EXHIBIT H

DEPARTMENT OF HUMAN SERVICES, MEDICAL SERVICES

SUBJECT: Pharmacy Manual #1-17 and Section 1 1-17

<u>DESCRIPTION</u>: CMS published the Covered Outpatient Drug final rule (CMS-2345-FC) on February 1, 2016 pertaining to reimbursement for covered outpatient drugs in the Medicaid program. It outlines key changes that states need to address when determining their reimbursement methodology for ingredient costs based on actual acquisition cost (AAC) plus a professional dispensing fee among other things.

PUBLIC COMMENT: No public hearing was held. The public comment period expired on May 15, 2017. The department received no comments.

This rule was filed and approved as an emergency rule on April 1, 2017. The emergency rule expires on July 29, 2017. The proposed effective date for the permanent rule is July 1, 2017.

FINANCIAL IMPACT: There is no financial impact.

LEGAL AUTHORIZATION: This rule is required in order to ensure compliance with CMS-2345-FC and 81 FR 5170. The Department of Human Services is authorized to "make rules and regulations and take actions as are necessary or desirable to carry out the provisions of this chapter [Public Assistance] and that are not inconsistent therewith." Arkansas Code Annotated § 20-76-201 (12). Arkansas Code § 20-77-107 specifically authorizes the department to "establish and maintain an indigent medical care program."



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

DEPARTMENT/AGENCY	Department of Human Services					
DIVISION	Division of Medical Services					
DIVISION DIRECTOR	Dawn Stehle					
CONTACT PERSON ADDRESS	Thomas Herndon PO Box 1437, Slot S295, Little Rock, AR 72203					
		thomas.hemdon@				
PHONE NO. 501-396-60		IL dhs.arkansas.gov				
NAME OF PRESENTER AT						
PRESENTER E-MAIL <u>ta</u>	ami.harlan@dhs.arkansas.gov					
	INSTRUCTIONS					
necessary. C. If you have a method of it of this Rule" below. D. Submit two (2) copies of the proof two (2) copies of tw	tion completely using layman terms. You may indexing your rules, please give the proposed ce this questionnaire and financial impact statem roposed rule and required documents. Mail or vis we Rules Review Section gislative Council gislative Research Mall, 5th Floor AR 72201	itation after "Short Title ent attached to the front r deliver to: ***********************************				
1. What is the short title of the	his rule? Pharmacy Manual #1-17 and Section	1 1-17				
2. What is the subject of the	proposed rule? Pharmacy Update					
3. Is this rule required to cor	mply with a federal statute, rule, or regulation?	Yes ⊠ No ☐ CMS-2345-FC; 81 FR				
If yes, please provide the	If yes, please provide the federal rule, regulation, and/or statute citation.					
Procedure Act?	the emergency provisions of the Administrative	Yes No 🖂				
ii yes, what is the effective	we date of the emergency rule? April 1, 2017					
When does the emergenc	y rule expire? July 29, 2017					

	the Administrative Procedure Act? Yes No Yes
5.	Is this a new rule? Yes No No No If yes, please provide a brief summary explaining the regulation.
	Does this repeal an existing rule? Yes No No If yes, a copy of the repealed rule is to be included with your completed questionnaire. If it is being replaced with a new rule, please provide a summary of the rule giving an explanation of what the rule does.
	Is this an amendment to an existing rule? Yes No No If yes, please attach a mark-up showing the changes in the existing rule and a summary of the substantive changes. Note: The summary should explain what the amendment does, and the mark-up copy should be clearly labeled "mark-up."
6.	Cite the state law that grants the authority for this proposed rule? If codified, please give the Arkansas Code citation. <u>Arkansas Statute 20-76-201</u>
ou the	What is the purpose of this proposed rule? Why is it necessary? CMS published the Covered atpatient Drug final rule (CMS-2345-FC) (81 FR 5170) on 02/01/2016, pertaining to reimbursement for covered to the things in the Medicaid program. It outlines key changes that States need to address when determining the reimbursement methodology for ingredient costs based on actual acquisition cost (AAC) plus a professional spensing fee among other things.
	Please provide the address where this rule is publicly accessible in electronic form via the Internet as required by Arkansas Code § 25-19-108(b). https://www.medicaid.state.ar.us/general/comment/comment.aspx
111	aps., www.medicaid.state.ar.us/general/comment/comment.aspx
9.	Will a public hearing be held on this proposed rule? Yes ☐ No ☒
	If yes, please complete the following:
	Date:
	Time:
	Place:
	O. When does the public comment period expire for permanent promulgation? (Must provide a date.) May 15, 2017
2	
	1. What is the proposed effective date of this proposed rule? (Must provide a date.)
J	uly 1, 2017 (Adopted by Federal Regulation April 1, 2017)
12	2. Do you expect this rule to be controversial? Yes \(\subseteq \text{No } \text{\infty}

