1	10A NCAC 13I	F.1003 is proposed for amendment as follows:
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3	10A NCAC 13	F .1003 MEDICATION LABELS
4	(a) Prescription	legend medications shall have a legible printed label with the following information:
5	(1)	the name of the resident for whom the medication is prescribed;
6	(2)	the most recent date of issuance;
7	(3)	the name of the prescriber;
8	(4)	the name and concentration of the medication, quantity dispensed, and prescription serial number;
9	(5)	directions for use stated and not abbreviated;
10	(6)	a statement of generic equivalency shall be indicated if a brand other than the brand prescribed is
11		dispensed;
12	(7)	the expiration date, unless dispensed in a single unit or unit dose package that already has ar
13		expiration date;
14	(8)	auxiliary statements as required of the medication;
15	(9)	the name, address and telephone number of the dispensing pharmacy; and
16	(10)	the name or initials of the dispensing pharmacist.
17	(b) For medicat	ion systems such as med paks and multi-paks when in which two or more <u>prescribed</u> solid oral dosage
18	forms are packaged and dispensed together, labeling shall be in accordance with Paragraph (a) of this Rule and the	
19	label or package	e shall also have a physical description or identification of each medication contained in the package.
20	(c) The facility	shall assure the container is relabeled by a licensed pharmacist or a dispensing practitioner at the
21	refilling of the medication when there is a change in the directions by the prescriber. The facility shall have a procedure	
22	for identifying direction changes until the container is correctly labeled. No person other than a licensed pharmacis	
23	or dispensing pr	ractitioner shall alter a prescription label.
24	(d) Non-prescr	ription medications shall have the manufacturer's label with the expiration date visible, unless the
25	container has been labeled by a licensed pharmacist or a dispensing practitioner. practitioner in accordance with	
26	Paragraph (a) of this Rule. Non-prescription medications in the original manufacturer's container shall be labeled with	
27	at least the resident's name and the name shall not obstruct any of the information on the container. Facility staff may	
28	label or write the resident's name on the container.	
29	(e) Medications, prescription and non-prescription, shall not be transferred from one container to another except whe	
30	prepared for a resident's leave of absence or administration to a resident.	
31	(f) Prescription medications leaving the facility shall be in a form packaged and labeled by a licensed pharmacist of	
32	a dispensing pro	netitioner. Non-prescription medications that are not packaged or labeled by a licensed pharmacist or
33	dispensing practitioner must be released in the original container and directions for administration must be provide	
34	to the resident or responsible party. The facility shall assure documentation of medications, including quantity	

History Note: Authority G.S. <u>131D-2</u> <u>131D-2.16</u>; 131D-4.5; 143B-165;

released and returned to the facility.

- Eff. July 1, 2005. <u>2005</u>;
- 2 <u>Amended Eff. April 1, 2015.</u>

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