

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL092-389	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 06/13/2019
--	---	---	---

NAME OF PROVIDER OR SUPPLIER WAKE COUNTY GROUP HOME #2	STREET ADDRESS, CITY, STATE, ZIP CODE 4808 WHITEHALL AVENUE RALEIGH, NC 27604
--	---

(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	INITIAL COMMENTS An annual survey was completed on 6/13/19. This facility is licensed for the following service category: 10A NCAC 27G .5600C Supervised Living for Adults with Developmental Disabilities.	V 000		
V 119	27G .0209 (D) Medication Requirements 10A NCAC 27G .0209 MEDICATION REQUIREMENTS (d) Medication disposal: (1) All prescription and non-prescription medication shall be disposed of in a manner that guards against diversion or accidental ingestion. (2) Non-controlled substances shall be disposed of by incineration, flushing into septic or sewer system, or by transfer to a local pharmacy for destruction. A record of the medication disposal shall be maintained by the program. Documentation shall specify the client's name, medication name, strength, quantity, disposal date and method, the signature of the person disposing of medication, and the person witnessing destruction. (3) Controlled substances shall be disposed of in accordance with the North Carolina Controlled Substances Act, G.S. 90, Article 5, including any subsequent amendments. (4) Upon discharge of a patient or resident, the remainder of his or her drug supply shall be disposed of promptly unless it is reasonably expected that the patient or resident shall return to the facility and in such case, the remaining drug supply shall not be held for more than 30 calendar days after the date of discharge.	V 119		

Division of Health Service Regulation
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE _____ TITLE _____ (X6) DATE _____

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL092-389	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 06/13/2019
--	---	---	---

NAME OF PROVIDER OR SUPPLIER WAKE COUNTY GROUP HOME #2	STREET ADDRESS, CITY, STATE, ZIP CODE 4808 WHITEHALL AVENUE RALEIGH, NC 27604
--	---

(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 119	<p>Continued From page 1</p> <p>This Rule is not met as evidenced by: Based on observation, record review and interviews the facility failed to assure medications were disposed of to guard against accidental ingestion for 1 of 3 audited clients (#5). The findings are:</p> <p>Observation on 6/13/19 at approximately 10:20 am of client #5's medications revealed Cetirizine 10 mg tablets with an expiration date of 3/1/19.</p> <p>Review on 6/13/19 of client #5's record revealed: - an admission date of 1/28/85 - an Individual Support Plan dated 2/21/19 with diagnoses including Moderate Intellectual Developmental Disability, Seasonal Allergies and Elevated Cholesterol - a physician's order dated 10/30/18 indicated the above medication was discontinued</p> <p>During an interview on 6/13/19, the Manager reported she was not sure why the medication was on site.</p> <p>During an interview on 6/13/19, the Registered Nurse reported he was not sure why the medication was on site.</p>	V 119		
V 121	<p>27G .0209 (F) Medication Requirements</p> <p>10A NCAC 27G .0209 MEDICATION REQUIREMENTS (f) Medication review: (1) If the client receives psychotropic drugs, the governing body or operator shall be responsible for obtaining a review of each client's drug regimen at least every six months. The review shall be to be performed by a pharmacist or physician. The on-site manager shall assure that</p>	V 121		

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL092-389	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 06/13/2019
--	---	---	---

NAME OF PROVIDER OR SUPPLIER WAKE COUNTY GROUP HOME #2	STREET ADDRESS, CITY, STATE, ZIP CODE 4808 WHITEHALL AVENUE RALEIGH, NC 27604
--	---

(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 121	<p>Continued From page 2</p> <p>the client's physician is informed of the results of the review when medical intervention is indicated. (2) The findings of the drug regimen review shall be recorded in the client record along with corrective action, if applicable.</p> <p>This Rule is not met as evidenced by: Based on record review and interview the facility failed to ensure the 6 month drug regimen review was not completed for 1 of 3 audited clients (#3) on psychotropic medications. The findings are:</p> <p>Review on 6/13/19 of client #3's record revealed: - admitted 05/10/99 - diagnoses anxiety disorder, mild mental retardation, seizure disorder - a physician's order dated 2/6/19 for abilify</p> <p>During an interview on 6/13/19, the Qualified Professional reported the agency was trying to identify a pharmacy to complete the drug regimen reviews. Review of April, May and June 2019 medication administration record revealed medication was given.</p>	V 121		