DEPARTMENT OF HEALTH, CENTER FOR HEALTH PROTECTION

SUBJECT: Immunization Reporting

<u>DESCRIPTION</u>: Act 541 of 2015 removed the adult consent requirement before immunization providers report an adult immunization to the statewide immunization information system (IIS). This change brings the IIS rules into alignment with the State's Health Information Exchange's (SHARE) consent requirements and eliminates the administrative burden on immunization providers to gain and document consent prior to reporting.

PUBLIC COMMENT: A public hearing was held on September 3, 2015. The Public Comment period expired on September 3, 2015. The Department did not receive any public comments.

The proposed effective date is February 10, 2016.

CONTROVERSY: This is not expected to be controversial.

FINANCIAL IMPACT: There is no financial impact. Prior rules permit immunization providers to report immunizations administered to adults but only after gaining consent. This rule will eliminate the administrative burden on immunization providers to collect and document consent prior to reporting. This change also aligns these rules with those of the State's Health Information Exchange (SHARE). There is no additional cost to the state, county, or municipal government, s the electronic reporting system already exists and is functional for capturing adult immunizations.

LEGAL AUTHORIZATION:

THIS RULE ENACTS ACT 541 OF 2015. The Act removed the adult consent requirement before immunization reports are sent to the statewide immunization information system.

Ark. Code Ann. § 20-15-1202 requires the Department to establish an immunization registry and the authority to adopt the rules to implement the registry.



Arkansas Department of Health

4815 West Markham Street ◆ Little Rock, Arkansas 72205-3867 ◆ Telephone (501) 661-2000

Governor Asa Hutchinson

Nathaniel Smith, MD, MPH, Director and State Health Officer

Summary of Rules and Regulations Pertaining to Immunization Reporting

It is proposed to revise the Rules and Regulations Pertaining to Immunization Reporting pursuant to the procedures of the Administrative Procedures Act process, as amended, by the authority of Act 434 of 1967 as amended. These requirements apply to reporting information to the statewide immunization registry under the authority of Ark. Code. Ann §§20-15-1201 – 1203.

Act 541 of 2015 removed the adult consent requirement before immunization providers report an adult immunization to the statewide immunization information system (IIS). This change brings the IIS Rules and Regs into alignment with the State's Health Information Exchange's (SHARE) consent requirements and eliminates the administrative burden on immunization providers to gain and document consent prior to reporting.

QUESTIONNAIRE FOR FILING PROPOSED RULES AND REGULATIONS WITH THE ARKANSAS LEGISLATIVE COUNCIL AND JOINT INTERIM COMMITTEE

DEPARTMENT/AGENCY	Arkansas Department of Health				
DIVISION	Center for Health Protection - Immunization and Communicable Disease Section				
DIVISION DIRECTOR	Renee Mallory				
CONTACT PERSON	Hilda Dupwe				
ADDRESS	4815 West Markham, MS48, Little Rock, AR 72205				
PHONE NO. 501-661-24 NAME OF PRESENTER A MEETING					
PRESENTER E-MAIL rol	pert.brech@arkansas.gov				
	INSTRUCTIONS				
 A. Please make copies of this form for future use. B. Please answer each question completely using layman terms. You may use additional sheets, if necessary. C. If you have a method of indexing your rules, please give the proposed citation after "Short Title of this Rule" below. D. Submit two (2) copies of this questionnaire and financial impact statement attached to the front of two (2) copies of the proposed rule and required documents. Mail or deliver to: Donna K. Davis Administrative Rules Review Section Arkansas Legislative Council Bureau of Legislative Research One Capitol Mall, 5th Floor Little Rock, AR 72201 					
•	~~·~~·~				
 What is the short title of th rule? 	nis Rules and Regulations Pertaining to Immunization Reporting				
2. What is the subject of the prule?	proposed Enhance the Utilization of Immunization Registry				
	aply with a federal statute, rule, or regulation? Yes \(\square\) No \(\square\) ederal rule, regulation, and/or statute citation.				
4. Was this rule filed under the Procedure Act? If yes, what is the effective rule?	he emergency provisions of the Administrative Yes \(\sum \) No \(\sum \) e date of the emergency				
When does the emergency expire?	rule				

	Will this emergency rule be promulgated under the permanent provisions of the Administrative Procedure Act?	Yes 🗌	No 🛚
5.	Is this a new rule? Yes No No If yes, please provide a brief summary explaining the regulation.		
	Does this repeal an existing rule? Yes No No No If yes, a copy of the repealed rule is to be included with your completed replaced with a new rule, please provide a summary of the rule giving a does.	questionnaire n explanation	e. If it is being of what the rule
rul	Is this an amendment to an existing e? Yes No No If yes, please attach a mark-up showing the changes in the existing rule changes. Note: The summary should explain what the amendment should be clearly labeled "mark-up."	and a summa	ry of the substantive mark-up copy
6.	Cite the state law that grants the authority for this proposed rule? If code citation. <u>Arkansas Code 20-15-1201 - 1203</u>	ified, please g	ive the Arkansas
rep adı	What is the purpose of this proposed rule? Why is it necessary? To creat porting adult immunizations to the registry consistent with Act 541 of 201 ministrative burden of gaining and documenting consent before reporting gistry. To align consent requirements with those set by the State's Health	5. To elimin adult immun	ate the izations given to the
8.	Please provide the address where this rule is publicly accessible in electrequired by Arkansas Code § 25-19-108(b). www.healthy.arkansas.gov	ronic form via	a the Internet as
9.	Will a public hearing be held on this proposed rule? Yes No If yes, please complete the following: Date: September 3, 2015 Time: 11:00 a.m. Arkansas Department of Health, Room Place: 2508	!	
	. When does the public comment period expire for permanent promulgati 00 a.m. on September 3, 2015	on? (Must pro	ovide a date.)
	. What is the proposed effective date of this proposed rule? (Must provide bruary 10, 2015	e a date.)	70.0
	Do you expect this rule to be controversial? Yes No If yes, please explain.		

