

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

PRINTED: 08/24/2018
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 535025	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED 07/10/2018
NAME OF PROVIDER OR SUPPLIER CHEYENNE HEALTH CARE CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 2700 E 12TH STREET CHEYENNE, WY 82001		
(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 10, 2018. Requirements for Long Term Care Facilities Section 42 CFR 483.70 (a) except as otherwise provided in this 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 (000) construction built in 1961. The building was equipped with a supervised, automatic wet and anti-freeze sprinkler system with an addressable fire alarm system. The facility had a capacity of 105 certified Medicare and Medicaid beds with a census of 96 residents.	K 000			
K 920 SS=D	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 care rooms, power strips meet other UL	K 920		8/13/18	

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

TITLE

(X6) DATE

Electronically Signed

08/09/2018

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 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 maintain electrical systems in accordance with the 2012 NFPA 101, Life Safety Code, and the 2011 NFPA 70, National Electrical Code. Failure to maintain electrical systems as required could cause a fire resulting in injury or death. The deficiency affected one (1) of the (96) residents. The findings were: Observation on 7/10/2018 at 10:41 AM in room 106 revealed an oxygen concentrator plugged into a power strip located under a resident's bed. Further observation revealed that the UL listing of the power strip could not be determined. Interview with the facility maintenance supervisor at the time of observation acknowledged the deficiency, and indicated he was aware of the requirement. Interview with the facility administrator at the time of exit acknowledged the deficiency. REF: 2012 NFPA 101, Sections: 19.5.1.1; 9.1.2 2011 NFPA 70, Section: 517.18(b)(2)	K 920	Corrective Action The concentrator was unplugged from the power strip on 7/10/18 and plugged into an outlet in the wall. The power strip was replaced with a power strip that was verified as UL listed. Identification of Others An audit was conducted on 7/27/18 of all resident rooms to see if any other concentrators were plugged into power strips and to verify that each power strip in the facility was UL listed. Systemic Change In-service education was provided to facility staff and completed prior to 8/9/18 regarding not plugging in concentrators into power strips. In-service training was conducted with the Maintenance Staff on 8/9/18 regarding the need for any power strip in the facility to be UL listed.		

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K 920	Continued From page 2	K 920	Monitoring The Maintenance Director/designee audits all rooms to ensure that no concentrators are plugged into power strips and verifies that any power strips in use for other items are UL listed. These audits will be conducted weekly for 4 weeks and then monthly for 2 additional months. The Maintenance Director will submit the findings from the audits to the monthly QAPI Committee meeting for review and any further recommendations.		

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E 000	Initial Comments An Emergency Preparedness survey was conducted by Healthcare Licensing and Surveys on July 10, 2018. Based upon the findings of the survey team, it was determined the facility was in compliance with all requirements.	E 000			

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07/31/2018

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