



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered

October 19, 2023

Administrator
The Green Prairie Rehabilitation Center
800 Second Avenue Northwest
Plainview, MN 55964

RE: CCN: 245345
Cycle Start Date: October 5, 2023

Dear Administrator:

On October 5, 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

The Green Prairie Rehabilitation Center

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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 January 5, 2024, (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 April 5, 2024, (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/ltc_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

The Green Prairie Rehabilitation Center

October 19, 2023

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dates specified for compliance or the imposition of remedies.

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:

Travis Z. Ahrens
Interim State Fire Safety Supervisor
Health Care & Correctional Facilities/Explosives
MN Department of Public Safety-Fire Marshal Division
445 Minnesota St., Suite 145
St. Paul, MN 55101
travis.ahrens@state.mn.us
Cell: 1-507-308-4189

Sincerely,

A handwritten signature in black ink that reads "Lori Hagen". The signature is written in a cursive style with a large, looping initial "L".

Lori Hagen, Compliance Analyst
Federal Enforcement
Health Regulation Division
Minnesota Department of Health
Telephone: 651-201-4306
E-Mail: Lori.Hagen@state.mn.us



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October 19, 2023

Administrator
The Green Prairie Rehabilitation Center
800 Second Avenue Northwest
Plainview, MN 55964

Re: Event ID: CH4F11

Dear Administrator:

The above facility survey was completed on October 5, 2023, for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules. At the time of the survey, the survey team from the Minnesota Department of Health - Health Regulation Division noted no violations of these rules promulgated under Minnesota Stat. section 144.653 and/or Minnesota Stat. Section 144A.10.

Electronically posted is the Minnesota Department of Health order form stating that no violations were noted at the time of this survey. The Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Please disregard the heading of the fourth column which states, "Provider's Plan of Correction." This applies to Federal deficiencies only. There is no requirement to submit a Plan of Correction.

Please contact me with any questions regarding this letter.

Sincerely,

A handwritten signature in black ink that reads 'Lori Hagen'.

Lori Hagen, Compliance Analyst
Federal Enforcement
Health Regulation Division
Minnesota Department of Health
Telephone: 651-201-4306
E-Mail: Lori.Hagen@state.mn.us

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

PRINTED: 11/20/2023
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245345	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 10/05/2023
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964
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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 10/2/23 through 10/5/23, a survey for compliance with Appendix Z, Emergency Preparedness Requirements for Long Term Care facilities, §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),</p>	E 041		10/17/23

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 10/25/2023
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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.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
(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
E 041	<p>Continued From page 1</p> <p>§485.625(e)(1) 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</p>	E 041		

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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
(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
E 041	<p>Continued From page 2</p> <p>material from the sources listed below. You may 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.</p> <p>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>	E 041		

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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
(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	
E 041	Continued From page 3 This REQUIREMENT is not met as evidenced by: Based on a 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, 8.3.4, 8.3.4.1, 8.4.9, 8.4.9.2. These deficient findings could have a widespread impact on the residents within the facility. Findings include: 1. On 09/27/2023 between 10:00 AM and 4:30 PM, it was revealed by a review of available documentation that no documentation was presented for review to confirm that 36-month - 4-hour load bank testing is occurring. 2. On 09/27/2023 between 10:00 AM and 4:30 PM, it was revealed by a review of available documentation that vendor inspection records identified to the facility that emergency generator needed a new air filter. There was no documentation presented to confirm replacement had yet occurred. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	E 041	Based on a review of available documentation and staff interviews, 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, 6.4.42 and NFPA 110 (2010 Edition), Standard for Emergency and Standby Power Systems, 8.3.4, 8.3.4.1, 8.4.9, 8.4.9.2. These deficient findings could have a widespread impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Facility 4 hr load bank testing was completed October 17, 2023. Facility has educated facility Regional Maintenance Director on frequency of testing. Regional Maintenance or designee will report on any upcoming or overdue tests during quarterly QAPI		
F 000	INITIAL COMMENTS On 10/2/23 through 10/5/23, 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,	F 000			

