

Division of Health Service Regulation

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: MHL054-159	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 08/14/2018
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NAME OF PROVIDER OR SUPPLIER MAPLEWOOD FACILITY	STREET ADDRESS, CITY, STATE, ZIP CODE 2002-G SHACKLEFORD ROAD KINSTON, NC 28502
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V 000	<p>INITIAL COMMENTS</p> <p>An annual and complaint survey was completed on August 14, 2018. One complaint was unsubstantiated (intake #NC001414170), two complaints were substantiated (intake #NC00141644 and #NC00141649). Deficiencies were cited.</p> <p>This facility is licensed for the following service category: 10A NCAC 27G .1900 Psychiatric Residential Treatment for Children and Adolescents.</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</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>needs; and (C) the disposition, including referrals and recommendations; (7) quality assurance and quality improvement activities, including: (A) composition and activities of a quality assurance and quality improvement committee; (B) written quality assurance and quality improvement plan; (C) methods for monitoring and evaluating the quality and appropriateness of client care, including delineation of client outcomes and utilization of services; (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; (E) strategies for improving client care; (F) review of staff qualifications and a determination made to grant treatment/habilitation privileges: (G) review of all fatalities of active clients who were being served in area-operated or contracted residential programs at the time of death; (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> <p>This Rule is not met as evidenced by:</p>	V 105		

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V 105	<p>Continued From page 2</p> <p>Based on record reviews and interviews the facility failed to develop and implement a written policy for adoption of standards of practice related to federal requirements for the reporting of events that result in the use of restraint or seclusion. The findings are:</p> <p>Review on 8/10/18 of LME-MCO (Local Management Entity-Managed Care Organization) communication Bulletin J287, "Clarifying the Reporting Standards for Psychiatric Residential Treatment Facilities[PRTF]" dated 5/11/18 revealed:</p> <ul style="list-style-type: none"> - "As a reminder, Serious Occurrences are any event that result in Restraint or Seclusion, Resident's Death, Any Serious Injury to a Resident, and a Resident's Suicide Attempt. NC [North Carolina] 483.374 specifies that facilities must report each Serious Occurrence to both the State Medicaid agency (Division of Medical Assistance - DMA) . . . " - "DMA receives reports of Serious Occurrences via the Incident Response and Improvement System (IRIS) managed by the Division of Mental Health, Developmental Disabilities and Substance Abuse Services . . . " <p>Review on 8/10/18 of the facility's "INCIDENT AND DEATH RESPONSE SYSTEM" policy last revised 11/1/17 revealed: "Upon learning of a Level II/III incident involving a consumer currently receiving services, [Licensee] shall document the event within the time frames specified in this policy using the DHHS [Department of Health and Human Services] Incident Response Improvement System (IRIS). Level II/III DHHS Incident and Death Report include:...b) Restrictive Intervention: additional documentation is required on the restrictive intervention details report. Level II any</p>	V 105		

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V 105	<p>Continued From page 3</p> <p>emergency, unplanned use or any planned use that exceeds Licensure Rules is administered by an unauthorized person, requires treatment by a licensed health professional. Level III any restrictive intervention that results in permanent physical or psychological impairment within 7 days. . ."</p> <p>Review on 8/10/18 of the facility's "LEVEL I INCIDENT REPORTING" policy effective 9/1/10 revealed that it did not address reporting of restrictive interventions.</p> <p>Review on 8/10/18 of the facility's "Consumer Death or Serious Occurrence/Sentinel Event" policy, last revised 11/1/17 revealed: "It is the policy of [Licensee] to define a Serious Occurrence/Sentinel Event as the death of a Consumer or any significant impairment of the physical condition of a Consumer as determined by [Licensee's] Primary Care Medical Director or other qualified Medical Personnel. This includes, but shall not be limited to, burns, lacerations, bone fractures, substantial hematomas, and injuries to internal organs, whether self-inflicted or inflicted by another person. Any allegation of abuse, neglect or exploitation shall also be considered a Serious Occurrence and reported and documented accordingly. Each Consumer Death or Serious Occurrence shall be reported and documented in accordance with Federal and State rules . . ."</p> <p>Review on 8/10/18 of Client #10's record revealed: - 11 year old male admitted to the facility 2/27/18. - Diagnoses included Oppositional Defiant disorder, Attention Deficit Hyperactivity Disorder, combined presentation. - Comprehensive Clinical Assessment dated</p>	V 105		

