



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered
August 7, 2023

Administrator
Gundersen Harmony Care Center
815 Main Avenue South
Harmony, MN 55939

RE: CCN: 245528
Cycle Start Date: May 11, 2023

Dear Administrator:

On July 13, 2023, the Minnesota Departments of Health and Public Safety completed a revisit to verify that your facility had achieved and maintained compliance. Based on our review, we have determined that your facility has achieved substantial compliance; therefore no remedies will be imposed.

Feel free to contact me if you have questions.

A handwritten signature in black ink, appearing to read 'Melissa Poepping'.

Melissa Poepping, Compliance Analyst
Federal Enforcement | Health Regulation Division
Minnesota Department of Health
P.O. Box 64900
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: Melissa.Poepping@state.mn.us



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
June 22, 2023

Administrator
Gundersen Harmony Care Center
815 Main Avenue South
Harmony, MN 55939

RE: CCN: 245528
Cycle Start Date: May 11, 2023

Dear Administrator:

On May 11, 2023, a survey was completed at your facility by the Minnesota Departments of Health and Public Safety, to determine if your facility was in compliance with Federal participation requirements for skilled nursing facilities and/or nursing facilities participating in the Medicare and/or Medicaid programs.

This survey found the most serious deficiencies in your facility to be widespread deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level F), as evidenced by the electronically attached CMS-2567 whereby corrections are required.

ELECTRONIC PLAN OF CORRECTION (ePoC)

Within **ten (10) calendar days** after your receipt of this notice, you must submit an acceptable ePOC for the deficiencies cited. An acceptable ePOC will serve as your allegation of compliance. Upon receipt of an acceptable ePOC, we will authorize a revisit to your facility to determine if substantial compliance has been achieved.

To be acceptable, a provider's ePOC must include the following:

- How corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- How the facility will identify other residents having the potential to be affected by the same deficient practice.
- What measures will be put into place, or systemic changes made, to ensure that the deficient practice will not recur.
- How the facility will monitor its corrective actions to ensure that the deficient practice is being corrected and will not recur.
- The date that each deficiency will be corrected.
- An electronic acknowledgement signature and date by an official facility representative.

The state agency may, in lieu of an onsite revisit, determine correction and compliance by accepting the facility's ePoC if the ePoC is reasonable, addresses the problem and provides evidence that the corrective action has occurred.

If an acceptable ePoC is not received within 10 calendar days from the receipt of this letter, we will recommend to the CMS Region V Office that one or more of the following remedies be imposed:

- Denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417);
- Civil money penalty (42 CFR 488.430 through 488.444).
- Termination of your facility's Medicare and/or Medicaid agreement (488.456(b)).

DEPARTMENT CONTACT

Questions regarding this letter and all documents submitted as a response to the resident care deficiencies (those preceded by an "F" and/or an "E" tag), i.e., the plan of correction should be directed to:

Judy Loecken, Unit Supervisor
St. Cloud B District Office
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Midtown Square
3333 Division Street, Suite 212
Saint Cloud, Minnesota 56301-4557
Email: judy.loecken@state.mn.us
Office: (320) 223-7300 Mobile: (320) 241-7797

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

The facility's ePoC will serve as your allegation of compliance upon the Department's acceptance. In order for your allegation of compliance to be acceptable to the Department, the ePoC must meet the criteria listed in the plan of correction section above. You will be notified by the Minnesota Department of Health, Licensing and Certification Program staff and/or the Department of Public Safety, State Fire Marshal Division staff, if your ePoC for the respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

Upon receipt of an acceptable ePoC, a Post Certification Revisit (PCR), of your facility will be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification.

If substantial compliance has been achieved, certification of your facility in the Medicare and/or Medicaid program(s) will be continued and remedies will not be imposed. Compliance is certified as of

the latest correction date on the approved ePoC, unless it is determined that either correction actually occurred between the latest correction date on the ePoC and the date of the first revisit, or correction occurred sooner than the latest correction date on the ePoC.