If yes, please explain.	3.99.d				
13. Please give the names Please provide their po	of persons, groups, osition (for or again	or organi st) if knov	zations that you vn.	expect to c	omment on these rules?

FINANCIAL IMPACT STATEMENT

PLEASE ANSWER ALL QUESTIONS COMPLETELY

DEPARTMENT		MENT	Department of Human Services						
DIVISION		N .	Division of Medical Services						
PEF	RSON	COMPLE	TING THIS ST	TATEMENT Ly	nn Burton		648		
TEI	LEPH	ONE NO.	501-682-1857	FAX NO. 501-4	04-4619 EMAIL: lynn.l	ourton@dhs.	arkansas.gov		
To Sta	comp	ly with Ark it and file tv	. Code Ann. § 2 wo copies with t	.5-15-204(e), plea	se complete the following and proposed rules.				
1.	Does	this propos	ed, amended, o	r repealed rule ha	ve a financial impact?	Yes 🗌	No 🖂		
2.	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.				ves to this rule, wastly rule considered	as this rule determined d?	Yes 🖂	No 🗌		
	If an	agency is p	roposing a mor	e costly rule, plea	se state the following:				
	(a)	How the a	dditional benefi	ts of the more cos	stly rule justify its additio	nal cost;			
	(b)	The reason	ı for adoption o	f 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;						welfare, and		
	(d) Whether the reason is within the scope of the agency's statutory authority; and if so, please explain.						f so, please		
4.	If th	e purpose o	this rule is to in	nplement a federal	rule or regulation, please s	state the follo	wing:		
	(a)	What is the	ne cost to imple	ment the federal r	ule or regulation?				
	Cu	rrent Fisc	ıl Year		Next Fiscal Year				
	Fee Ca Sp	neral Rever deral Funds sh Funds ecial Rever her (Identif	\$0 nue		General Revenue Federal Funds Cash Funds Special Revenue Other (Identify)	\$0 \$0			

Total	\$0	Total	\$0
(b) What is the	ne additional cost of the st	ate rule?	
Current Fis	cal Year	Next Fiscal Year	
General Reve Federal Fund Cash Funds Special Reve Other (Ident	enue	Cash Funds	
Total	-	Total	3
proposed, am they are affect	ended, or repealed rule? eted.	al year to any private individual, entit Identify the entity(ies) subject to the p	proposed rule and explain how
Current Fiscal	<u>Year</u>	Next Fisca	Year
\$	<u> </u>	\$	
		cal year to state, county, and municipor grant? Please explain how the gove	
Current Fiscal	Year	Next Fisca	l Year
\$ 0		\$ 0	
Methodology.	ance/savings was included This is just the change to additional impact.	l in State Plan Amendment 2016-003 the manuals reflecting that change in	- Pharmacy Pricing methodology. Changing the
or obligatio private enti	n of at least one hundred	to Questions #5 and #6 above, is therethousand dollars (\$100,000) per year government, county government, murbined?	to a private individual,
		Yes No 🛛	
time of filir	ng the financial impact sta	k. Code Ann. § 25-15-204(e)(4) to file tement. The written findings shall be and shall include, without limitation, t	filed simultaneously
(1) a statem	nent of the rule's basis and	d purpose;	
	blem the agency seeks to a s required by statute;	address with the proposed rule, includ	ing 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.

Summary for Pharmacy 1-17 & Section 1 1-17

CMS published the Covered Outpatient Drug final rule (CMS-2345-FC) (81 FR 5170) on 02/01/2016, pertaining to reimbursement for covered outpatient drugs in the Medicaid program. It outlines key changes that States need to address when determining their reimbursement methodology for ingredient costs based on actual acquisition cost (AAC) plus a professional dispensing fee among other things.