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V 105	<p>Continued From page 4</p> <p>2/27/18 included history of violence towards others, impulsive behaviors, verbal aggression, homicidal ideation, elopement, self-injurious behaviors, stealing, property damage, sexualized behaviors.</p> <p>- "Crisis Prevention and Intervention Plan" dated 2/27/18, included: "Restrictive Interventions: Every attempt will be made to de-escalate the crisis prior to the use of physical restraint or seclusion. Restrictive Intervention should be used when (Client #10) is at imminent risk of, or in the process of injuring self or others. Type: Physical Restraint 1. Duration Limit: The use of Physical Restraint will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 10 minutes has elapsed. . . . Type: Seclusion 1. Duration Limit: The use of Seclusion will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 1 hour elapsed. . . ."</p> <p>- "Consumer Safety Plan" signed 2/27/18 included: "PRTF Setting: . . . Staff will utilize restrictive interventions to de-escalate imminent risk situations that place the consumer and/or others in jeopardy once least restrictive interventions have been exhausted and proven ineffective. Restrictive interventions include: NCI [North Carolina Interventions] techniques, seclusion and chemical intervention. . . ."</p> <p>- "Order for Emergency Safety Interventions" dated 5/28/18, signed by physician 8/9/18, for use of physical restraint.</p> <p>Review on 8/10/18 of facility's Level I and Level II Incident Reports completed 5/1/18 - 8/10/18 revealed: - Level I Incident Report dated 5/18/18; ". . .</p>	V 105		

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V 105	<p>Continued From page 5</p> <p>Consumer (Client #10) was placed in a restraint." - No Level II Incident Report of the events that led to the use of the physical restraint for Client #10. - A Level II Incident Report dated 7/18/18 documented allegation of abuse by a staff member.</p> <p>Review on 8/10/18 of Client #11's record and facility's Level I and Level II Incident Reports completed 5/1/18 - 8/10/18 revealed: - 10 year old male admitted to the facility 11/16/17. - Diagnoses included Disruptive Mood Dysregulation Disorder, Attention Deficit Hyperactivity Disorder, combined presentation, Post Traumatic Stress Disorder. - Comprehensive Clinical Assessment dated 11/16/17 included history of elopement, physical and verbal aggression, elopement, property destruction, self injurious behaviors, suicide threats. - "Person Centered Profile" dated 11/16/17 included: "Strategies for crisis response and stabilization . . ."Restrictive Interventions: Every attempt will be made to de-escalate the crisis prior to the use of physical restraint or seclusion. Restrictive Intervention should be used when (Client #11) is at imminent risk of, or in the process of injuring self or others. Type: Physical Restraint 1. Duration Limit: The use of Physical Restraint will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 10 minutes has elapsed. . . . Type: Seclusion 1. Duration Limit: The use of Seclusion will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 1 hour elapsed. . . ."</p>	V 105		

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V 105	<p>Continued From page 6</p> <ul style="list-style-type: none"> - "Consumer Safety Plan" signed 11/16/17 included: "PRTF Setting: . . . Staff will utilize restrictive interventions to de-escalate imminent risk situations that place the consumer and/or others in jeopardy once least restrictive interventions have been exhausted and proven ineffective. Restrictive interventions include: NCI techniques, seclusion and chemical intervention. . . ." - 29 "Order for Emergency Safety Interventions" completed between May 1 and August 10, 2018, and signed by the physician, for the use of physical restraints with no corresponding Level I or Level II incident reports. - 11 "Order for Emergency Safety Interventions" completed between 5/1/18 and 8/10/18, signed by the physician, for the use of physical restraints, with corresponding Level I incident reports. - Level I Incident Report dated 5/14/18 for "Planned use of appropriately administered Seclusion or Restraint"; "Medical Progress Notes" written by the Registered Nurse and signed 5/14/18 documented the use of Ativan injections due to "out of control" physical aggression toward staff and peers and destructive behaviors. - No Level II Incident Report of the events that led to the use of physical restraints or chemical restraint for Client #11. - Level II Incident Report dated 7/22/18 documented incident of alleged abuse by a staff member. <p>Review on 8/10/18 of Client #13's record revealed:</p> <ul style="list-style-type: none"> - 14 year old male admitted to the facility 6/26/18. - Diagnoses included Conduct Disorder, Post Traumatic Stress Disorder, Oppositional Defiant Disorder, Attention Deficit Hyperactivity Disorder, combined presentation, Autism Spectrum Disorder, Disruptive Mood Dysregulation 	V 105		