FAILURE TO ACHIEVE SUBSTANTIAL COMPLIANCE BY THE THIRD OR SIXTH MONTH AFTER THE LAST DAY OF THE SURVEY

If substantial compliance with the regulations is not verified by August 11, 2023 (three months after the identification of noncompliance), the CMS Region V Office must deny payment for new admissions as mandated by the Social Security Act (the Act) at Sections 1819(h)(2)(D) and 1919(h)(2)(C) and Federal regulations at 42 CFR Section 488.417(b).

In addition, if substantial compliance with the regulations is not verified by November 11, 2023 (six months after the identification of noncompliance) your provider agreement will be terminated. This action is mandated by the Social Security Act at Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

Please note that this notice does not constitute formal notice of imposition of alternative remedies or termination of your provider agreement. Should the Centers for Medicare & Medicaid Services determine that termination or any other remedy is warranted, it will provide you with a separate formal notification of that determination.

INFORMAL DISPUTE RESOLUTION (IDR) / INDEPENDENT INFORMAL DISPUTE RESOLUTION (IIDR)

In accordance with 42 CFR 488.331, you have one opportunity to question cited deficiencies through an informal dispute resolution process. You are required to send your written request, along with the specific deficiencies being disputed, and an explanation of why you are disputing those deficiencies, to:

Nursing Home Informal Dispute Process
Minnesota Department of Health
Health Regulation Division
P.O. Box 64900
St. Paul, Minnesota 55164-0900

This request must be sent within the same ten days you have for submitting an ePoC for the cited deficiencies. All requests for an IDR or IIDR of federal deficiencies must be submitted via the web at: https://mdhprovidercontent.web.health.state.mn.us/lrc_idr.cfm

You must notify MDH at this website of your request for an IDR or IIDR within the 10 calendar day period allotted for submitting an acceptable electronic plan of correction. A copy of the Department's informal dispute resolution policies are posted on the MDH Information Bulletin website at: https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html

Please note that the failure to complete the informal dispute resolution process will not delay the dates specified for compliance or the imposition of remedies.

Gundersen Harmony Care Center

June 22, 2023

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Questions regarding all documents submitted as a response to the Life Safety Code deficiencies (those preceded by a "K" tag), i.e., the plan of correction, request for waivers, should be directed to:

William Abderhalden, Fire Safety Supervisor
Deputy State Fire Marshal
Health Care/Corrections Supervisor – Interim
Minnesota Department of Public Safety
445 Minnesota Street, Suite 145
St. Paul, MN 55101-5145
Cell: (507) 361-6204
Email: william.abderhalden@state.mn.us
Fax: (651) 215-0525

Feel free to contact me if you have questions.

Sincerely,



Kamala Fiske-Downing
Minnesota Department of Health
Health Regulation Division
Telephone: (651) 201-4112
Email: Kamala.Fiske-Downing@state.mn.us

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

PRINTED: 07/07/2023
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245528	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 05/11/2023
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NAME OF PROVIDER OR SUPPLIER GUNDERSEN HARMONY CARE CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 815 MAIN AVENUE SOUTH HARMONY, MN 55939
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(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
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E 000	<p>Initial Comments</p> <p>On 5/8/23 through 5/11/23, a survey for compliance with Appendix Z, Emergency Preparedness Requirements, §483.73(b)(6) was conducted during a standard recertification survey. The facility was NOT in compliance.</p> <p>The facility's plan of correction (POC) will serve as your allegation of compliance upon the Department's acceptance. Because you are enrolled in ePOC, your signature is not required at the bottom of the first page of the CMS-2567 form.</p> <p>Upon receipt of an acceptable electronic POC, an onsite revisit of your facility may be conducted to validate substantial compliance with the regulation has been attained.</p>	E 000		
E 041 SS=F	<p>Hospital CAH and LTC Emergency Power CFR(s): 483.73(e)</p> <p>§482.15(e) Condition for Participation: (e) Emergency and standby power systems. The hospital must implement emergency and standby power systems based on the emergency plan set forth in paragraph (a) of this section and in the policies and procedures plan set forth in paragraphs (b)(1)(i) and (ii) of this section.</p> <p>§483.73(e), §485.625(e), §485.542(e) (e) Emergency and standby power systems. The [LTC facility CAH and REH] must implement emergency and standby power systems based on the emergency plan set forth in paragraph (a) of this section.</p> <p>§482.15(e)(1), §483.73(e)(1), §485.542(e)(1), §485.625(e)(1)</p>	E 041		6/26/23

