

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL076-131	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 04/07/2022
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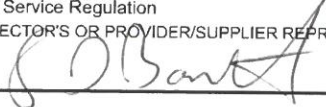
NAME OF PROVIDER OR SUPPLIER PATH OF HOPE, INC-ALPHA HOUSE	STREET ADDRESS, CITY, STATE, ZIP CODE 373 HILL STREET ASHEBORO, NC 27203
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(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING 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	INITIAL COMMENTS An annual survey was completed on April 7, 2022. Deficiency cited. This facility is licensed for the following service category: 10A NCAC 27G. 5600E Supervised Living for Substance Abuse Adults The survey sample consisted of audits of 3 current clients.	V 000		
V 105	27G .0201 (A) (1-7) Governing Body Policies 10A NCAC 27G .0201 GOVERNING BODY POLICIES (a) The governing body responsible for each facility or service shall develop and implement written policies for the following: (1) delegation of management authority for the operation of the facility and services; (2) criteria for admission; (3) criteria for discharge; (4) admission assessments, including: (A) who will perform the assessment; and (B) time frames for completing assessment. (5) client record management, including: (A) persons authorized to document; (B) transporting records; (C) safeguard of records against loss, tampering, defacement or use by unauthorized persons; (D) assurance of record accessibility to authorized users at all times; and (E) assurance of confidentiality of records. (6) screenings, which shall include: (A) an assessment of the individual's presenting problem or need; (B) an assessment of whether or not the facility can provide services to address the individual's needs; and (C) the disposition, including referrals and	V 105	<i>Waiting to see rule requiring CLIA if the test we use do not require CLIA.</i>	5-15-22

RECEIVED
APR 25 2022

DHSR-MH Licensure Sect

Division of Health Service Regulation
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE



TITLE

Executive Director

(X6) DATE

4-13-22

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V 105	<p>Continued From page 1</p> <p>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	Continued From page 2 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: The facility presented no evidence of a CLIA Waiver. Observation on 4/5/22 of the facility's drug testing kits revealed: -There was a clear cup that labeled the group of different drugs. -Client's had to dispensed urine in the clear cup. -After clients dispensed in the cup, they would pull the strip to show the results. -The results identified positive or negative and what drug was in the client's system. -The alcohol kit was a 4-minute screening test. -Client's would put the strip in their mouth and wait 4 minutes to check the results. Interview on 4/7/22 with the Program Director revealed: -Confirmed the facility conducted random drug and alcohol test. -Every client admitted and attended the program received a random drug and alcohol test. -Clients that returned after a 24 and 48 hours leave pass was tested. -Confirmed there was no evidence the facility had a CLIA Waiver.	V 105		