

Chapter 44

2019 Regular Session

Subject Pharmacy Licensure and Peritoneal Dialysis

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Overview

This act exempts from pharmacy licensure requirements manufacturers and other entities that distribute dialysate or devices to perform home peritoneal dialysis on patients with end-stage renal disease, if specified criteria are met.

Summary

Section Description

1 Pharmacy licensure requirements.

Amends § 151.19, subd. 1. Provides an exemption from pharmacy licensure requirements for a drug manufacturer, wholesale drug distributor, or a third-party logistics provider, that distributes dialysate or devices to perform home peritoneal dialysis on patients with end-stage renal disease, if:

- 1) the manufacturer or its agent leases or owns the facility from which the dialysate or devices will be delivered;
- 2) the dialysate is comprised of dextrose or icodextrin and has been approved by the Food and Drug Administration;
- 3) the dialysate is stored and delivered in original, sealed, and unopened packaging;
- 4) the dialysate or devices are delivered only upon receipt of a physician's order by a pharmacy, and the prescription is reviewed and processed by a pharmacist meeting specified criteria;
- 5) records and other information are maintained by the manufacturer for a minimum of three years and made available to the board upon request; and
- 6) the manufacturer or agent delivers the dialysate or devices directly to the patient or patient's designee, for self-administration of the dialysis therapy; or a health care provider or institution, for administration or delivery of dialysis therapy to the patient.



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