Ohio Revised Code

Section 4729.55 Terminal distributor license requirements.

Effective: October 3, 2023
Legislation: House Bill 33

No license shall be issued to an applicant for licensure as a terminal distributor of dangerous drugs unless the applicant has furnished satisfactory proof to the state board of pharmacy that:

(A) The applicant is equipped as to land, buildings, and equipment to properly carry on the business of a terminal distributor of dangerous drugs within the category of licensure approved by the board.

(B) A pharmacist, licensed health professional authorized to prescribe drugs, other person authorized by the board, animal shelter or county dog warden licensed under section 4729.531 of the Revised Code, or laboratory will maintain supervision and control over the possession and custody of dangerous drugs and controlled substances that may be acquired by or on behalf of the applicant.

(C) Adequate safeguards are assured to prevent the sale or other distribution of dangerous drugs by any person other than a pharmacist or licensed health professional authorized to prescribe drugs.

(D) Adequate safeguards are assured that the applicant will carry on the business of a terminal distributor of dangerous drugs in a manner that allows pharmacists and pharmacy interns employed by the terminal distributor to practice pharmacy in a safe and effective manner.

(E) If the applicant, or any agent or employee of the applicant, has been found guilty of violating section 4729.51 of the Revised Code, the "Federal Food, Drug, and Cosmetic Act," 52 Stat. 1040 (1938), 21 U.S.C.A. 301, the federal drug abuse control laws, Chapter 2925., 3715., 3719., or 4729. of the Revised Code, or any rule of the board, adequate safeguards are assured to prevent the recurrence of the violation.

(F) If the application is made on behalf of an animal shelter or county dog warden, at least one of the agents or employees of the animal shelter or county dog warden is certified in compliance with section 4729.532 of the Revised Code.

(G) In the case of an applicant who is a retail seller of peritoneal dialysis solutions in original packages labeled as required by the "Federal Food, Drug, and Cosmetic Act," 52 Stat. 1040 (1938), 21 U.S.C.A. 301, the applicant will maintain supervision and control over the possession, custody, and retail sale of the peritoneal dialysis solutions.

(H) In the case of an applicant who is a pain management clinic, the applicant meets the requirements to receive a license with a pain management clinic classification issued under section 4729.552 of the Revised Code.