

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

PRINTED: 08/16/2019
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 535034	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED 07/25/2019
NAME OF PROVIDER OR SUPPLIER WESTWARD HEIGHTS CARE CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 150 CARING WAY LANDER, WY 82520		
(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) COMPLETION DATE
K 000	INITIAL COMMENTS A Life Safety Code Survey was conducted by Healthcare Licensing and Surveys on July 25th, 2019. Requirements for Long Term Care Facilities Section 42 CFR 483.70 (a) except as otherwise provided in the section, the facility must meet the applicable provisions of the 2012 Edition of the Life Safety Code, Existing Health Care, of the National Fire Protection Association. The facility was a fully sprinklered, one story building of Type V (111) construction built in 1978 with two additions built in 1996. The building was equipped with a supervised automatic wet sprinkler system with an anti-freeze branch and an addressable fire alarm system. The facility had a capacity of 60 certified Medicare and Medicaid beds with a census of 50 residents.	K 000			
K 920 SS=E	Electrical Equipment - Power Cords and Extens CFR(s): NFPA 101 Electrical Equipment - Power Cords and Extension Cords Power strips in a patient care vicinity are only used for components of movable patient-care-related electrical equipment (PCREE) assemblies that have been assembled by qualified personnel and meet the conditions of 10.2.3.6. Power strips in the patient care vicinity may not be used for non-PCREE (e.g., personal electronics), except in long-term care resident rooms that do not use PCREE. Power strips for PCREE meet UL 1363A or UL 60601-1. Power strips for non-PCREE in the patient care rooms (outside of vicinity) meet UL 1363. In non-patient	K 920			8/23/19

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

TITLE

(X6) DATE

Electronically Signed

08/14/2019

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.

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K 920	Continued From page 1 care rooms, power strips meet other UL standards. All power strips are used with general precautions. Extension cords are not used as a substitute for fixed wiring of a structure. Extension cords used temporarily are removed immediately upon completion of the purpose for which it was installed and meets the conditions of 10.2.4. 10.2.3.6 (NFPA 99), 10.2.4 (NFPA 99), 400-8 (NFPA 70), 590.3(D) (NFPA 70), TIA 12-5 This REQUIREMENT is not met as evidenced by: Based on observation and staff interview, the facility failed to properly use power strips in accordance with the 2012 NFPA 99, Health Care Facilities Code. Failure to properly use power strips could result in injury or death. The deficiency affected two (2) of three (3) smoke compartments. The findings were: Observation on 07/25/19 at 12:15 AM and throughout the survey revealed resident rooms 110, 117 and 206 had Patient Care Related Electrical Equipment located in the room with a power strip located in the patient care area, and resident room 206 had the PCREE plugged directly into the power strip. Resident rooms utilizing patient care related electrical equipment shall not have a power strip. Interview with the maintenance director at the time of the observation acknowledged the deficiency, and indicated they were unaware of the requirement. Interview with the director of nursing at the time of exit acknowledged the deficiency.	K 920	Correct deficiency to individual: Deficiency to resident living in 110 was corrected by moving the non-patient care items out of the healthcare area and also providing an approved power strip for the personal electronics. Deficiency to residents living in 117 was corrected by relocated non-patient care items out of the healthcare area and to the opposite side of the room. Deficiency to residents living in 206 was corrected by offering one of the residents a room move so that more outlets were available for the healthcare equipment and thus allowing the non-patient care equipment to be plugged in outside of the healthcare area. How others are identified in similar situations: The maintenance supervisor or designee will complete 100% audit of each room, identifying non-healthcare items plugged in the healthcare area as well as identifying any unapproved power strips		

