Kansas Administrative Regulations

State Board of Pharmacy

Article 7.—Miscellaneous Provisions

68-7-14. Prescription labels.

(a) The label of each drug or device shall be typed or machine-printed and shall include the following information:

(1) The name, address, and telephone number of the pharmacy or registered facility dispensing the prescription;

(2) the name of the prescriber;

(3) the full name of the patient;

(4) the identification number assigned to the prescription by the dispensing pharmacy or registered facility;

(5) the date the prescription was filled or refilled, whichever is most recent;

(6) adequate directions for use of the drug or device;

(7) the beyond-use date of the drug or device dispensed. If the drug is removed from the original manufacturer’s packaging, the beyond-use date shall not exceed:

(A) one year from the date of dispensing;

(B) the manufacturer’s expiration date; or

(C) the maximum rating of the packaging materials,

whichever is less.

(8) the name of the drug or device dispensed;

(9) if a generic drug or device is dispensed, the name of the manufacturer or an easily identified abbreviation of the manufacturer's name;

(10) the strength of the drug;

(11) the contents in terms of weight, measure, or numerical count; and

(12) necessary auxiliary labels and storage instructions, if needed.

(b) A registered facility shall be permitted to label or relabel only those drugs or devices originally dispensed from the registered facility’s location.

(Authorized by K.S.A. 65-1630; implementing K.S.A. 2025 Supp. 65-1626a; effective, E-77-39, July 22, 1976; effective Feb. 15, 1977; amended May 1, 1978; amended May 1, 1980; amended May 1, 1988; amended June 6, 1994; amended March 20, 1995; amended April 28, 2000; amended Oct. 23, 2009; amended Aug. 14, 2026.)