

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

PRINTED: 06/19/2024

FORM APPROVED

OMB NO. 0938-039

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER 155443		X2) MULTIPLE CONSTRUCTION A. BUILDING -- B. WING _____		X3) DATE SURVEY COMPLETED 05/29/2024	
NAME OF PROVIDER OR SUPPLIER WATERS OF MUNCIE, THE				STREET ADDRESS, CITY, STATE, ZIP COD 2400 CHATEAU DR MUNCIE, IN 47303			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIE (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) COMPLETION DATE
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: 05/29/24</p> <p>Facility Number: 000310 Provider Number: 155443 AIM Number: 100288970</p> <p>At this Emergency Preparedness survey, The Waters of Muncie was found in compliance with Emergency Preparedness Requirements for Medicare and Medicaid Participating Providers and Suppliers, 42 CFR 483.73. The facility has a capacity of 72 and had a census of 50 at the time of this survey.</p> <p>Quality Review completed on 05/31/24</p>			E 0000	<p>Preparation and/or execution of this plan of correction in general, or this corrective action does not constitute an admission agreement by this facility of the facts alleged or conclusions set forth in this statement of deficiencies. The plan of correction and specific corrective actions are prepared and/or executed in compliance with state and federal laws. This plan of correction constitutes our credible allegation of compliance with all regulatory requirements. Our date of compliance is 06/03/2024.</p> <p>Facility respectfully requests a desk review.</p>		
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: 05/29/24</p> <p>Facility Number: 000310 Provider Number: 155443 AIM Number: 100288970</p> <p>At this Life Safety Code survey, The Waters of Muncie was found not in compliance with Requirements for Participation in</p>			K 0000	<p>Preparation and/or execution of this plan of correction in general, or this corrective action does not constitute an admission agreement by this facility of the facts alleged or conclusions set forth in this statement of deficiencies. The plan of correction and specific corrective actions are prepared and/or executed in compliance with state and federal laws. This plan of correction constitutes our credible allegation of compliance with all regulatory</p>		

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

Brenda Darlien Alfrey

Executive Director

06/19/2024

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 0363 SS=E Bldg. 01	<p>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 V111 construction and was fully sprinklered. The facility has a fire alarm system with smoke detection in the corridors, areas open to the corridors and battery powered smoke detection in the resident sleeping rooms. The facility has a capacity of 72 and had a census of 50 at the time of this survey.</p> <p>All areas where the residents have customary access were sprinklered. All areas providing facility services were sprinklered.</p> <p>Quality Review completed on 05/31/24</p> <p>NFPA 101 Corridor - Doors Corridor - Doors Doors protecting corridor openings in other than required enclosures of vertical openings, exits, or hazardous areas resist the passage of smoke and are made of 1 3/4 inch solid-bonded core wood or other material capable of resisting fire for at least 20 minutes. Doors in fully sprinklered smoke compartments are only required to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible materials have positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Clearance between bottom of door and floor</p>				<p>requirements. Our date of compliance is 06/03/2024. Facility respectfully requests a desk review.</p>		

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	<p>covering is not exceeding 1 inch. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5 lbf is applied. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 19.3.6.3.6 are permitted. Door frames shall be labeled and made of steel or other materials in compliance with 8.3, unless the smoke compartment is sprinklered. Fixed fire window assemblies are allowed per 8.3. In sprinklered compartments there are no restrictions in area or fire resistance of glass or frames in window assemblies.</p> <p>19.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485</p> <p>Show in REMARKS details of doors such as fire protection ratings, automatics closing devices, etc.</p> <p>Based on observation and interview, the facility failed to ensure 1 of around 50 sets of resident room doors to the corridor would close completely and latch into the door frame. This deficient practice could affect as many as 2 residents, 2 staff and 2 visitors in the facility.</p> <p>Findings include:</p> <p>Based on observations made during a tour of the facility on 05/29/24 at 1:12 p.m. with the facility Administrator and the Maintenance Director present, resident room # 304 failed to latch into the door frame. Based on interview at the time of observations, the Maintenance Director acknowledged the resident room door as not fully</p>			K 0363	<p>K363 – It is the intent of the facility to provide resident room doors to the corridor that will close completely and latch into the door frame to meet set standards.</p> <p>1 CORRECTIVE ACTIONS TAKEN:</p> <p>a On 5/29/2024 the Maintenance Supervisor/designee made repairs to resident room #304 door to ensure door latches into the door frame to meet set standards. The Administrator verified the repairs on 5/29/2024.</p> <p>2 ALL OTHERS WITH POTENTIAL TO BE AFFECTED:</p>		06/03/2024

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	<p>closing and latching into the doorframe adding that he would look at it as soon as possible.</p> <p>This item was again discussed at the exit conference with the facility Administrator and the Maintenance Director on 05/29/24 at 1:55 p.m.</p> <p>3.1-19(b)</p>		<p>a All residents and all staff and visitors have the potential to be affected but none were. The Maintenance Supervisor/designee inspected all doors and found no other negative findings.</p> <p>3 MEASURES TO PREVENT REOCCURRENCE:</p> <p>a On 5/29/2024 the Administrator in serviced the Maintenance Supervisor/and all other staff members on the requirement to provide resident room doors that would close completely and latch fully into the frame to meet set standards.</p> <p>b Maintenance Supervisor/designee will inspect all doors throughout the facility monthly to ensure resident room doors will close completely and latch fully into the frame as a part of the facility's Preventive Maintenance Program and document those inspection results as appropriate. If any issues are discovered, they will be addressed and resolved immediately. The Maintenance Supervisor/designee will review with the Administrator the inspection results.</p> <p>c The Administrator will monitor adherence to the Preventative Maintenance schedule and validate the Preventative Maintenance documentation is in place.</p> <p>4 MONITORING CORRECTIVE ACTION:</p> <p>a The inspection results will</p>		

