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# A BILL FOR AN ACT

RELATING TO THE PRACTICE OF PHARMACY.

**BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF HAWAII:**

1           SECTION 1. Chapter 461, Hawaii Revised Statutes, is  
2 amended by adding a new section to be appropriately designated  
3 and to read as follows:

4           "§461-       Distribution of dialysate drugs and devices.

5           (a) The license, registration, and permit requirements of this  
6 chapter shall not apply to a manufacturer, wholesale  
7 distributor, or third-party logistics provider, to the extent  
8 the manufacturer, wholesale distributor, or third-party  
9 logistics provider is engaged in the distribution of dialysate  
10 drugs or devices necessary to perform home dialysis on patients  
11 with end-stage renal disease; provided that the following  
12 criteria are met:

13           (1) The dialysate drugs or devices are approved by the  
14           United States Food and Drug Administration, as  
15           required by federal law;

16           (2) The dialysate drugs or devices are lawfully held by a  
17           manufacturer or a manufacturer's agent that is



1           properly registered with the board as a manufacturer  
2           or wholesale distributor;  
3       (3) The dialysate drugs or devices are held and delivered  
4           in their original, sealed, and labeled packaging from  
5           the manufacturing facility;  
6       (4) The dialysate drugs or devices are delivered only by  
7           the manufacturer or the manufacturer's agent and only  
8           upon receipt of a physician's order; and  
9       (5) The manufacturer or the manufacturer's agent delivers  
10           the dialysate drugs or devices directly to a:  
11           (A) Patient with end stage renal disease, or the  
12           patient's designee, for the patient's self-  
13           administration of dialysis therapy; or  
14           (B) Health care provider or institution for  
15           administration or delivery of dialysis therapy to  
16           a patient with end stage renal disease.  
17       (b) For the purposes of this section:  
18       "Manufacturer" shall have the same meaning as in section  
19       328-112.  
20       "Third-party logistics provider" means an entity that  
21       provides or coordinates warehousing or other logistics services



1 on behalf of a manufacturer, wholesale distributor, or dispenser  
2 of a product.

3 "Wholesale distributor" shall have the same meaning as in  
4 section 328-112."

5 SECTION 2. New statutory material is underscored.

6 SECTION 3. This Act shall take effect on June 30, 3000.



**Report Title:**

Practice of Pharmacy; Dialysate Drugs or Devices; Manufacturers;  
Wholesalers; Third-Party Logistics Providers; Exemption

**Description:**

Exempts manufacturers, wholesale distributors, and third-party logistics providers of home dialysate drugs or devices from the license, registration, and permit requirements for pharmacies, under certain conditions. Effective 6/30/3000. (HD1)

*The summary description of legislation appearing on this page is for informational purposes only and is not legislation or evidence of legislative intent.*

