV preexposure prophylaxis be issued by a primary care provider; and	
30	(D) To the extent possible, the pharmacist documents the	
31	services provided by the pharmacist in the record system.	
32	(3) Under a statewide protocol, a pharmacist shall dispense a	
33	course of HIV postexposure prophylaxis if the pharmacist:	
33 34	<u>course of HIV postexposure prophylaxis if the pharmacist:</u> (A) Screens the patient and determines the exposure to HIV	

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1	(B) Provides HIV testing or determines the patient is:	
2	(i) Willing to undergo HIV testing consistent with	
3	federal guidelines; or	
4	(ii) Unwilling to undergo HIV testing but otherwise	
5	eligible for HIV postexposure prophylaxis;	
6	(C) Provides counseling to the patient on the ongoing use	
7	of HIV postexposure prophylaxis, which shall include education about:	
8	(i) Side effects;	
9	(ii) Safety during pregnancy and breastfeeding;	
10	(iii) Adherence to recommended dosing;	
11	(iv) The importance of timely testing and treatment	
12	for HIV, renal function, hepatitis B, hepatitis C, sexually transmitted	
13	diseases, and pregnancy for individuals of childbearing capacity; and	
14	(v) The availability of HIV preexposure prophylaxis	
15	for a person who is at a substantial risk of acquiring HIV; and	
16	(D) To the extent possible, documents the services	
17	provided by the pharmacist in the record system.	
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19	SECTION 3. Arkansas Code § 23-92-506(b)(6) and (7), concerning	
20	prohibited practices of a pharmacy benefits manager, are amended to read as	
21	follows:	
22	(6) Make or permit any reduction of payment for pharmacist	
23	services by a pharmacy benefits manager or a healthcare insurer directly or	
24	indirectly to a pharmacy under a reconciliation process to an effective rate	
25	of reimbursement, including without limitation generic effective rates, brand	
26	effective rates, direct and indirect remuneration fees, or any other	
27	reduction or aggregate reduction of payment; <del>or</del>	
28	(7)(A) Prohibit a pharmacist from dispensing HIV preexposure	
29	prophylaxis or HIV postexposure prophylaxis under a state protocol.	
30	(B) As used in subdivision (b)(7) of this section, "HIV"	
31	means the human immunodeficiency virus or any other identified causative	
32	agent of acquired immunodeficiency syndrome (AIDS); or	
33	(7)(8) Do any combination of the actions listed in subdivisions	
34	(b)(1)-(6)(b)(1)-(7) of this section.	
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1	SECTION 4. Arkansas Code Title 23, Chapter 99, Subchapter 11, is	
2	amended to add an additional section to read as follows:	
3	23-99-1120. HIV preexposure prophylaxis and HIV postexposure	
4	prophylaxis.	
5	(a) As used in this section:	
6	(1) "AIDS" means acquired immunodeficiency syndrome; and	
7	(2) "HIV" means the human immunodeficiency virus or any other	
8	identified causative agent of acquired immunodeficiency syndrome.	
9	(b) Except as provided in subsection (c) of this section, a health	
10	benefit plan or healthcare insurer shall not require prior authorization or	
11	step therapy for antiretroviral drugs that are medically necessary for the	
12	prevention of HIV or AIDS, including HIV preexposure prophylaxis and HIV	
13	postexposure prophylaxis.	
14	(c) If the United States Food and Drug Administration approves one (1)	
15	or more therapeutic equivalents of a drug, device, or product for the	
16	prevention of HIV or AIDS, a health benefit plan or healthcare insurer is not	
17	required to cover all therapeutically equivalent versions without prior	
18	authorization or step therapy if at least one (1) therapeutically equivalent	
19	version is covered without prior authorization or step therapy.	
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22	Referred by Representative Love	
23	Prepared by: JMB/JMB	
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