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 06/26/2023
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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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E 041	<p>Continued From page 1</p> <p>Emergency generator location. The generator must be located in accordance with the location requirements found in the Health Care Facilities Code (NFPA 99 and Tentative Interim Amendments TIA 12-2, TIA 12-3, TIA 12-4, TIA 12-5, and TIA 12-6), Life Safety Code (NFPA 101 and Tentative Interim Amendments TIA 12-1, TIA 12-2, TIA 12-3, and TIA 12-4), and NFPA 110, when a new structure is built or when an existing structure or building is renovated.</p> <p>482.15(e)(2), §483.73(e)(2), §485.625(e)(2), §485.542(e)(2) Emergency generator inspection and testing. The [hospital, CAH and LTC facility] must implement the emergency power system inspection, testing, and [maintenance] requirements found in the Health Care Facilities Code, NFPA 110, and Life Safety Code.</p> <p>482.15(e)(3), §483.73(e)(3), §485.625(e)(3), §485.542(e)(2) Emergency generator fuel. [Hospitals, CAHs and LTC facilities] that maintain an onsite fuel source to power emergency generators must have a plan for how it will keep emergency power systems operational during the emergency, unless it evacuates.</p> <p>*[For hospitals at §482.15(h), LTC at §483.73(g), REHs at §485.542(g), and and CAHs §485.625(g):] The standards incorporated by reference in this section are approved for incorporation by reference by the Director of the Office of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. You may obtain the material from the sources listed below. You may</p>	E 041		

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E 041	<p>Continued From page 2</p> <p>inspect a copy at the CMS Information Resource Center, 7500 Security Boulevard, Baltimore, MD or at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call 202-741-6030, or go to: http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html. If any changes in this edition of the Code are incorporated by reference, CMS will publish a document in the Federal Register to announce the changes.</p> <p>(1) National Fire Protection Association, 1 Batterymarch Park, Quincy, MA 02169, www.nfpa.org, 1.617.770.3000.</p> <p>(i) NFPA 99, Health Care Facilities Code, 2012 edition, issued August 11, 2011.</p> <p>(ii) Technical interim amendment (TIA) 12-2 to NFPA 99, issued August 11, 2011.</p> <p>(iii) TIA 12-3 to NFPA 99, issued August 9, 2012.</p> <p>(iv) TIA 12-4 to NFPA 99, issued March 7, 2013.</p> <p>(v) TIA 12-5 to NFPA 99, issued August 1, 2013.</p> <p>(vi) TIA 12-6 to NFPA 99, issued March 3, 2014.</p> <p>(vii) NFPA 101, Life Safety Code, 2012 edition, issued August 11, 2011.</p> <p>(viii) TIA 12-1 to NFPA 101, issued August 11, 2011.</p> <p>(ix) TIA 12-2 to NFPA 101, issued October 30, 2012.</p> <p>(x) TIA 12-3 to NFPA 101, issued October 22, 2013.</p> <p>(xi) TIA 12-4 to NFPA 101, issued October 22, 2013.</p> <p>(xiii) NFPA 110, Standard for Emergency and Standby Power Systems, 2010 edition, including TIAs to chapter 7, issued August 6, 2009..</p> <p>This REQUIREMENT is not met as evidenced by:</p>	E 041		

