

**68-7-14 Prescription labels.** (a) The label of each drug or device shall, at a minimum, be typed or machine-printed and shall include the following information:

(1) The name, address, and telephone number of the pharmacy or facility dispensing the prescription. If shared order filling is utilized, the label shall also include the name and address of the other pharmacy participating in the shared service agreement;

(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 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 one year from the date of dispensing, the time remaining on the manufacturer's expiration date, or the maximum rating of the packaging materials, whichever is less;

(8) either the generic name of the drug or device and its manufacturer or the brand name or corresponding generic name, whichever was dispensed of the drug or device;

(9) ~~the name of the manufacturer or distributor of the drug or device, or an easily identified abbreviation of the manufacturer's or distributor'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.~~

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

(Authorized by K.S.A. 65-1630; implementing K.S.A. 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 P-\_\_\_\_\_.)

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