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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) Labeling of Prescription prescription legend medications medications, except for medications prepared for a 5 resident's leave of absence in accordance with Rule .1010(d)(4) of this Section, shall have a be legible {printed} label 6 and include with the following information: 7 the name of the resident for whom the medication is prescribed; (1)8 (2)the most recent date of issuance; 9 (3) the name of the prescriber; 10 (4)the name and concentration of the medication, quantity dispensed, and prescription serial number; 11 directions for use stated and not abbreviated; unabbreviated directions for use stated; (5) 12 a statement of generic equivalency shall be indicated if a brand other than the brand prescribed is (6) 13 dispensed; 14 (7) the expiration date, unless dispensed in a single unit or unit dose package that already has an 15 expiration date; 16 (8) auxiliary statements information as required of the medication; 17 (9) the name, address address, and telephone number of the dispensing pharmacy; and 18 (10)the name or initials of the dispensing pharmacist. 19 (b) For medication systems such as med paks and multi paks when in which two or more prescribed solid oral dosage 20 forms are packaged and dispensed together, labeling shall be in accordance with Paragraph (a) of this Rule and the 21 label or package shall also have a physical description or identification of each medication contained in the package. 22 (c) The facility shall assure any changes in directions of a resident's medication by the prescriber are on the container is relabeled by a licensed pharmacist or a dispensing practitioner at the refilling of the medication when there is a 23 24 change in the directions by the prescriber. by the pharmacist or dispensing practitioner. The facility shall have a 25 procedure for identifying direction changes until the container is correctly labeled. labeled in accordance with 26 Paragraph (a) of this Rule. No person other than a licensed pharmacist or dispensing practitioner shall alter a 27 prescription label. 28 (d) Non-prescription medications shall have the manufacturer's label with the expiration date visible, unless the 29 container has been labeled by a licensed pharmacist or a dispensing practitioner. practitioner in accordance with 30 Paragraph (a) of this Rule. Non-prescription medications in the original manufacturer's container shall be labeled with 31 at least the resident's name and the name shall not obstruct any of the information on the container. Facility staff may 32 label or write the resident's name on the container. 33 (e) Medications, prescription and non-prescription, shall not be transferred from one container to another except when 34 prepared for a resident's leave of absence or administration to a resident. 35 (f) Prescription medications leaving the facility shall be in a form packaged and labeled by a licensed pharmacist or 36 a dispensing practitioner. Non-prescription medications that are not packaged or labeled by a licensed pharmacist or

37 dispensing practitioner must be released in the original container and directions for administration must be provided

| 1 | to the resident or responsible party. The facility shall assure documentation of medications, including quantity released |
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| 2 | and returned to the facility.   |

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| 4 | History Note: | Authority G.S. <del>131D-2</del> <u>131D-2.16</u> ; 131D-4.5; 143B-165; |
| 5 |               | Eff. July 1, <del>2005</del> . <u>2005:</u>                             |
| 6 |               | <u>Amended Eff. April 1, 2015.</u>                                      |