Stricken language would be deleted from and underlined language would be added to present law. Act 722 of the Regular Session

1	State of Arkansas	As Engrossed: H3/17/17	
2	91st General Assembly	A Bill	
3	Regular Session, 2017		SENATE BILL 361
4			
5	By: Senator Flippo		
6			
7	For An Act To Be Entitled		
8	AN ACT TO CREATE AN EXEMPTION FROM THE LAWS REGARDING		
9	THE PRACTICE OF PHARMACY FOR DIALYSATE OR DEVICES		
10	NECESSARY FOR HOME PERITONEAL KIDNEY DIALYSIS IN		
11	CERTAIN S	SITUATIONS; AND FOR OTHER PURPOSES.	
12			
13			
14		Subtitle	
15	TO (	CREATE AN EXEMPTION FROM THE LAWS	
16	REGA	ARDING THE PRACTICE OF PHARMACY FOR	
17	DIALYSATE OR DEVICES NECESSARY FOR HOME		
18	PERITONEAL KIDNEY DIALYSIS IN CERTAIN		
19	SIT	UATIONS.	
20			
21			
22	BE IT ENACTED BY THE	GENERAL ASSEMBLY OF THE STATE OF ARKAN	NSAS:
23			
24	SECTION 1. Ark	ansas Code Title 17, Chapter 92, Subcl	napter l, is
25	amended to add an add	litional section to read as follows:	
26	<u>17-92-115. Exe</u>	emption for home peritoneal kidney dia	<u>lysis.</u>
27	<u>(a) The provis</u>	sions of §§ 17-92-101, 17-92-103, 17-92	<u>2-105, 17-92-205,</u>
28	<u>17-92-206, 17-92-303, 17-92-401, 17-92-402, 17-92-404, 17-92-405, 17-92-409,</u>		
29	17-92-410, 17-92-411, and 17-92-902 do not apply to the sale or distribution		
30	of dialysate or devices necessary to perform home peritoneal kidney dialysis		
31	to patients with end stage renal disease if:		
32	(1) The dialysate composed of dextrose or icodextrin or devices		
33	are:		
34	<u>(A)</u>	Approved or cleared by the United St	tates Food and Drug
35	Administration as required by federal law;		
36	<u>(B)</u>	Lawfully held by a manufacturer or a	<u>a third-party</u>



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As Engrossed: H3/17/17

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1	logistics provider of the manufacturer that is properly registered with the
2	Arkansas State Board of Pharmacy as a wholesale distributor or medical device
3	provider;
4	(C) Held and delivered in original, sealed packaging from
5	the manufacturing facility; and
6	(D) Delivered only by the manufacturer or a third-party
7	logistics provider of the manufacturer and only upon receipt of a physician's
8	order by a licensed pharmacy and the transmittal of an order from a licensed
9	pharmacy to the manufacturer or a third party logistics provider of the
10	manufacturer; and
11	(2) The manufacturer or a third-party logistics provider of the
12	manufacturer delivers the dialysate or devices directly to:
13	(A) A patient with end stage renal disease or a designee
14	for the self-administration of the dialysis therapy; or
15	(B) A healthcare provider or institution for
16	administration or delivery of the dialysis therapy to a patient with end
17	stage renal disease.
18	(b)(1) The board shall retain oversight of all other drugs for home
19	peritoneal kidney dialysis with the exception of dialysate as described in
20	subdivision (a)(l) of this section.
21	(2) All records of sales and distribution of dialysate to
22	patients under this section shall be retained according to state law and rule
23	of the board.
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25	/s/Flippo
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28	APPROVED: 03/28/2017
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