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K 920	Continued From page 2	K 920	and replacing them immediately with an appropriate power strip. Measures/systemic changes to prevent recurrence: Education will be provided by Administrator or designee to all team members who work in the resident rooms related to the proper use of outlets in the healthcare area and also the use of approved power strips for non-healthcare items outside of the patient care area. Weekly room audits will be completed each week for the next 60 days and prn thereafter by the maintenance supervisor or designee to check for healthcare items being plugged into appropriate receptacles, identifying any healthcare equipment that is plugged into a power strip, and checking for appropriate power strips. Education will be provided on the spot and the items will be corrected at the time they are discovered. The results of the audits will be reported at monthly Safety Meetings by the Maintenance Supervisor or designee		
K 923 SS=D	Gas Equipment - Cylinder and Container Storage CFR(s): NFPA 101 Gas Equipment - Cylinder and Container Storage Greater than or equal to 3,000 cubic feet Storage locations are designed, constructed, and ventilated in accordance with 5.1.3.3.2 and 5.1.3.3.3. >300 but <3,000 cubic feet Storage locations are outdoors in an enclosure or within an enclosed interior space of non- or	K 923			8/23/19

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K 923	Continued From page 3 limited- combustible construction, with door (or gates outdoors) that can be secured. Oxidizing gases are not stored with flammables, and are separated from combustibles by 20 feet (5 feet if sprinklered) or enclosed in a cabinet of noncombustible construction having a minimum 1/2 hr. fire protection rating. Less than or equal to 300 cubic feet In a single smoke compartment, individual cylinders available for immediate use in patient care areas with an aggregate volume of less than or equal to 300 cubic feet are not required to be stored in an enclosure. Cylinders must be handled with precautions as specified in 11.6.2. A precautionary sign readable from 5 feet is on each door or gate of a cylinder storage room, where the sign includes the wording as a minimum "CAUTION: OXIDIZING GAS(ES) STORED WITHIN NO SMOKING." Storage is planned so cylinders are used in order of which they are received from the supplier. Empty cylinders are segregated from full cylinders. When facility employs cylinders with integral pressure gauge, a threshold pressure considered empty is established. Empty cylinders are marked to avoid confusion. Cylinders stored in the open are protected from weather. 11.3.1, 11.3.2, 11.3.3, 11.3.4, 11.6.5 (NFPA 99) This REQUIREMENT is not met as evidenced by: Based on observation and staff interview, the facility failed to protect oxygen storage in accordance with the 2012 NFPA 99, Health Care Facilities Code. Failure to properly protect oxygen storage could result in injury or death in the event of a fire. The deficiency affected one (1) of three (3) smoke compartments. The findings were:	K 923	Correct deficiency to individual: In order to correct the deficiency to all individuals, a keypad locking system will be installed on both oxygen storage rooms in the smoke compartment. How others are identified and correction to others in similar situations: There are no other oxygen storage rooms		

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K 923	Continued From page 4 Observation on 07/25/19 at 12:56 PM revealed that the oxygen storage room contained 2,250 cubic feet of stored oxygen in cylinders and the room was incapable of being locked. Oxygen storage locations shall have doors that can be secured against unauthorized entry. Interview with the maintenance director at the time of the observation acknowledged the deficiency, and indicated they were unaware of the requirement. Interview with the administrator at the time of exit acknowledged the deficiency. Reference: 2012 NFPA 99 11.3.2.1			K 923	in the facility. Measures/systemic changes to prevent reoccurrence: Education to be provided by the Administrator or designee to all facility leadership regarding this regulation of having ability to secure the oxygen storage rooms with greater than 300 cubic feet and less than 3000 cubic feet. Monitor permanent solution: Maintenance supervisor or designee will provide weekly checks 60 day and then monthly checks of the keypad lock to ensure the oxygen storage rooms are secure. The checks will be reported at the monthly Safety Meeting by the Maintenance Supervisor or designee.		

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E 000	Initial Comments An Emergency Preparedness survey was conducted by Healthcare Licensing and Surveys on July 25, 2019. Based upon the findings of the survey team, it was determined that the facility was in compliance with all requirements.	E 000			

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