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E 041	<p>Continued From page 3</p> <p>Based on observation, review of available documentation and staff interview, the facility failed to test the on-site emergency generator system per NFPA 99 (2012 edition), Health Care Facilities Code, section 6.4.4.1.1.3, 6.4.4.2 and NFPA 110 (2010 edition), Standard for Emergency and Standby Power Systems, 8.4.1, 8.4.2 This deficient finding could have a widespread impact on the residents within the facility.</p> <p>Findings include:</p> <ol style="list-style-type: none"> 1. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by a review of available documentation, that the most recent documented weekly generator inspection was dated as being completed on 03/23/2023. 2. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by a review of available documentation, that the most recent documented monthly generator inspection was dated as being completed in February 2023. <p>An interview with the Maintenance Director verified these deficient findings at the time of discovery.</p>	E 041	<p>E041 Gundersen Harmony Care Center will continue to implement the emergency power system inspection, testing and [maintenance] requirements found in the Health Care Facilities Code, NFPA 11- and Life Safety Code. The Facilities Mechanic was re-educated to the need to complete weekly and monthly inspections. Along with this the facilities mechanic added to his calendar a reminder to complete the weekly inspection of the emergency generator and log this in his logbook. With this, the Facilities Mechanic added to his calendar a reminder to complete the monthly generator run test and log this in his logbook. Administrator will audit weekly x 1 month and then monthly x 6 months. Results will be reported to the QAA committee monthly. Date Completed: 6.26.23</p>	
F 000	<p>INITIAL COMMENTS</p> <p>On 5/8/23 through 5/11/23, a standard recertification survey was conducted at your facility. A complaint investigation was also conducted. Your facility was IN compliance with the requirements of 42 CFR 483, Subpart B, Requirements for Long Term Care Facilities.</p> <p>In addition to the recertification survey, the</p>	F 000		

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F 000	<p>Continued From page 4 following complaints were reviewed</p> <p>The following complaints were reviewed with no deficiency issued. H55282048C (MN89678), H5528020C (MN78572), H55282011C (MN91980), H55284618C (MN86737), H5528019C (MN80692)</p> <p>The facility is enrolled in ePOC, therefore a signature is not required at the bottom of the first page of the CMS-2567 form. Although no plan of correction is required, the facility must acknowledge receipt of the electronic documents.</p>	F 000		

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

F5528033

PRINTED: 07/20/2023
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245528	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING B. WING _____	(X3) DATE SURVEY COMPLETED 05/09/2023
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NAME OF PROVIDER OR SUPPLIER GUNDERSEN HARMONY CARE CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 815 MAIN AVENUE SOUTH HARMONY, MN 55939
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K 000	<p>INITIAL COMMENTS</p> <p>FIRE SAFETY</p> <p>An annual Life Safety Code survey was conducted by the Minnesota Department of Public Safety, State Fire Marshal Division on 05/09/2023. At the time of this survey, GUNDERSON HARMONY CARE CENTER was found not in compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2012 edition of National Fire Protection Association (NFPA) 101, Life Safety Code (LSC), Chapter 19 Existing Health Care and the 2012 edition of NFPA 99, Health Care Facilities Code.</p> <p>THE FACILITY'S POC WILL SERVE AS YOUR ALLEGATION OF COMPLIANCE UPON THE DEPARTMENT'S ACCEPTANCE. YOUR SIGNATURE AT THE BOTTOM OF THE FIRST PAGE OF THE CMS-2567 FORM WILL BE USED AS VERIFICATION OF COMPLIANCE.</p> <p>UPON RECEIPT OF AN ACCEPTABLE POC, AN ONSITE REVISIT OF YOUR FACILITY MAY BE CONDUCTED TO VALIDATE THAT SUBSTANTIAL COMPLIANCE WITH THE REGULATIONS HAS BEEN ATTAINED IN ACCORDANCE WITH YOUR VERIFICATION.</p> <p>PLEASE RETURN THE PLAN OF CORRECTION FOR THE FIRE SAFETY DEFICIENCIES (K-TAGS) TO:</p> <p>IF PARTICIPATING IN THE E-POC PROCESS, A PAPER COPY OF THE PLAN OF CORRECTION IS NOT REQUIRED.</p>	K 000		
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LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 06/26/2023
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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 000	<p>Continued From page 1</p> <p>Healthcare Fire Inspections State Fire Marshal Division 445 Minnesota St., Suite 145 St. Paul, MN 55101-5145, OR</p> <p>By email to: FM.HC.Inspections@state.mn.us</p> <p>THE PLAN OF CORRECTION FOR EACH DEFICIENCY MUST INCLUDE ALL OF THE FOLLOWING INFORMATION:</p> <ol style="list-style-type: none"> 1. A detailed description of the corrective action taken or planned to correct the deficiency. 2. Address the measures that will be put in place to ensure the deficiency does not reoccur. 3. Indicate how the facility plans to monitor future performance to ensure solutions are sustained. 4. Identify who is responsible for the corrective actions and monitoring of compliance. 5. The actual or proposed date for completion of the remedy. <p>GUNDERSON HARMONY CARE CENTER is a 1 story building with no basement.</p> <p>The building was constructed at 2 different times. The original building was constructed in 1963 and was determined to be of Type II (111) construction. In 1967 an addition was constructed and was determined to be of Type II(111) construction.</p>	K 000		

