

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL011-287	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 05/18/2018
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NAME OF PROVIDER OR SUPPLIER DAWN FORREST HOME	STREET ADDRESS, CITY, STATE, ZIP CODE 29 GRANDVIEW CIRCLE ASHEVILLE, NC 28806
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(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETE DATE
V 000	<p>INITIAL COMMENTS</p> <p>An annual survey was completed on 5/18/18. A deficiency was cited.</p> <p>This facility is licensed for the following service category: 10A NCAC 27G .5600F Supervised Living for Individuals of all Disability Groups/Alternative Family Living.</p>	V 000		
V 118	<p>27G .0209 (C) Medication Requirements</p> <p>10A NCAC 27G .0209 MEDICATION REQUIREMENTS</p> <p>(c) Medication administration:</p> <p>(1) Prescription or non-prescription drugs shall only be administered to a client on the written order of a person authorized by law to prescribe drugs.</p> <p>(2) Medications shall be self-administered by clients only when authorized in writing by the client's physician.</p> <p>(3) Medications, including injections, shall be administered only by licensed persons, or by unlicensed persons trained by a registered nurse, pharmacist or other legally qualified person and privileged to prepare and administer medications.</p> <p>(4) A Medication Administration Record (MAR) of all drugs administered to each client must be kept current. Medications administered shall be recorded immediately after administration. The MAR is to include the following:</p> <p>(A) client's name;</p> <p>(B) name, strength, and quantity of the drug;</p> <p>(C) instructions for administering the drug;</p> <p>(D) date and time the drug is administered; and</p> <p>(E) name or initials of person administering the drug.</p> <p>(5) Client requests for medication changes or checks shall be recorded and kept with the MAR file followed up by appointment or consultation</p>	V 118		

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LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE _____ TITLE _____ (X6) DATE _____

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V 118	<p>Continued From page 1 with a physician.</p> <p>This Rule is not met as evidenced by: Based on record review and interviews, the facility failed to keep the MAR current and failed to follow the written order of a physician affecting 1 of 2 sampled clients (Client #1). The findings are:</p> <p>Record review on 5/18/18 for Client #1 revealed: -Admission date of 12/1/99 with diagnoses of Moderate Intellectual Disability and High Blood Pressure. -Physician ordered medications included: -Loratidine 10mg once daily for allergies. -Polyethylene Glycol 1 capful with water/juice twice daily for constipation. -Both Loratidine and Polyethylene Glycol were discontinued 9/22/17. Review on 5/18/18 of March-May 2018 MARs revealed: -Loratidine was initialed as administered at April 1-30. -Polyethylene Glycol was initialed as administered at April 1-30.</p> <p>Interview on 5/18/18 with AFL Caregiver revealed: -She typed and printed the MARs each month for her clients. -She must have pulled an old MAR forward to use for April. -Client #1 had not taken Loratidine or Polyethylene Glycol for some time and did not receive it in April despite her initials on the MAR. -She now uses the "dispill packs" which did not contain Loratadine.</p>	V 118		

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V 118	Continued From page 2 -She would pay better attention to her documentation.	V 118		