

Department of Health & Human Services
Centers for Medicare & Medicaid Services

Printed: 07/30/2026
Form Approved OMB
No. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 415078	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 05/26/2026
NAME OF PROVIDER OR SUPPLIER Coventry Operations RI LLC DbA Respiratory and Reh		STREET ADDRESS, CITY, STATE, ZIP CODE 10 Woodland Drive Coventry, RI 02816	
For information on the nursing home's plan to correct this deficiency, please contact the nursing home or the state survey agency.			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (Each deficiency must be preceded by full regulatory or LSC identifying information)		
F 0695 Level of Harm - Immediate jeopardy to resident health or safety Residents Affected - Few	Provide safe and appropriate respiratory care for a resident when needed. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on surveyor observation, clinical record review, and staff interview, the facility failed to provide necessary respiratory care and services in accordance with professional standards of practice for a resident who requires mechanical ventilation (a life-support therapy that assists or replaces natural breathing by delivering oxygen and removing carbon dioxide through a ventilator) who experienced chest pain and was sent via rescue to an acute care hospital for evaluation, Resident ID #1, who ultimately expired in transit to the hospital. Further, the facility failed to ensure that respiratory staff provide respiratory care consistent with professional standards of practice for 4 of 6 respiratory staff reviewed, Staff A, C, D, and E to ensure that a physician's order for mechanical ventilation settings were in place for 1 of 1 resident reviewed for non-invasive ventilation (NIV, a machine that provides breathing support using an external mask or other component, eliminating the need for an invasive surgical airway or breathing tube), Resident ID #1. Additionally, the facility failed to ensure that a resident who requires mechanical ventilation receives the mechanical ventilation settings as prescribed by the provider for 2 of 5 residents reviewed, Resident ID #s 1 and 3. Findings are as follows: Review of a complaint received by the Rhode Island Department of Health on [DATE] alleges in part, "[Resident ID #1]. had chest pains so [s/he] agreed to go to the hospital in the ambulance. We [family member] first got a call from a doctor from [the acute care hospital] telling us that [Resident ID #1] had not made it to the hospital and died on [his/her] way. Additionally, the complaint alleges that the facility failed to ensure that Resident ID #1 was provided with the necessary respiratory equipment prior to leaving the facility via rescue and that Resident ID #1 had been receiving the incorrect ventilator settings. 1a. Review of an undated facility policy titled, Ventilator Patient External Transport Procedure states in part, ".ensure safe transport of mechanically ventilated patients outside the facility for hospital transfer while maintaining adequate ventilation and patient stability. All ventilated patients requiring external transport must be assessed by Respiratory Therapy prior to departure to ensure appropriate equipment. Indications. Hospital transfer. Emergency transport. Required Equipment * Portable ventilator with charged battery. Transport Procedure. Transfer patient to portable ventilator if required. According to an article published in April of 2021 in the American Journal of the Medical Sciences titled, Non-Rebreather Mask [NRM, a type of oxygen mask that delivers high concentrations of oxygen to individuals who can breathe on their own]: A Bridge Worth Crossing? states in part, ".NRM does not provide any positive pressure to help support the patient mechanically. Record review revealed Resident ID #1 was admitted to the facility in March of 2026 with diagnoses including, but not limited to, duchenne or [NAME] muscular dystrophy (a severe, progressive genetic disorder characterized by the wasting away of skeletal, heart, and respiratory muscles), dependence on respirator (ventilator) status, and chronic respiratory failure. Review of the care plan revealed a focus area dated [DATE] indicating that Resident ID #1 has an alteration in respiratory status and is ventilator dependent. Further, the care plan revealed that s/he receives ventilation support using non-invasive ventilation via mouthpiece ventilation equipment during the daytime. Additionally, s/he uses nasal pillows [soft inserts that fit directly into the nostrils to provide pressurized air from the ventilator machine via a flexible tube to help keep the airway open, improve (continued on next page)		

