Rules For: Family Care Homes Type of Rule: Permanent Amendment MCC Action: Final Adoption

9/17/14

10A NCAC 13G .1003 is amended as published in 29:08 NCR, pp. 907-909 as follows: 1 2 3 10A NCAC 13G .1003 **MEDICATION LABELS** 4 (a) Prescription legend medications shall have a legible printed label with the following information: the name of the resident for whom the medication is prescribed; 5 (1)(2) the most recent date of issuance; 6 7 (3) the name of the prescriber; the name and concentration of the medication, quantity dispensed, and prescription serial number; 8 (4) 9 (5) directions for use stated and not abbreviated; a statement of generic equivalency shall be indicated if a brand other than the brand prescribed is (6) 10 11 dispensed; the expiration date, unless dispensed in a single unit or unit dose package that already has an 12 **(7)** 13 expiration date; auxiliary statements as required of the medication; 14 (8) (9) the name, address and telephone number of the dispensing pharmacy; and 15 16 (10)the name or initials of the dispensing pharmacist. (b) For medication systems such as med paks and multi-paks when in which two or more prescribed solid oral dosage 17 forms are packaged and dispensed together, labeling shall be in accordance with Paragraph (a) of this Rule and the 18 label or package shall also have a physical description or identification of each medication contained in the package. 19 (c) The facility shall assure the container is relabeled by a licensed pharmacist or a dispensing practitioner at the 20 refilling of the medication when there is a change in the directions by the prescriber. The facility shall have a procedure 21 for identifying direction changes until the container is correctly labeled. No person other than a licensed pharmacist 22 or dispensing practitioner shall alter a prescription label. 23 24 (d) Non-prescription medications shall have the manufacturer's label with the expiration date visible, unless the container has been labeled by a licensed pharmacist or a dispensing practitioner. practitioner in accordance with 25 Paragraph (a) of this Rule. Non-prescription medications in the original manufacturer's container shall be labeled with 26 at least the resident's name and the name shall not obstruct any of the information on the container. Facility staff may 27 28 label or write the resident's name on the container. (e) Medications, prescription and non-prescription, shall not be transferred from one container to another except when 29 prepared for a resident's leave of absence or administration to a resident. 30 (f) Prescription medications leaving the facility shall be in a form packaged and labeled by a licensed pharmacist or a 31 dispensing practitioner. Non prescription medications that are not packaged or labeled by a licensed pharmacist or 32 dispensing practitioner must be released in the original container and directions for administration must be provided 33 to the resident or responsible party. The facility shall assure documentation of medications, including quantity 34

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released and returned to the facility.

1	Note: Dispensing of medications is restricted to pharmacists or other health care practitioners that are approved by				
2	the North Carolina Board of Pharmacy. Repackaging or providing more than one dose of a prescription medication				
3	including unit dose prescription medications, for subsequent administration is an act of dispensing.				
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5	History Note:	Authority G.S. 131D-2 <u>131D</u>) <u>-2.16</u> ; 131D-4.5;	143B-165; S.L. 1999-0334	
6		Temporary Adoption Eff. De	cember 1, 1999;		
7		Eff. July 1, 2000. <u>2000;</u>		1.12.12.2.2. 2012年至10月1日 - 11.12.12.12.12.12.12.12.12.12.12.12.12.1	The state of the s
Ω.		Amended Fff April 1 2015			

Rules For: Family Care Homes Type of Rule: Permanent Amendment MCC Action: Final Adoption

1/27/15

10A NCAC 13G .1010 is amended with changes as published in 29:08 NCR, pp. 907-909 as follows:

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10A NCAC 13G .1010 PHARMACEUTICAL SERVICES

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- 5 (a) A family care home shall allow the residents the right to choose a pharmacy provider as long as the pharmacy will provides services that are in compliance with the facility's medication management policies and procedures.
- 7 (b) There shall be a current, written agreement with a licensed pharmacist or a prescribing practitioner for pharmaceutical care services according to Rule .1009 of this Section. The written agreement shall include a statement
- 9 of the responsibility of each party.

during the absence:

- 10 (c) The facility shall assure the provision of pharmaceutical services to meet the needs of the residents including
 11 procedures that assure the accurate ordering, receiving and administering of all medications prescribed on a routine,
 12 emergency, or as needed basis.
- (d) The facility shall assure the provision of medication for residents on temporary leave from the facility or involved 13 in day activities out of the facility. (Medications prepared for a resident's temporary leave of absence shall be 14 packaged in a manner that facilitates safe administration and enables the resident or resident's responsible person to 15 identify the correct medication and correct administration time for each medication. The amount of medications 16 necessary to cover the duration of the resident's absence may be taken from the supply of medication already dispensed 17 to the resident and prepared by a medication aide, or licensed health professional with authority to administer or 18 dispense medications. The following information for each medication prepared for the resident's absence shall be 19 provided verbally and in writing to the resident or the person who is designated as the resident's responsible person 20
 - (1) the name and strength of the drug;
 - (2) the directions for administration as prescribed by the resident's physician; and
 - (3) any cautionary information from the original prescription package.
 - For medications removed from the resident's supply of medications, the name of the resident and the information provided in Subparagraphs (1) and (2) shall be provided directly on the container containing the medication. The facility shall maintain documentation of medications provided for the resident's leave of absence, including the quantity released from the facility, the quantity returned to the facility, and the name of the individual who prepared the medication for the resident's leave of absence.}
 - The facility shall have written policies and procedures for a resident's temporary leave of absence. The policies and procedures shall facilitate safe administration by assuring that upon receipt of the medication for a leave of absence the resident or resident's responsible person is able to identify the medication, dosage, and administration time for each medication provided for the temporary leave of absence. The policies and procedures shall include at least the following provisions:
 - (1) The amount of resident's medications provided shall be sufficient and necessary to cover the duration of the resident's absence. For the purposes of this Rule, sufficient and necessary means the provision of the amount of medication to be administered during the leave of absence or the

1		provision of only a current dose pack, card, or container if the current dose pack, card, or container			
2		has enough medication for the planned absence;			
3	<u>(2)</u>	Written and verbal instructions for each medication to be released for the resident's absence shall			
4		be provided to the resident or the person who is designated as the resident's responsible person			
5		during the absence and shall include at least:			
6		(A) the name and strength of the medication;			
7		(B) the directions for administration as prescribed by the resident's physician.			
8	,	(C) any cautionary information from the original prescription package if the			
9		information is not on the container released for the leave of absence; and			
10	<u>(3)</u>	Labeling of the resident's medication container for the leave of absence shall be legible and include			
11	at least the name of the resident and the name and strength of the medication.				
12	The facility shall maintain documentation of medications provided for the resident's leave of absence, including the				
13	quantity released from the facility and the quantity returned to the facility. The documentation of the quantities o				
14	medications released from and returned to the facility for a resident's leave of absence shall be verified by signature				
15	of the facility sta	ff and resident or resident's responsible person upon the medications' release from and return to the			
16	facility.				
17	(e) The facility shall assure that accurate records of the receipt, use and disposition of medications are maintained in				
18	the facility and readily available for review.				
19	(f) A facility with 12 or more beds shall have a written agreement with a pharmacy provider for dispensing services				
20	The written agreement shall include a statement of the responsibility of each party.				
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22	History Note:	Authority G.S. 131D-2 <u>131D-2.16</u> ; 131D-4.5; 143B-165;			
23		Eff. July 1, 2005. <u>2005;</u>			
24		Amended Eff. April 1, 2015.			