DEPARTMENT OF HEALTH AND HUMAN SERVICES
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
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F 000	<p>Continued From page 4 Requirements for Long Term Care Facilities.</p> <p>The following complaints were reviewed with NO deficiencies cited: H53455947C (MN91311), H53455948C (MN89928)</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		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00672	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 10/05/2023
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTE	STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964
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2 000	<p>Initial Comments</p> <p>*****ATTENTION*****</p> <p>NH LICENSING CORRECTION ORDER</p> <p>In accordance with Minnesota Statute, section 144A.10, this correction order has been issued pursuant to a survey. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a fine for each violation not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health.</p> <p>Determination of whether a violation has been corrected requires compliance with all requirements of the rule provided at the tag number and MN Rule number indicated below. When a rule contains several items, failure to comply with any of the items will be considered lack of compliance. Lack of compliance upon re-inspection with any item of multi-part rule will result in the assessment of a fine even if the item that was violated during the initial inspection was corrected.</p> <p>You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance.</p> <p>INITIAL COMMENTS: On 10/2/23 through 10/5/23, a licensing survey was conducted at your facility by surveyors from the Minnesota Department of Health (MDH). Your facility was IN compliance with the MN State Licensure</p> <p>The following complaints were reviewed during</p>	2 000		
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Minnesota Department of Health LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 10/25/23
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Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00672	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 10/05/2023
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTE	STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964
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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) COMPLETE DATE
2 000	<p>Continued From page 1</p> <p>the survey: H53455947C (MN91311), H53455948C (MN89928)</p> <p>Minnesota Department of Health is documenting the State Licensing Correction Orders using Federal software.</p> <p>The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of state form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.</p>	2 000		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245345	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____	(X3) DATE SURVEY COMPLETED 10/03/2023
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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964
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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 10/03/2023. At the time of this survey, THE GREEN PRAIRIE REHABILITATION 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 10/25/2023
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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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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
(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	<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>THE GREEN PRAIRIE REHABILITATION CENTER is a 2 story building, with partial basement</p> <p>The building was constructed at (3) different times.</p> <p>The original building was constructed in 1968 and was determined to be of Type II (222)</p>	K 000		

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NAME OF PROVIDER OR SUPPLIER THE GREEN PRAIRIE REHABILITATION CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 800 SECOND AVENUE NORTHWEST PLAINVIEW, MN 55964		
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K 000	Continued From page 2 construction. In 1992, addition was constructed to the (Dining Kitchen area) that was determined to be of Type II (222) construction. Another addition was added in 2005 to the chapel area that was determined to be of Type II (222) Because the original building and the (2) addition are of the same type of construction and meet the construction type allowed for existing buildings, the facility was surveyed as one building. The building is protected by a full fire sprinkler system. The facility has a fire alarm system with full corridor smoke detection and spaces open to the corridors that is monitored for automatic fire department notification. The facility has a capacity of 50 beds and had a census of 34 at the time of the survey.	K 000		
K 324 SS=F	The requirement at 42 CFR, Subpart 483.70(a) is NOT MET as evidenced by: Cooking Facilities CFR(s): NFPA 101 Cooking Facilities Cooking equipment is protected in accordance with NFPA 96, Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, unless: * residential cooking equipment (i.e., small appliances such as microwaves, hot plates, toasters) are used for food warming or limited cooking in accordance with 18.3.2.5.2, 19.3.2.5.2 * cooking facilities open to the corridor in smoke compartments with 30 or fewer patients comply with the conditions under 18.3.2.5.3, 19.3.2.5.3, or	K 324		10/3/23

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K 324	<p>Continued From page 3</p> <p>* cooking facilities in smoke compartments with 30 or fewer patients comply with conditions under 18.3.2.5.4, 19.3.2.5.4.</p> <p>Cooking facilities protected according to NFPA 96 per 9.2.3 are not required to be enclosed as hazardous areas, but shall not be open to the corridor.</p> <p>18.3.2.5.1 through 18.3.2.5.4, 19.3.2.5.1 through 19.3.2.5.5, 9.2.3, TIA 12-2</p> <p>This REQUIREMENT is not met as evidenced by: Based on a review of available documentation and staff interview, the facility failed to maintain proper inspection scheduling associated to range hood fire suppression system and other protective measures per NFPA 101 (2012 edition), Life Safety Code section 19.3.2.5.3(9), 9.2.3, and NFPA 96 (2014 edition), Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, section 11.2.1. These deficient findings could have a widespread impact on the residents within the facility.</p> <p>Findings Include:</p> <p>On 10/03/2023 between 11:00 AM and 3:00 PM, it was revealed by observation that the residential style stove located the Physical Therapy Area did not have full protective hardware</p> <p>An interview with the Maintenance Director verified these deficient findings at the time of discovery.</p>	K 324	<p>Based on a review of available documentation and staff interview, the facility failed to maintain proper inspection scheduling associated to range hood fire suppression system and other protective measures per NFPA 101 (2012 edition), Life Safety Code section 19.3.2.5.3 (9), 9.2.3, and NFPA 96 (2014 edition), Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, section 11.2.1. These deficient findings could have a widespread impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Therapy Oven immediately placed out of order. Electrician notified of needing required full protective hardware placed.</p>	