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K 000	Continued From page 2 Because the original building and addition meet the construction type allowed for existing buildings, the facility was surveyed as one building as allowed in the 2012 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC), Chapter 19 Existing Health Care Occupancies. The facility is fully protected throughout by an automatic sprinkler system and has a fire alarm system with smoke detection in the corridors, spaces open to the corridors, that is monitored for automatic fire department notification. There are two occupancies in the building. The nursing home (I-2) and an outpatient clinic (B) with proper fire separation. The facility has a capacity of 43 beds and had a census of 30 at the time of the survey.	K 000		
K 291 SS=C	The requirement at 42 CFR, Subpart 483.70(a) is NOT MET as evidenced by: Emergency Lighting CFR(s): NFPA 101 Emergency Lighting Emergency lighting of at least 1-1/2-hour duration is provided automatically in accordance with 7.9.18.2.9.1, 19.2.9.1 This REQUIREMENT is not met as evidenced by: Based on a review of available documentation and staff interview, the facility failed to maintain emergency lighting testing documentation per NFPA 101 (2012 edition), Life Safety Code, section 19.2.9.1, 7.9, and 7.9.3.1.1(5). This deficient finding could have a widespread impact	K 291	K291 Gundersen Harmony Care Center will continue to ensure emergency lighting of at least 1 ½ hour duration is provided automatically in accordance with 7.9. The Facilities Mechanic along with the facility's electrical vendor completed the annual 90	6/12/23

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K 291	Continued From page 3 on the residents within the facility. Findings include: On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by review of available documentation that it was unclear as to who completed the testing, and if annual 90 min. testing occurred. An interview with Maintenance Director verified this deficient finding at the time of discovery.	K 291	minute emergency lighting test on 6.12.23 and all emergency lights passed. A calendar reminder was added to the Facilities Mechanic calendar to ensure completion moving forward annually and to the Administrator's calendar for auditing purposes. Date Completed: 6.12.23	
K 355 SS=F	Portable Fire Extinguishers CFR(s): NFPA 101 Portable Fire Extinguishers Portable fire extinguishers are selected, installed, inspected, and maintained in accordance with NFPA 10, Standard for Portable Fire Extinguishers. 18.3.5.12, 19.3.5.12, NFPA 10 This REQUIREMENT is not met as evidenced by: Based on observation, review of available documentation and staff interview, the facility failed to properly inspect, and maintain documentation of portable fire extinguishers in accordance with NFPA 101 (2012 edition), Life Safety Code, sections 19.3.5.12, 9.7.4.1, and NFPA 10 (2010 edition), Standard for Portable Fire Extinguishers, section 7.2.4.3, 7.2.4.4, 7.2.4.5. These deficient findings could have a widespread impact on the residents within the facility. Findings include: On 05/09/2023 between 10:00 AM and 2:00 PM,	K 355	K355 Gundersen Harmony care center will continue to ensure that portable fire extinguishers are selected, installed, inspected, and maintained in accordance with NFPA 10, Standard for Portable Fire Extinguishers 18.3.5.12, 19.3.5.12, NFPA 10. The facilities mechanic had completed his monthly inspections of the fire extinguishers and logged this in his logbook. However, being new in his position he overlooked signing on the tags. Thus, he was re-educated on the need to sign off on the tag as well. Administrator will audit monthly x 6 months. Results will be reported to the	6/26/23

