

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 155751	X2) MULTIPLE CONSTRUCTION A. BUILDING -- _____ B. WING _____	X3) DATE SURVEY COMPLETED 04/01/2025
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NAME OF PROVIDER OR SUPPLIER MEADOW LAKES	STREET ADDRESS, CITY, STATE, ZIP CODE 200 MEADOW LAKE DR MOORESVILLE, IN 46158
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E 0000 Bldg. --	<p>An Emergency Preparedness Survey was conducted by the Indiana Department of Health in accordance with 42 CFR 483.73.</p> <p>Survey Date: 04/01/25</p> <p>Facility Number: 004831 Provider Number: 155751 AIM Number: 200809750</p> <p>At this Emergency Preparedness survey, Meadow Lakes was found in compliance with Emergency Preparedness Requirements for Medicare and Medicaid Participating Providers and Suppliers, 42 CFR 483.73</p> <p>The facility has 137 certified beds. At the time of the survey, the census was 119.</p> <p>Quality Review completed on 04/04/25</p>	E 0000		
K 0000 Bldg. 01	<p>A Life Safety Code Recertification and State Licensure Survey was conducted by the Indiana Department of Health in accordance with 42 CFR 483.90(a).</p> <p>Survey Date: 04/01/25</p> <p>Facility Number: 004831 Provider Number: 155751 AIM Number: 200809750</p> <p>At this Life Safety Code survey, Meadow Lakes was found not in compliance with Requirements</p>	K 0000	The submission of this plan of correction does not indicate an admission by Meadow Lakes Nursing Facility that the findings and allegations contained herein are an accurate and true representation of the quality of care and environment provided to the residents of this facility. This facility recognizes its obligation to provide legally and medically necessary care and service in a safe environment for its residents	

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
Mac McCallum	Executive Director	04/15/2025

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined other safeguards provide sufficient protection to the patients. (see instructions.) Except for nursing homes, the findings stated above are disclosable 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 disclosed 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.

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K 0324 SS=E Bldg. 01	<p>for Participation in Medicare/Medicaid, 42 CFR Subpart 483.90(a), Life Safety from Fire and the 2012 Edition of the National Fire Protection Association (NFPA) 101, Life Safety Code (LSC), Chapter 19, Existing Health Care Occupancies and 410 IAC 16.2.</p> <p>This one-story facility was determined to be of Type V (111) construction and fully sprinklered. The facility has a fire alarm system with smoke detection in the corridors and in all areas open to the corridor. The facility has smoke detectors hard wired to the fire alarm system in all resident sleeping rooms. The facility has a capacity of 137 and had a census of 119 at the time of this visit.</p> <p>All areas where the residents have customary access were sprinklered and all areas providing facility services were sprinklered.</p> <p>Quality Review completed on 04/04/25</p> <p>NFPA 101 Cooking Facilities</p> <p>Based on observation and interview, the facility failed to provide an approved method for returning cooking appliances to where they were when the kitchen hood extinguishing equipment was designed and installed for 1 of 1 kitchen hood extinguishing system. NFPA 96, Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations Section 2011 Edition Section 12.1.2.2, states cooking appliances requiring protection shall not be moved, modified, or rearranged without prior re-evaluation of the fire-extinguishing system by the system installer or servicing agent, unless otherwise allowed by</p>	K 0324	<p>in an economic and safe manner. The facility hereby maintains it is in substantial compliance with the requirements of participation for Nursing facilities. To this end, this plan of correction shall serve as the credible allegation of compliance with all state requirements governing the management of this facility. It is thus submitted as a matter of statute only.</p> <p>This facility respectfully requests from the Department a desk review. If anything, further is needed the facility will provide department documentation upon request for paper compliance/desk review.</p> <p><u>K324 Cooking Facilities:</u> Director of plant operations immediately placed floor markings to identify and return cooking appliances to where they were when the kitchen hood extinguishing equipment was designed and installed for 1 of 1 kitchen hood extinguishing system.</p> <p><u>Method to Assess:</u> The director of plant operations will</p>	04/02/2025

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	<p>the design of the fire extinguishing system. Section 12.1.2.3 states the fire-extinguishing system shall not require reevaluation where the cooking appliances are moved for the purposes of maintenance and cleaning, provided the appliances are returned to approved design location prior to cooking operations, and any disconnected fire-extinguishing system nozzles attached to the appliances are reconnected in accordance with the manufacturer's listed design manual. Section 12.1.2.3.1 states an approved method shall be provided that will ensure that the appliance is returned to an approved design location. The deficient practice could affect as many as 32 residents, 6 staff, and 2 visitors in the facility.</p> <p>Findings include:</p> <p>Based on observations made during a tour of the facility with the Maintenance Director on 04/01/25 at 12:25 p.m., the four (4) burner stove and the flat grill which was located on the cooking line under the hood in the kitchen was not provided with an approved method that would ensure that the appliance was returned to an approved design location after it had been moved for maintenance and/or cleaning. Based on interview on 04/01/25 at 12:28 p.m., the Maintenance Director stated that he was not aware an approved method should be provided to ensure that the appliance was returned to an approved design location after maintenance or cleaning and that he would have something done to the kitchen stove or floor to meet code compliance as soon as possible.</p> <p>This item was discussed with the Maintenance Director and the facility Administrator at the exit conference on 04/01/25.</p>		<p>visually inspect floor return markings daily to ensure that identification and return is completed in alignment with NFPA 96, Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations Section 2011 Edition Section 12.1.2.2 .</p> <p><u>Systematic Process:</u> The director of plant operations or designee will complete Floor/Audit checks daily until substantial compliance is maintained. Quality Assurance Executive Director/Designee will present the results of any visual inspection to the QAPI committee for further recommendations and will continue until QAPI team determines substantial compliance has been achieved.</p>	