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K 345 K 345 SS=F	<p>Continued From page 4</p> <p>Fire Alarm System - Testing and Maintenance CFR(s): NFPA 101</p> <p>Fire Alarm System - Testing and Maintenance A fire alarm system is tested and maintained in accordance with an approved program complying with the requirements of NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm and Signaling Code. Records of system acceptance, maintenance and testing are readily available. 9.6.1.3, 9.6.1.5, NFPA 70, NFPA 72 This REQUIREMENT is not met as evidenced by: Based on a review of available documentation and staff interview, the facility failed to conduct sensitivity testing of the fire alarm system per NFPA 101 (2012 edition), Life Safety Code, sections 19.3.4.1, 9.6.1.3, and NFPA 72 (2010 edition), National Fire Alarm and Signaling Code, section 14.4.5.3. This deficient finding could have a patterned impact on the residents within the facility.</p> <p>Findings include:</p> <p>On 10/03/2023 between 11:00 AM and 3:00 PM, it was revealed by a review of available documentation that the most recent fire alarm sensitivity testing was completed 06/28/2021.</p> <p>An interview with the Maintenance Director verified this deficient finding at the time of discovery.</p>	K 345 K 345	<p>Based on review of available documentation and staff interview, the facility failed to conduct sensitivity testing of the fire alarm system per NFPA 101 (2012 edition), Life Safety Code, sections 19.3.4.1, 9.6.1.3, and NFPA 72 (2012 edition), National Fire Alarm and Signaling Code, section 14.4.5.3. This deficient finding could have a patterned impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Facility sensitivity test was completed October 5, 2023 Facility has educated facility Regional Maintenance Director on frequency of testing. Regional Maintenance or designee will report on any upcoming or overdue tests during quarterly QAPI</p>	10/5/23

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K 353 K 353 SS=F	<p>Continued From page 5</p> <p>Sprinkler System - Maintenance and Testing CFR(s): NFPA 101</p> <p>Sprinkler System - Maintenance and Testing Automatic sprinkler and standpipe systems are inspected, tested, and maintained in accordance with NFPA 25, Standard for the Inspection, Testing, and Maintaining of Water-based Fire Protection Systems. Records of system design, maintenance, inspection and testing are maintained in a secure location and readily available.</p> <p>a) Date sprinkler system last checked _____</p> <p>b) Who provided system test _____</p> <p>c) Water system supply source _____</p> <p>Provide in REMARKS information on coverage for any non-required or partial automatic sprinkler system. 9.7.5, 9.7.7, 9.7.8, and NFPA 25 This REQUIREMENT is not met as evidenced by: Based on a review of available documentation, and staff interview the facility failed to maintain the sprinkler system in accordance with NFPA 101 (2012 edition), Life Safety Code, sections 4.6.12, 9.7.5, 9.7.6, NFPA 25 (2011 edition) Standard for the Inspection, Testing, and Maintenance of Water-Based Fire Protection Systems, section(s), 5.1.1.1, 5.1.1.2, 5,2,5, 13.7.1, 4.3, 4.4 These deficient findings could have a widespread impact on the residents within the facility.</p> <p>Findings include: On 10/03/2023 between 11:00 AM and 3:00 PM,</p>	K 353 K 353	<p>Based on the review of the available documentation, and staff interview the facility failed to maintain the sprinkler system in accordance with NFPA 101 (2012 edition), Life Safety Code, sections 4.6.12, 9.7.5, 9.7.6, NFPA 25 (2011 edition) Standard for the Inspection, Testing, and Maintenance of Water-Based Fire Protection Systems, section(s) 5.1.1.1, 5.1.1.2, 5,2,5, 13.7.1, 4.3, 4.4. These deficient findings could have a widespread impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance.</p>	10/3/23

