GENERAL ASSEMBLY OF NORTH CAROLINA SESSION 2015

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SENATE DRS35085-MGqq-62 (03/03)

Short Title:	Create Permit Exemptions/Home Renal Products.	(Public)		
Sponsors:	Senator Hise (Primary Sponsor).			
Referred to:				
	A BILL TO BE ENTITLED			
AN ACT AMENDING THE PHARMACY PRACTICE ACT TO CREATE A PHARMACY				
PERMIT EXEMPTION AND A DEVICE AND MEDICAL EQUIPMENT PERMIT				
EXEMPTION FOR THE DISPENSING AND DELIVERY OF HOME RENAL				
PRODUC	CTS.			
The General Assembly of North Carolina enacts:				
	ECTION 1. Article 4A of Chapter 90 of the General Statutes is ame	nded by		
_	section to read:			
	2. Pharmacy permit exemption for dispensing and delivery of hon	<u>ne renal</u>		
	oducts.			
	ation or facility within or outside this State from which dialysate of			
	perform home renal dialysis are dispensed and delivered to a patient in this			
-	the pharmacy permit requirements established by G.S. 90-85.21, provided	1 that all		
_	criteria are met:	4 T d		
<u>(1</u>)	The dialysate or drugs have been approved or cleared by United Sta and Drug Administration.	tes Food		
<u>(2</u>)		nt of the		
(2	manufacturer that is properly registered with the Board as a manufacturer			
	a wholesaler, or both, as required by the Board.	cturer or		
<u>(3</u>)	- · · · · · · · · · · · · · · · · · · ·	original.		
75	sealed packaging from the manufacturing facility.	<u></u>		
<u>(4</u>		agent of		
<u> </u>	the manufacturer and only upon receipt of a physician's order.			
<u>(5</u>		lysate or		
	drugs directly to either or both of the following:			
	a. A patient with chronic kidney failure or a designee of the pa	tient, for		
	self-administration of the dialysis therapy.			
	<u>b.</u> <u>A health care provider, or health care facility licensed under</u>	Chapter		
	122C, 131D, or 131E of the General Statutes, for administration			
	delivery of the dialysis therapy to a patient with chronic	<u>kidney</u>		
~-	failure.			
SECTION 2. G.S. 90-85.22 reads as rewritten:				
"§ 90-85.22. Device and medical equipment permits; exemptions.				

This section shall not apply to either any of the following:

Gen	General Assembly of North Carolina Session 201				
	(1)	A pharmaceutical manufacturer registered with the Food and Drug			
		Administration.			
	(2)	A wholly owned subsidiary of a pharmaceutical manufacturer registered			
		with the Food and Drug Administration.			
	<u>(3)</u>	The dispensing and delivery of home renal products in accordance with the			
		criteria specified in G.S. 90-85.21C."			
	SEC.	FION 3. This act becomes affective October 1, 2015			