FINANCIAL IMPACT STATEMENT

PLEASE ANSWER ALL QUESTIONS COMPLETELY

DEPARTMENT		IMENT	Arkansas Depa	artment of Heal	th			
DF	VISIC	N	Immunization	and Communic	able Disea	se Section		
PE	RSON	N COMPLE	TING THIS ST	FATEMENT	Hilda Dup	owe		
TE	LEPH	HONE NO.	501-661-2493	_ FAX NO. <u>501</u>	1-661-2300	EMAIL: Hile	da.Dupwe@	Arkansas.gov
To Sta	comp ateme	oly with Ark. nt and file tw	Code Ann. § 25 to copies with the	5-15-204(e), plone questionnaire	ease comple and propo	ete the followin osed rules.	g Financial I	mpact
SF	IORT	TITLE OF	THIS RULE	Rules and Re	gulations F	Pertaining to Imi	munization F	Reporting
1.	Does	this propos	ed, amended, or	repealed rule h	ave a finan	icial impact?	Yes 🗌	No 🔀
2.	Is the rule based on the best reasonably obtainable scientific, technical, economic, or other evidence and information available concerning the need for, consequences of, and alternatives to the rule? Yes No						No 🗌	
3. In consideration of the alternatives to this rule, v by the agency to be the least costly rule consider				e determined	Yes 🔀	No 🗌		
	If an	agency is pr	roposing a more	costly rule, ple	ase state th	e following:		
	(a) How the additional benefits of the more costly rule justify its additional cost;							
	(b)	The reason	for adoption of	the more costly	rule;			
	(c)	Whether the more costly rule is based on the interests of public health, safety, or welfare, and if so, please explain; and;						
	(d)	(d) Whether the reason is within the scope of the agency's statutory authority; and if so, please explain.				so, please		
4.	If the	purpose of the	his rule is to imp	lement a federal	rule or regi	ılation, please sta	ate the follow	ing:
	(a) What is the cost to implement the federal rule or regulation?							
	<u>Cur</u>	rent Fiscal	<u>Year</u>		<u>Ne</u>	xt Fiscal Year		
Federal Funds Cash Funds					Fed Cas Spe	neral Revenue leral Funds sh Funds scial Revenue ler (Identify)		

	Total	0	Total	0	
	(b) What is the ad	lditional cost of the state rule?			
	Current Fiscal Year		Next Fiscal Year		
	General Revenue Federal Funds Cash Funds Special Revenue		Federal Funds Cash Funds Special Revenue		
	Other (Identity)		Other (Identity)	Acres 1	
	Total	0	Total	0	
5.	What is the total es the proposed, amer explain how they a	stimated cost by fiscal year to anded, or repealed rule? Identify re affected.	y private individual, entit the entity(ies) subject to t	y and business subject to he proposed rule and	
<u>C</u> 1	urrent Fiscal Year		Next Fiscal Ye	<u>ear</u>	
	NA		\$ <u>NA</u>		
bu im	t only after gaining munization provide	tions permit immunization provi consent. This proposed rule will ers to collect and document cons- with those of the State's Health I	ll eliminate the administra ent prior to reporting. Thi	ative burden on s change also aligns these	
6.	What is the total e implement this rul affected.	estimated cost by fiscal year to st le? Is this the cost of the program	ate, county, and municipa m or grant? Please explai	al government to n how the government is	
	urrent Fiscal Year		Next Fiscal Ye	e <u>ar</u>	
No	NA o additional cost is a pturing adult immu	anticipated as the electronic reponizations.	\$ <u>NA</u> rting system already exis	ts and is functional for	
				-	
7.	With respect to the agency's answers to Questions #5 and #6 above, is there a new or increased cost or obligation of at least one hundred thousand dollars (\$100,000) per year to a private individual, private entity, private business, state government, county government, municipal government, or to two (2) or more of those entities combined?				
			Yes ☐ No ⊠		
If YES, the agency is required by Ark. Code Ann. § 25-15-204(e)(4) to file written findings time of filing the financial impact statement. The written findings shall be filed simultaneous with the financial impact statement and shall include, without limitation, the following:				filed simultaneously	
	(1) a statement of	the rule's basis and purpose;			
	(2) the problem the a rule is requir	e agency seeks to address with the	he proposed rule, includir	ng a statement of whether	

- (3) a description of the factual evidence that:
 - (a) justifies the agency's need for the proposed rule; and
 - (b) describes how the benefits of the rule meet the relevant statutory objectives and justify the rule's costs;
- (4) a list of less costly alternatives to the proposed rule and the reasons why the alternatives do not adequately address the problem to be solved by the proposed rule;
- (5) a list of alternatives to the proposed rule that were suggested as a result of public comment and the reasons why the alternatives do not adequately address the problem to be solved by the proposed rule;
- (6) a statement of whether existing rules have created or contributed to the problem the agency seeks to address with the proposed rule and, if existing rules have created or contributed to the problem, an explanation of why amendment or repeal of the rule creating or contributing to the problem is not a sufficient response; and
- (7) an agency plan for review of the rule no less than every ten (10) years to determine whether, based upon the evidence, there remains a need for the rule including, without limitation, whether:
 - (a) the rule is achieving the statutory objectives;
 - (b) the benefits of the rule continue to justify its costs; and
 - (c) the rule can be amended or repealed to reduce costs while continuing to achieve the statutory objectives.