

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:  <b>MHL034168</b>	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____  B. WING _____	(X3) DATE SURVEY COMPLETED  <b>R</b> <b>09/21/2021</b>
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NAME OF PROVIDER OR SUPPLIER  <b>DAVIS HOUSE AT BETHABARA</b>	STREET ADDRESS, CITY, STATE, ZIP CODE <b>2020 CLYDE HAYES DRIVE</b> <b>WINSTON SALEM, NC 27106</b>
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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 and follow up survey was completed on 9/21/21. Deficiencies were cited.</p> <p>This facility is licensed for the following service category: 10A NCAC 27G. 5600C Supervised Living for Adults with Developmental Disabilities.</p>	V 000		
V 117	<p>27G .0209 (B) Medication Requirements</p> <p>10A NCAC 27G .0209 MEDICATION REQUIREMENTS</p> <p>(b) Medication packaging and labeling:</p> <p>(1) Non-prescription drug containers not dispensed by a pharmacist shall retain the manufacturer's label with expiration dates clearly visible;</p> <p>(2) Prescription medications, whether purchased or obtained as samples, shall be dispensed in tamper-resistant packaging that will minimize the risk of accidental ingestion by children. Such packaging includes plastic or glass bottles/vials with tamper-resistant caps, or in the case of unit-of-use packaged drugs, a zip-lock plastic bag may be adequate;</p> <p>(3) The packaging label of each prescription drug dispensed must include the following:</p> <p>(A) the client's name;</p> <p>(B) the prescriber's name;</p> <p>(C) the current dispensing date;</p> <p>(D) clear directions for self-administration;</p> <p>(E) the name, strength, quantity, and expiration date of the prescribed drug; and</p> <p>(F) the name, address, and phone number of the pharmacy or dispensing location (e.g., mh/dd/sa center), and the name of the dispensing practitioner.</p>	V 117		

Division of Health Service Regulation LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
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V 117	<p>Continued From page 1</p> <p>This Rule is not met as evidenced by: Based on observations, record reviews, and interviews, the facility failed to ensure prescription medications were dispensed with the prescriber's name, the current dispensing date, the name, strength, quantity and expiration date of the prescribed medication, the name, address and phone number of the pharmacy and the name of the dispensing practitioner for 1 of 6 clients (#5). The findings are:</p> <p>Review on 9/21/21 of client #5's Medication Administration Record (MAR) dated 9/1/21-9/21/21 revealed:</p> <ul style="list-style-type: none"> <li>- Sodium Citrate and Citric Acid Solution 500 mg/334 mg per 5 ML: Give 10 ML (milliliter) twice daily.</li> <li>- Carnitor SF (Sugar Free) Solution: Give 15 ML by mouth 3 times a day.</li> </ul> <p>Observations at approximately 10:57 am on 9/21/21 of client #5's medications revealed:</p> <ul style="list-style-type: none"> <li>- There were no labels on the bottles for: Sodium Citrate and Citric Acid Solution nor Carnitor SF.</li> </ul> <p>Interview on 9/21/21 with the Qualified Professional revealed:</p> <ul style="list-style-type: none"> <li>- Both Sodium Citrate and Citric Acid Solution, and Carnitor SF were delivered in large bags with the labels.</li> <li>- The large bags with labels had been thrown away.</li> </ul>	V 117		
V 118	27G .0209 (C) Medication Requirements	V 118		

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V 118	<p>Continued From page 2</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 with a physician.</p> <p> </p> <p>This Rule is not met as evidenced by: Based on observations, record reviews, and interviews, the facility failed to ensure</p>	V 118		

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V 118	<p>Continued From page 3</p> <p>medications were administered as ordered affecting 1 of 6 clients (#6). The findings are:</p> <p>Review on 9/20/21 of client #6's physician's orders dated 7/14/21 revealed:</p> <ul style="list-style-type: none"> <li>- Zyrtec 10 mg: take 1 tablet daily.</li> </ul> <p>Observations at approximately 2:21 pm on 9/21/21 of client #6's medications revealed:</p> <ul style="list-style-type: none"> <li>- Client #6's bottle of Zyrtec was not in his medication container.</li> </ul> <p>Interview on 9/21/21 with the Qualified Professional revealed:</p> <ul style="list-style-type: none"> <li>- Client #6 had stayed with his father/legal guardian from 9/17/21-9/18/21.</li> <li>- His father/legal guardian did not bring back client #6's Zyrtec on 9/18/21.</li> <li>- Client #6 had not taken his Zyrtec for the past 3 days.</li> </ul>	V 118		