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K 355	Continued From page 4 it was revealed by observation, that the tags on the fire extinguishers were missing multiple inspection dates and initials of staff conducting the inspection. An interview with the Maintenance Director verified this deficient finding at the time of discovery.	K 355	QAA committee monthly. Date Completed: 6.26.23	
K 712 SS=F	Fire Drills CFR(s): NFPA 101 Fire Drills Fire drills include the transmission of a fire alarm signal and simulation of emergency fire conditions. Fire drills are held at expected and unexpected times under varying conditions, at least quarterly on each shift. The staff is familiar with procedures and is aware that drills are part of established routine. Where drills are conducted between 9:00 PM and 6:00 AM, a coded announcement may be used instead of audible alarms. 19.7.1.4 through 19.7.1.7 This REQUIREMENT is not met as evidenced by: Based on a review of available documentation and staff interview, the facility failed to conduct fire drills per NFPA 101 (2012 edition), Life Safety Code, sections 19.7.1, and 19.7.1.6. This deficient finding could have a widespread impact on the residents within the facility. Findings include: On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by a review of available documentation, that no documentation was presented to confirm that the monthly fire drills	K 712	K712: Gundersen Harmony Care Center will continue to ensure that fire drills include the transmission of a fire alarm signal and simulation of emergency fire conditions. Fire drills are held at expected and unexpected times under varying conditions, at least quarterly on each shift. The staff is familiar with procedures and is aware that drills are part of established routine. Where drills are conducted between 9:00 PM and 6:00 AM, a coded announcement may be used instead of audible alarms. Moving forward,	6/26/23

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K 712	Continued From page 5 had been completed for: 1st shift - second, third and fourth quarters; 2nd shift - first and third quarters; 3rd shift - second, third, and fourth quarters. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 712	Gundersen Harmony Care Center will ensure that monthly fire drills are conducted and documented. The facilities mechanic was re-educated on K712 and the need for a monthly fire drill. Administrator will conduct an audit monthly x 6 months of ensuring fire drills are completed monthly. Results will be reported to the QAA committee monthly. Completion Date: 6.26.23	
K 918 SS=F	Electrical Systems - Essential Electric Syste CFR(s): NFPA 101 Electrical Systems - Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of	K 918		6/26/23

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K 918	<p>Continued From page 6</p> <p>maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations.</p> <p>6.4.4, 6.5.4, 6.6.4 (NFPA 99), NFPA 110, NFPA 111, 700.10 (NFPA 70)</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on observation, review of available documentation and staff interview, the facility failed to test the on-site emergency generator system per NFPA 99 (2012 edition), Health Care Facilities Code, section 6.4.4.1.1.3, 6.4.4.2 and NFPA 110 (2010 edition), Standard for Emergency and Standby Power Systems, 8.4.1, 8.4.2 This deficient finding could have a widespread impact on the residents within the facility.</p> <p>Findings include:</p> <ol style="list-style-type: none"> 1. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by a review of available documentation, that the most recent documented weekly generator inspection was dated as being completed on 03/23/2023. 2. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by a review of available documentation, that the most recent documented monthly generator inspection was dated as being completed in February 2023. <p>An interview with the Maintenance Director verified these deficient findings at the time of discovery.</p>	K 918	<p>K918: Gundersen Harmony Care Center will continue to implement the emergency power system inspection, testing and [maintenance] requirements found in the Health Care Facilities Code, NFPA 11- and Life Safety Code. The Facilities Mechanic was re-educated to the need to complete weekly and monthly inspections. Along with this the facilities mechanic added to his calendar a reminder to complete the weekly inspection of the emergency generator and log this in his logbook. With this, the Facilities Mechanic added to his calendar a reminder to complete the monthly generator run test and log this in his logbook. Administrator will audit weekly x 1 month and then monthly x 6 months. Results will be reported to the QAA committee monthly. Date Completed: 6.26.23</p>	

