3. If a unit of use drug distribution system is used, each dose of medication shall be individually packaged in a hermetically sealed, tamper-proof container, and shall carry full manufacturer's disclosure information on each discrete dose. Disclosure information shall include, but not be limited to, the following: product name and strength, lot number, expiration date, and manufacturer's or distributor's name.

(c) Single use and disposable items shall not be reused.

(d) No stock supply of prescription medications shall be maintained, unless prior approval is obtained from the Bureau.

(e) Discontinued or expired medications shall be destroyed within 30 days in the facility, or, if unopened and properly labeled, returned to the pharmacy. All medication destruction in the facility shall be witnessed and documented by two persons, each of whom shall be either the administrator, the registered nurse or the pharmacist.

New Rule, R.2000 d.202, effective May 15, 2000.
See: 32 N.J.R. 739(a), 32 N.J.R. 1763(b).

SUBCHAPTER 14. CARBON MONOXIDE ALARMS

5:27–14.1 Carbon monoxide alarms

(a) Carbon monoxide alarms shall be installed and maintained in full operating condition in the following locations:

1. Single station carbon monoxide alarms shall be installed and maintained in the immediate vicinity of every sleeping room in buildings that contain a fuel-burning appliance or that have an attached garage.

2. As an alternative to the requirements above, carbon monoxide alarms may be installed in the locations specified in the Uniform Construction Code (N.J.A.C. 5:23–3.20) with the approval of the Bureau.

(b) Carbon monoxide alarms shall be manufactured, listed and labeled in accordance with UL 2034 and shall be installed in accordance with the requirements of this subchapter and NFPA 720. Carbon monoxide alarms shall be battery-operated, hard-wired or of the plug-in type.