

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL084-098	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 03/10/2022
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NAME OF PROVIDER OR SUPPLIER NEW LONDON GROUP HOME	STREET ADDRESS, CITY, STATE, ZIP CODE 40163 HIGHWAY 740 NEW LONDON, NC 28127
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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 March 10, 2022. Deficiency cited.</p> <p>This facility is licensed for the following service category: 10A NCAC 27G. 5600C Supervised Living for Adults with Developmental Disabilities</p> <p>The survey sample consisted of audits of 3 current clients.</p>	V 000		
V 105	<p>27G .0201 (A) (1-7) Governing Body Policies</p> <p>10A NCAC 27G .0201 GOVERNING BODY POLICIES</p> <p>(a) The governing body responsible for each facility or service shall develop and implement written policies for the following:</p> <p>(1) delegation of management authority for the operation of the facility and services;</p> <p>(2) criteria for admission;</p> <p>(3) criteria for discharge;</p> <p>(4) admission assessments, including:</p> <p>(A) who will perform the assessment; and</p> <p>(B) time frames for completing assessment.</p> <p>(5) client record management, including:</p> <p>(A) persons authorized to document;</p> <p>(B) transporting records;</p> <p>(C) safeguard of records against loss, tampering, defacement or use by unauthorized persons;</p> <p>(D) assurance of record accessibility to authorized users at all times; and</p> <p>(E) assurance of confidentiality of records.</p> <p>(6) screenings, which shall include:</p> <p>(A) an assessment of the individual's presenting problem or need;</p> <p>(B) an assessment of whether or not the facility can provide services to address the individual's needs; and</p>	V 105		

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

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V 105	<p>Continued From page 1</p> <p>(C) the disposition, including referrals and recommendations;</p> <p>(7) quality assurance and quality improvement activities, including:</p> <p>(A) composition and activities of a quality assurance and quality improvement committee;</p> <p>(B) written quality assurance and quality improvement plan;</p> <p>(C) methods for monitoring and evaluating the quality and appropriateness of client care, including delineation of client outcomes and utilization of services;</p> <p>(D) professional or clinical supervision, including a requirement that staff who are not qualified professionals and provide direct client services shall be supervised by a qualified professional in that area of service;</p> <p>(E) strategies for improving client care;</p> <p>(F) review of staff qualifications and a determination made to grant treatment/habilitation privileges:</p> <p>(G) review of all fatalities of active clients who were being served in area-operated or contracted residential programs at the time of death;</p> <p>(H) adoption of standards that assure operational and programmatic performance meeting applicable standards of practice. For this purpose, "applicable standards of practice" means a level of competence established with reference to the prevailing and accepted methods, and the degree of knowledge, skill and care exercised by other practitioners in the field;</p>	V 105		

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V 105	<p>Continued From page 2</p> <p>This Rule is not met as evidenced by: Based on record review and interview, the facility failed to develop and implement adoption of standards that ensured operational and programmatic performance meeting applicable standards of practice for random drug testing instrument including the CLIA (Clinical Laboratory Improvement Amendments) waiver. The findings are:</p> <p>Review on 3/10/22 of Client #3's record revealed: -Admission date of 2/21/22. -Diagnoses of Borderline Personality Disorder, Bipolar Disorder, Current Episode Depressed, Moderate, Major Depressive Disorder, Recurrent Unspecified, Mild Intellectual Developmental Disability, Type 2 Diabetes Mellitus Without Complications and Gastro-Esophageal Reflux Disease Without Esophagitis. -Physician order dated 2/21/22 included the following orders: -Blood Glucose Test - Check Blood Glucose three times a week. -BD Pen Needle 31Gx 5mm - use as directed with Levemir Flex pen. -Levemir Injection - Inject 20 unit subcutaneously at bedtime. -Trulicity inject 1.5/0-5 - inject.5ml subcutaneously once a week.</p> <p>Review on 3/10/22 of the facility's documents revealed: -There was no evidence of a CLIA waiver.</p> <p>Observation on 3/10/22 of Client #1's diabetic medication revealed: -The injection was in a locked box in the refrigerator.</p>	V 105		

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V 105	<p>Continued From page 3</p> <p>-Documentation of blood sugar check was recorded.</p> <p>Interview on 3/10/22 with the Program Manager revealed:</p> <p>-Confirmed staff administered client #3's blood sugar checks and injections.</p> <p>-Confirmed the facility did not have a CLIA Waiver.</p>	V 105		