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K 920 SS=D	<p>Electrical Equipment - Power Cords and Extens CFR(s): NFPA 101</p> <p>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 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 manage usage of relocatable power taps in accordance with NFPA 99 (2012 edition), Health Care Facilities Code, section 10.2.3.6, and NFPA 70, (2011 edition), National Electrical Code, sections 110.3(B), 400.8 (1) and UL 1363. This deficient finding could have an isolated impact on the residents within the facility.</p> <p>Findings include:</p>	K 920	<p>K920 Gundersen Harmony Care Center will continue to ensure that power strips in non-patient care rooms 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. The facilities mechanic</p>	6/26/23

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K 920	Continued From page 8 On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by observation, that in the Business Office relocatable power taps were daisy-chained together. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 920	separated the daisy chained cords in the business office and plugged the cords into different outlets. The Facilities Mechanic will complete a monthly walk through monthly x 6 months and then quarterly thereafter a safety walkthrough will be completed by members of the leadership team to ensure extension cords are not being used improperly. Results will be reported to the QAA committee monthly. Date Completed: 6.26.23		
K 923 SS=E	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 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,	K 923		6/23/23	

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K 923	<p>Continued From page 9</p> <p>where the sign includes the wording as a minimum "CAUTION: OXIDIZING GAS(ES) STORED WITHIN NO SMOKING."</p> <p>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:</p> <p>Based on observation and staff interview, the facility failed to maintain proper medical gas storage and management per NFPA 99 (2012 edition), Health Care Facilities Code, sections 11.6.5, 11.6.5.2, 11.6.5.3. This deficient finding could have a patterned impact on the residents within the facility.</p> <p>Findings include:</p> <ol style="list-style-type: none"> 1. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by observation that in the Med Gas Storage Room #1, there were cylinders that were missing empty / full hanging identifier tags. Missing these indicators there was no way to visually assess whether cylinders were full or empty. 2. On 05/09/2023 between 10:00 AM and 2:00 PM, it was revealed by observation that in the Med Gas Storage Room #1, there were cylinders that were missing the visual full cylinder indicator consisting of the common white dust cap or the newer white safe wrap. Missing these indicators there was no way to visually assess whether 	K 923	<p>K923 Gundersen Harmony Care Center will continue to ensure that empty cylinders are segregated from full cylinders. Gundersen Harmony Care Center had full cylinders segregated from empty cylinders at the time of the survey. The oxygen vendor had switched from the common white plastic wrap to a white paper wrap and the paper wrap had gotten loose during transportation. Along with this some of the empty/full hanging identifier tags had fallen off as well. The facility requested the oxygen vendor to remove those cylinders with the white paper wrap that had loosened and become displaced on the cylinder. This was completed on 6.23.23. The oxygen vendor was also reminded on 6.23.23 to ensure the empty/full hanging identifier tags are in place on delivery as well. DON or designee will audit segregation of empty and full cylinders to include white plastic wrap being securely on the full cylinders and empty/full identifier tags to be on cylinders as well. This audit will be</p>	

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K 923	Continued From page 10 cylinders were full or empty. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 923	monthly x 6 months. Results will be reported to the QAA committee meeting monthly. Date Completed: 6.23.23		