1	INTERIM STUDY PROPOSAL 2019-174	
2	State of Arkansas	
3	92nd General Assembly A Bill JMB/JM	В
4	Second Extraordinary Session, 2020 HOUSE BIL	L
5		
6	By: Representative Love	
7	Filed with: House Committee on Public Health, Welfare, and Lab)01
8	pursuant to A.C.A. §10-3-21	17
9	For An Act To Be Entitled	
10	AN ACT TO AUTHORIZE PHARMACISTS TO DISPENSE 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 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 to read as follows:	
24	(h) Under a statewide protocol, a pharmacist	
25	may initiate therapy and administer or dispense, or both, drugs that include	1
26	Naloxone, and nicotine replacement therapy products, HIV preexposure	
27	prophylaxis, and HIV postexposure prophylaxis;	
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29	SECTION 2. Arkansas Code § 17-92-101, concerning definitions regardin	Ŭ
30	pharmacy and pharmacists, is amended to add an additional subdivision to rea	.d
31	as follows:	
32	(26) "HIV" means the human immunodeficiency virus or any other	
33	identified causative agent of acquired immunodeficiency syndrome (AIDS);	
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35	SECTION 2. Arkansas Code § 17-92-115 is amended to read as follows:	

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
11	patient's choice, if the patient does not have a primary care provider; and
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,
24	for HIV preexposure prophylaxis or HIV postexposure prophylaxis, or both,
25	under a statewide protocol, a pharmacist shall:
26	(A) Within twelve (12) months of initiating therapy and
27	administering or dispensing, or both, complete a training program approved by
28	the Arkansas State Board of Pharmacy on the use of HIV preexposure
29	prophylaxis and HIV postexposure prophylaxis, which shall include information
30 31	about:
32	(i) Financial assistance programs for HIV preexposure prophylaxis and HIV postexposure prophylaxis; and
33	
34	(ii) Relevant federal guidelines regarding HIV preexposure prophylaxis and HIV postexposure prophylaxis; and
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	<pre>prophylaxis if:</pre>
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:
34	(A) Screens the patient and determines the exposure to HIV
35	occurred within the previous seventy-two (72) hours and the patient otherwise
36	meets the clinical criteria for HIV postexposure prophylaxis;

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; or
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	$\frac{(b)(1)-(6)}{(b)(1)-(7)}$ 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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