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
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F 0695 Level of Harm - Immediate jeopardy to resident health or safety Residents Affected - Few	follow the prescribed ventilator settings. He further indicated that staff do not have the autonomy to adjust ventilator settings without first speaking to a provider.2b. Record review revealed Resident ID #3 was readmitted to the facility in April of 2026 with a diagnosis including, but not limited to, dependence on respirator (ventilator) status.Review of a care plan focus area dated [DATE] revealed that Resident ID #3 is ventilator dependent with an intervention to use the ventilator for respiratory support per the physician's order.Review of a physician's order dated [DATE] revealed in part, the following ventilator setting:-FiO2: 3 liters of oxygenDuring a surveyor observation and simultaneous interview on [DATE] at approximately 1:40 PM of Resident ID #3, in the presence of Staff C, Resident ID #3 was noted lying in his/her bed connected to his/her ventilator. Further, his/her ventilator settings were observed, and s/he was noted to be receiving only room air, and was not receiving the 3 liters of oxygen, as prescribed.During a surveyor interview immediately following the above observation with Staff C, she was unable to provide evidence that Resident ID #3 had his/her ventilator settings set to include the 3 liters of oxygen, as prescribed.During a surveyor interview on [DATE] at 12:53 PM with the DNS, she was unable to provide evidence that the facility provided necessary respiratory care and services in accordance with professional standards of practice and per the facility's policies.Cross reference F 842		

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F 0842 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Some	Safeguard resident-identifiable information and/or maintain medical records on each resident that are in accordance with accepted professional standards. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on clinical record review and staff interview, the facility failed to maintain complete and accurate resident medical records in accordance with accepted professional standards and practices for 1 of 1 resident reviewed for non-invasive ventilation (NIV, a machine that provides breathing support using an external mask or other component, eliminating the need for an invasive surgical airway or breathing tube), Resident ID #1. Findings are as follows:Review of a complaint received by the Rhode Island Department of Health on 5/19/2026 alleges, in part, that Resident ID #1 had been receiving the incorrect ventilator settings.Record review revealed Resident ID #1 was admitted to the facility in March of 2026 with diagnoses including, but not limited to, duchenne or [NAME] muscular dystrophy (a severe, progressive genetic disorder characterized by the wasting away of skeletal, heart, and respiratory muscles), dependence on respirator (ventilator) status, and chronic respiratory failure.Review of the care plan revealed a focus area dated 3/5/2026 indicating that Resident ID #1 has an alteration in respiratory status and is ventilator dependent. Further, the care plan revealed that s/he receives ventilation support using non-invasive ventilation via mouthpiece ventilation equipment during the daytime. Additionally, s/he uses nasal pillows [soft inserts that fit directly into the nostrils to provide pressurized air from the ventilator machine via a flexible tube to help keep the airway open, improve breathing, and prevent an individual from not breathing] at nighttime. Further, interventions include, but are not limited to, ventilator check assessments to be conducted every six hours and to use the ventilator for respiratory support per the physician's order.Review of an undated facility policy titled, Non-Invasive Ventilator (NIV) Policy & Procedure states in part, .Purpose.Promote patient safety and improve respiratory support management.Procedure.Verify physician/provider order and prescribed settings.Record review revealed Resident ID #1 utilized the following three ventilators while at the facility between 3/2/2026 through 3/14/2026:-Astral (daytime ventilator)-LTV (nighttime ventilator)-VOCSN (nighttime ventilator)Record review revealed the following physician's order for the resident's LTV ventilator with the prescribed settings:-Tidal volume (VT, the amount of air inhaled or exhaled during a normal breath) 1100-Respiratory rate (RR, the number of breaths a person takes per minute) 14-Positive end-expiratory pressure (PEEP, the pressure in the lungs above atmospheric pressure at the end of exhalation that prevents the lungs from collapsing when breathing out) 6-Fraction of inspired oxygen (FiO2, refers to the oxygen concentration; room air is considered 21% oxygen) 21%-Pressure support (PS, a mode of mechanical ventilation that assists patients by providing a preset level of pressure during inhalation) 8Record review revealed that the facility discontinued using the LTV ventilator and began using the VOCSN ventilator at nighttime beginning on 3/4/2026 through 3/14/2026.Review of the March 2026 Treatment Administration Record revealed that staff continued to document that Resident ID #1 was receiving his/her LTV ventilator at the prescribed settings as denoted above, on the following dates, after it was discontinued on 3/4/2026:-3/4-3/5-3/6-3/8-3/9-3/10-3/11-3/12-3/14/2026During a surveyor interview on 5/22/2026 at approximately 11:25 AM with the Physician Assistant, he revealed that he expects staff to document accurately.During a surveyor interview on 5/22/2026 at 12:53 PM with the Director of Nursing Services, she was unable to provide evidence that the facility maintained accurate medical records for Resident ID #1.Cross reference F 695		