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K 0921 SS=E Bldg. 01	<p>3.1-19(b)</p> <p>NFPA 101 Electrical Equipment - Testing and Maintenance</p> <p>Based on record review, observation, and interview, the facility failed to conduct the required maintenance and maintain complete documentation of inspections for Patient Care Related Electrical Equipment (PCREE). NFPA 99 2012 edition, sections 10.3 and 10.5 states the physical integrity, resistance, leakage current, and touch current tests for fixed and portable PCREE is performed as required in 10.3. Testing intervals are established with policies and protocols. All PCREE used in patient care rooms is tested in accordance with 10.3.5.4 or 10.3.6 before being put into service and after any repair or modification. Any system consisting of several electrical appliances demonstrates compliance with NFPA 99 as a complete system. Service manuals, instructions, and procedures provided by the manufacturer include information as required by 10.5.3.1.1 and are considered in the development of a program for electrical equipment maintenance. Electrical equipment instructions and maintenance manuals are readily available, and safety labels and condensed operating instructions on the appliance are legible. A record of electrical equipment tests, repairs, and modifications is maintained for a period of time to demonstrate compliance in accordance with the facility's policy. Personnel responsible for the testing, maintenance and use of electrical appliances receive continuous training. This deficient practice could affect all residents.</p> <p>Findings include:</p> <p>Based on record review on 04/01/25 at 10:05 a.m.</p>	K 0921	<p><u>K921 Electrical Equipment - Testing and Maintenance:</u></p> <p>Director of plant operations or Designees conducted the required maintenance and audit documentation of inspections for Patient Care Related Electrical Equipment (PCREE) on 4.7.2025 for the entire facility. This audit was completed in accordance of NFPA 99 2012 edition, sections 10.3 and 10.5 states the physical integrity, resistance, leakage current, and touch current tests for fixed and portable PCREE is performed as required in 10.3.</p> <p><u>Method to Assess:</u> The director of plant operations will inspect all PCREE used in patient care rooms in accordance with 10.3.5.4 or 10.3.6 before being put into service and after any repair or modification. This inspection will occur as needed and prior to use to ensure all compliance related to manufacture guidelines and requirements is maintained. All records and services will be upheld and maintained by the director of maintenance.</p> <p><u>Systematic Process:</u> The director of plant operations or designee will complete</p>	04/07/2025
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K 0927 SS=E Bldg. 01	<p>with the Maintenance Director present, there was no documentation for the testing of Patient Care Related Electrical Equipment (PCREE), such as electric beds, nebulizers, oxygen concentrators, air pumps for air mattresses, and other electrical medical equipment. Based on interview on 04/01/25 at 10:07 a.m., the Maintenance Director stated that he was unaware of the requirement for PCREE testing and would begin doing so as soon as he could. Based on observations made during a tour of the facility, it was noted the facility provided PCREE such as electric beds, air pumps for air mattresses, and other electrical medical equipment that was present and in use within the facility.</p> <p>This finding was reviewed with the Facility Administrator, the Maintenance Director, and Field Maintenance Supervisor during the exit conference on 04/01/25.</p> <p>3.1-19(b)</p> <p>NFPA 101 Gas Equipment - Transfilling Cylinders</p> <p>Based on observation and interview, the facility failed to ensure 1 of 1 oxygen storage room, where oxygen transferring takes place, was provided with properly working mechanical ventilation for the room. This deficient practice could affect as many as 8 staff working in the area of the oxygen storage and transferring room.</p> <p>Findings include:</p> <p>Based on observations made on 04/01/25 at 12:10 p.m. during a tour of the facility with the Maintenance Director, the Field Maintenance Supervisor, and the facility Administrator, the oxygen storage/transfer room had six large liquid</p>	K 0927	<p>PCREE/Audit checks as needed until substantial compliance is maintained. Audits include external condition, ground wire resistance, touch current grounded, and touch current non-grounded. Quality Assurance Executive Director/Designee will present the results of any visual inspection to the QAPI committee for further recommendations and will continue until QAPI team determines substantial compliance has been achieved.</p> <p><u>K927 Gas Equipment - Transfilling Cylinders</u></p> <p>Director of plant operations contacted facility vendor Safe Care for further inspection and repair/replacement of ceiling exhaust fan.</p> <p><u>Method to Assess:</u> The director of plant operations will visually inspect all mechanical ventilation rooms daily for proper functioning to ensure that the transfilling to liquid oxygen containers or to portable containers under 50 psi complies</p>	04/18/2025

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	<p>oxygen tanks. There was a mechanically ventilated exhaust fan in the ceiling of this room, however, it was not working at the time of observation. The Maintenance Director tested this by placing a tissue paper up against the fan to see if the fan took hold of it, but it did not. Based on an interview on 04/01/25 at 12:15 p.m., the Maintenance Director agreed that the exhaust fan was not working and stated that he would have it replaced as soon as possible.</p> <p>This finding was reviewed with the Facility Administrator, the Maintenance Director, and the Field Maintenance Supervisor during the exit conference on 04/01/25.</p> <p>3.1-19(b)</p>		<p>with conditions under 11.5.2.3.2 (NFPA 99). 11.5.2.2 (NFPA 99) <u>Systematic Process:</u> The director of plant operations or designee will complete Daily Audit checks to ensure proper functioning of exhaust units. Quality Assurance Executive Director/Designee will present the results of any visual inspection to the QAPI committee for further recommendations and will continue until QAPI team determines substantial compliance has been achieved.</p>		