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K 353	Continued From page 6 it was revealed by a review of available documentation that there was no documentation presented to confirm that fire sprinkler system - 3rd quarter 2023 inspection had occurred. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 353	This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Facility fire sprinkler system quarter 3 inspection completed September 15, 2023 Facility has educated facility Regional Maintenance Director on frequency of testing. Regional Maintenance or designee will report on any upcoming or overdue tests during quarterly QAPI	
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, a review of available documentation and staff interview, the facility failed to properly inspect, and maintain 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 7.1.1, 7.1.2.2, 7.2.1.2, 7.2.4.3, 7.2.4.4, 7.2.4.5,, 7.3.1.1.1, 7.3.2.4 These deficient findings could have a widespread impact on the residents within the facility. Findings include: 1. On 10/03/2023 between 11:00 AM and 3:00	K 355	Based on observations, a review of available documentation and staff interview, the facility failed to properly inspect, and maintain 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 7.1.1, 7.1.1.2.2, 7.2.1.2, 7.2.4.3, 7.2.4.4, 7.2.4.5, 7.3.1.1.1, 7.3.2.4. These deficient findings could have a widespread impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not	10/23/23

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K 355	Continued From page 7 PM, it was revealed by a review of available documentation that there was no documentation presented for review to confirm either the facility or vendor had a current listing of fire extinguishers. 2. On 10/03/2023 between 11:00 AM and 3:00 PM, it was revealed by observation, that the fire extinguisher located in Physical Therapy was missing the annual hang-tag An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 355	constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Facility map marked with fire extinguisher locations within the facility and placed in Emergency Preparedness Binders. All Fire extinguishers checked to be in compliance. Education provided to staff on facility map location. Administrator or designee will audit compliance of operation weekly x4, monthly x3 months, then quarterly thereafter. Audit results will be reviewed by QAPI committee for further recommendations.		
K 918 SS=F	Electrical Systems - Essential Electric System 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	K 918		10/17/23	

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K 918	<p>Continued From page 8</p> <p>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 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 a 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, 8.3.4, 8.3.4.1, 8.4.9, 8.4.9.2. These deficient findings could have a widespread impact on the residents within the facility.</p> <p>Findings include:</p> <ol style="list-style-type: none"> 1. On 10/3/2023 between 10:00 AM and 4:30 PM, it was revealed by a review of available documentation that no documentation was presented for review to confirm that 36-month - 4-hour load bank testing is occurring. 2. On 10/3/2023 between 10:00 AM and 4:30 PM, it was revealed by a review of available documentation that vendor inspection records identified to the facility that emergency generator 	K 918	<p>Based on a 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, 8.3.4, 8.3.4.1, 8.4.9, 8.4.9.2. These deficient findings could have a widespread impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Facility 4 hr load bank testing was completed October 17, 2023. Facility has educated facility Regional Maintenance Director on frequency of testing. Regional Maintenance or designee will</p>	

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K 918	Continued From page 9 needed a new air filter. There was no documentation presented to confirm replacement had yet occurred. An interview with the Maintenance Director verified these deficient findings at the time of discovery.	K 918	report on any upcoming or overdue tests during quarterly QAPI	
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 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 electrical devices	K 920	Based on observations and staff interview, the facility failed to manage	10/3/23

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K 920	<p>Continued From page 10</p> <p>in accordance with NFPA 99 (2012 edition), Health Care Facilities Code, section 10.2.3.6, 10.2.4, 10.5.2.3 and NFPA 70, (2011 edition), National Electrical Code, sections 110.3(B), 400.8 (1) and UL 1363. These deficient findings could have an isolated impact on the residents within the facility.</p> <p>Findings include:</p> <p>On 10/03/2023 between 11:00 AM and 3:00 PM, it was revealed by observation that in the corridor adjacent to Physical Therapy and extension cord was in use and extended to the exterior of building to power and appliance (sump pump).</p> <p>An interview with the Maintenance Director verified these deficient findings at the time of discovery.</p>	K 920	<p>usage electrical devices in accordance with NFPA 99 (2012 edition), Health Care Facilities Code, section 10.2.3.6, 10.2.4, 10.5.2.3 and NFPA 70, (2011 edition), National Electrical Code, sections 110.3(B), 400.8 (1) and UL 1363. These deficient findings could have an isolated impact on the residents within the facility. Please accept the following as the facility's credible allegation of compliance. This Plan of Correction does not constitute any admission of guilt or liability by the facility and is submitted only in response to the regulatory requirements. Extension cord immediately removed, and facility common areas, resident rooms and offices checked. Electrician notified to place outdoor outlet near outdoor appliance requiring power. Facility has educated facility Regional Maintenance Director on extension cords. Maintenance Director or designee will audit compliance weekly x 4weeks, monthly x 3months and then quarterly thereafter. Audit results will be reviewed by QAPI committee for further recommendations.</p>	