

1 10A NCAC 13F .1003 is amended with changes as published in 29:08 NCR, pp. 907-909 as follows:

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3 **10A NCAC 13F .1003 MEDICATION LABELS**

4 (a) Prescription legend medications dispensed by the pharmacy shall have a legible ~~{ printed }~~ label with the following
5 information:

- 6 (1) the name of the resident for whom the medication is prescribed;
- 7 (2) the most recent date of issuance;
- 8 (3) the name of the prescriber;
- 9 (4) the name and concentration of the medication, quantity dispensed, and prescription serial number;
- 10 (5) directions for use stated and not abbreviated;
- 11 (6) a statement of generic equivalency shall be indicated if a brand other than the brand prescribed is
12 dispensed;
- 13 (7) the expiration date, unless dispensed in a single unit or unit dose package that already has an
14 expiration date;
- 15 (8) auxiliary statements as required of the medication;
- 16 (9) the name, address and telephone number of the dispensing pharmacy; and
- 17 (10) the name or initials of the dispensing pharmacist.

18 (b) For medication systems ~~such as med-paks and multi-paks when in which~~ two or more prescribed solid oral dosage
19 forms are packaged and dispensed together, labeling shall be in accordance with Paragraph (a) of this Rule and the
20 label or package shall also have a physical description or identification of each medication contained in the package.

21 (c) The facility shall assure the container is relabeled by a licensed pharmacist or a dispensing practitioner at the
22 refilling of the medication when there is a change in the directions by the prescriber. The facility shall have a procedure
23 for identifying direction changes until the container is correctly labeled. No person other than a licensed pharmacist
24 or dispensing practitioner shall alter a prescription label.

25 (d) Non-prescription medications shall have the manufacturer's label with the expiration date visible, unless the
26 container has been labeled by a licensed pharmacist or a dispensing ~~practitioner.~~ practitioner in accordance with
27 Paragraph (a) of this Rule. Non-prescription medications in the original manufacturer's container shall be labeled with
28 at least the resident's name and the name shall not obstruct any of the information on the container. Facility staff may
29 label or write the resident's name on the container.

30 (e) Medications, prescription and non-prescription, shall not be transferred from one container to another except when
31 prepared for a resident's leave of absence or administration to a resident.

32 ~~(f) Prescription medications leaving the facility shall be in a form packaged and labeled by a licensed pharmacist or~~
33 ~~a dispensing practitioner. Non-prescription medications that are not packaged or labeled by a licensed pharmacist or~~
34 ~~dispensing practitioner must be released in the original container and directions for administration must be provided~~
35 ~~to the resident or responsible party. The facility shall assure documentation of medications, including quantity released~~
36 ~~and returned to the facility.~~

1 *History Note: Authority G.S. ~~131D-2~~ 131D-2.16; 131D-4.5; 143B-165;*
2 *Eff. July 1, ~~2005~~. 2005;*
3 *Amended Eff. April 1, 2015.*