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K 0914 SS=E Bldg. 01	NFPA 101 Electrical Systems - Maintenance and Testing Electrical Systems - Maintenance and Testing Hospital-grade receptacles at patient bed locations and where deep sedation or general anesthesia is administered, are tested after initial installation, replacement or servicing. Additional testing is performed at intervals defined by documented performance data. Receptacles not listed as hospital-grade at these locations are tested at intervals not exceeding 12 months. Line isolation monitors (LIM), if installed, are tested at intervals of less than or equal to 1 month by actuating the LIM test switch per 6.3.2.6.3.6, which		be presented by the Maintenance Supervisor/designee to the Administrator monthly and the Administrator will present the inspection results at the monthly Quality Assurance/Performance Improvement (QA/PI) meeting. Inspection results and system components will be reviewed by the QA/PI Committee with subsequent plans of correction developed and implemented as deemed necessary to ensure compliance is maintained. This plan of correction constitutes our credible allegation of compliance with all regulatory requirements. Our date of compliance is 06.03.24 Facility respectfully requests a desk review.		

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	<p>activates both visual and audible alarm. For LIM circuits with automated self-testing, this manual test is performed at intervals less than or equal to 12 months. LIM circuits are tested per 6.3.3.3.2 after any repair or renovation to the electric distribution system. Records are maintained of required tests and associated repairs or modifications, containing date, room or area tested, and results.</p> <p>6.3.4 (NFPA 99) Based on record review, observation, and interview, the facility failed to ensure all nonhospital-grade electrical receptacles at resident room locations were tested at least annually. NFPA 99, Health Care Facilities Code 2012 Edition, Section 6.3.4.1.3 states receptacles not listed as hospital-grade, at patient bed locations and in locations where deep sedation or general anesthesia is administered, shall be tested at intervals not exceeding 12 months. Additionally, Section 6.3.3.2, Receptacle Testing in Patient Care Rooms requires the physical integrity of each receptacle shall be confirmed by visual inspection. The continuity of the grounding circuit in each electrical receptacle shall be verified. Correct polarity of the hot and neutral connections in each electrical receptacle shall be confirmed; and retention force of the grounding blade of each electrical receptacle (except locking-type receptacles) shall be not less than 115 grams (4 ounces). This deficient practice could affect as many as 25 residents, 4 staff, and 2 visitors within the facility.</p> <p>Findings include:</p> <p>Based on record review on 05/29/24 with the Maintenance Director at 11:00 a.m., when asked to review the annual receptacle retention</p>			K 0914	<p>K914– It is the intent of the facility to ensure all non-hospital grade electrical receptacles at resident room locations are tested at least annually to meet set standards.</p> <p>1 CORRECTIVE ACTIONS TAKEN:</p> <p>a On 6/3/2024 the Maintenance Supervisor/designee conducted the annual electrical receptacle inspection and documented the results in the facilities Life safety binder to meet set standards. The Administrator verified repairs on 6/3/2024.</p> <p>2 ALL OTHERS WITH POTENTIAL TO BE AFFECTED:</p> <p>a All residents and all staff and visitors have the potential to be affected but none were.</p> <p>3 MEASURES TO PREVENT REOCCURRENCE:</p> <p>a On 6/3/2024 the Administrator in serviced the Maintenance Supervisor/designee on the requirement the annual electrical receptacle inspection and testing must be completed annually and documented in the</p>		06/03/2024

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	<p>documentation, the Maintenance Director stated that he had not completed all the resident rooms stating that he was only half finished. The last completed receptacle retention test was conducted on 04/03/23 and was more than twelve months old. During a tour of the facility from 11:02 a.m. to 1:45 p.m., the facility's 50 resident rooms had roughly four to six electrical receptacles in each room. Based on interview at the time of the observation, the Maintenance Director indicated all of the electrical receptacles in the resident rooms were not hospital-grade and also indicated there was no up to date documentation of annual testing per NFPA 99, Receptacle Testing requirements for review.</p> <p>This item was again discussed at the exit conference with the facility Administrator and the Maintenance Director on 05/29/24 at 1:55 p.m.</p> <p>3.1-19(b)</p>				<p>life safety binder to meet set standards.</p> <p>b Maintenance Supervisor/designee will ensure the annual electrical receptacle inspection and testing is completed and documented as a part of the facility's Preventive Maintenance Program and document those inspection results as appropriate. If any issues are discovered, they will be addressed and resolved immediately. The Maintenance Supervisor/designee will review with the Administrator the inspection results.</p> <p>c The Administrator will monitor adherence to the Preventative Maintenance schedule and validate the Preventative Maintenance documentation is in place.</p> <p>4 MONITORING</p> <p>CORRECTIVE ACTION:</p> <p>a The inspection results will be presented by the Maintenance Supervisor/designee to the Administrator monthly and the Administrator will present the inspection results at the monthly Quality Assurance/Performance Improvement (QA/PI) meeting. Inspection results and system components will be reviewed by the QA/PI Committee with subsequent plans of correction developed and implemented as deemed necessary to ensure compliance is maintained.</p> <p>This plan of correction</p>		

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