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V 105	<p>Continued From page 7</p> <p>Disorder.</p> <p>- "Person Centered Profile" dated 6/26/18 included: "Strategies for crisis response and stabilization . . . "Restrictive Interventions: Every attempt will be made to de-escalate the crisis prior to the use of physical restraint or seclusion. Restrictive Intervention should be used when (Client #13) is at imminent risk of, or in the process of injuring self or others. Type: Physical Restraint 1. Duration Limit: The use of Physical Restraint will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 10 minutes has elapsed. . . . Type: Seclusion 1. Duration Limit: The use of Seclusion will be immediately discontinued at any indication of Consumer risk or distress, or immediately when the Consumer gains control over at-risk behaviors, or when 1 hour elapsed. . . ."</p> <p>- "Consumer Safety Plan" signed 6/26/18 included: "PRTF Setting: . . . Staff will utilize restrictive interventions to de-escalate imminent risk situations that place the consumer and/or others in jeopardy once least restrictive interventions have been exhausted and proven ineffective. Restrictive interventions include: NCI techniques, seclusion and chemical intervention. . . ."</p> <p>- 4 "Order for Emergency Safety Interventions" completed between 6/26/18 and 8/10/18, signed by the physician, for the use of physical restraints.</p> <p>Review on 8/10/18 of facility's Level I and Level II Incident Reports completed 5/1/18 - 8/10/18 revealed no Level I or Level II incident reports of the events that led to the use of physical restraints for Client #13.</p> <p>Interview on 8/14/18 the Director of PRTF</p>	V 105		

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V 105	Continued From page 8 Services stated Federal guidelines required PRTF reporting of "Serious Occurrences and Sentinel Events". The definition of "Serious Occurrence" did not include the use of restrictive interventions, including physical restraint, chemical restraint, or seclusion. The Licensee was seeking legal clarification of the reporting requirements.	V 105		
V 367	27G .0604 Incident Reporting Requirements 10A NCAC 27G .0604 INCIDENT REPORTING REQUIREMENTS FOR CATEGORY A AND B PROVIDERS (a) Category A and B providers shall report all level II incidents, except deaths, that occur during the provision of billable services or while the consumer is on the providers premises or level III incidents and level II deaths involving the clients to whom the provider rendered any service within 90 days prior to the incident to the LME responsible for the catchment area where services are provided within 72 hours of becoming aware of the incident. The report shall be submitted on a form provided by the Secretary. The report may be submitted via mail, in person, facsimile or encrypted electronic means. The report shall include the following information: (1) reporting provider contact and identification information; (2) client identification information; (3) type of incident; (4) description of incident; (5) status of the effort to determine the cause of the incident; and (6) other individuals or authorities notified or responding. (b) Category A and B providers shall explain any	V 367		

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V 367	<p>Continued From page 9</p> <p>missing or incomplete information. The provider shall submit an updated report to all required report recipients by the end of the next business day whenever:</p> <p>(1) the provider has reason to believe that information provided in the report may be erroneous, misleading or otherwise unreliable; or</p> <p>(2) the provider obtains information required on the incident form that was previously unavailable.</p> <p>(c) Category A and B providers shall submit, upon request by the LME, other information obtained regarding the incident, including:</p> <p>(1) hospital records including confidential information;</p> <p>(2) reports by other authorities; and</p> <p>(3) the provider's response to the incident.</p> <p>(d) Category A and B providers shall send a copy of all level III incident reports to the Division of Mental Health, Developmental Disabilities and Substance Abuse Services within 72 hours of becoming aware of the incident. Category A providers shall send a copy of all level III incidents involving a client death to the Division of Health Service Regulation within 72 hours of becoming aware of the incident. In cases of client death within seven days of use of seclusion or restraint, the provider shall report the death immediately, as required by 10A NCAC 26C .0300 and 10A NCAC 27E .0104(e)(18).</p> <p>(e) Category A and B providers shall send a report quarterly to the LME responsible for the catchment area where services are provided. The report shall be submitted on a form provided by the Secretary via electronic means and shall include summary information as follows:</p> <p>(1) medication errors that do not meet the definition of a level II or level III incident;</p> <p>(2) restrictive interventions that do not meet</p>	V 367		

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V 367	<p>Continued From page 10</p> <p>the definition of a level II or level III incident; (3) searches of a client or his living area; (4) seizures of client property or property in the possession of a client; (5) the total number of level II and level III incidents that occurred; and (6) a statement indicating that there have been no reportable incidents whenever no incidents have occurred during the quarter that meet any of the criteria as set forth in Paragraphs (a) and (d) of this Rule and Subparagraphs (1) through (4) of this Paragraph.</p> <p>This Rule is not met as evidenced by: Based on record reviews and interviews the facility failed to submit Level II incident reports as required. The findings are:</p> <p>Refer to Tag v105 for details.</p>	V 367		