

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

ID: 3YDJ

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

Facility ID: 00916

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245409		3. NAME AND ADDRESS OF FACILITY (L3) EDENBROOK OF ROCHESTER (L4) 1875 19TH STREET NORTHWEST (L5) ROCHESTER, MN (L6) 55901			4. TYPE OF ACTION: <u>7</u> (L8) 1. Initial 2. Recertification 3. Termination 4. CHOW 5. Validation 6. Complaint 7. On-Site Visit 9. Other 8. Full Survey After Complaint																
2.STATE VENDOR OR MEDICAID NO. (L2) 843242200		5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9) 01/13/2015			7. PROVIDER/SUPPLIER CATEGORY <u>02</u> (L7) 01 Hospital 05 HHA 09 ESRD 13 PTIP 22 CLIA 02 SNF/NF/Dual 06 PRTF 10 NF 14 CORF 03 SNF/NF/Distinct 07 X-Ray 11 ICF/IID 15 ASC 04 SNF 08 OPT/SP 12 RHC 16 HOSPICE																
6. DATE OF SURVEY 03/16/2022 (L34)		8. ACCREDITATION STATUS: <u> </u> (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other			FISCAL YEAR ENDING DATE: (L35) 12/31																
11. LTC PERIOD OF CERTIFICATION From (a) : To (b) :		10.THE FACILITY IS CERTIFIED AS: X A. In Compliance With Program Requirements Compliance Based On: <u> </u> 1. Acceptable POC			And/Or Approved Waivers Of The Following Requirements: <u> </u> 2. Technical Personnel <u> </u> 6. Scope of Services Limit <u> </u> 3. 24 Hour RN <u> </u> 7. Medical Director <u> </u> 4. 7-Day RN (Rural SNF) <u> </u> 8. Patient Room Size <u>X</u> 5. Life Safety Code <u> </u> 9. Beds/Room																
12.Total Facility Beds 81 (L18)		13.Total Certified Beds 81 (L17)			B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: A, 5* (L12)																
14. LTC CERTIFIED BED BREAKDOWN <table border="0"> <tr> <td>18 SNF</td> <td>18/19 SNF</td> <td>19 SNF</td> <td>ICF</td> <td>IID</td> </tr> <tr> <td></td> <td>81</td> <td></td> <td></td> <td></td> </tr> <tr> <td>(L37)</td> <td>(L38)</td> <td>(L39)</td> <td>(L42)</td> <td>(L43)</td> </tr> </table>					18 SNF	18/19 SNF	19 SNF	ICF	IID		81				(L37)	(L38)	(L39)	(L42)	(L43)	15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)	
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16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE): Edenbrook of Rochester 245409 is requesting a temporary waiver for K-916 for installing a remote generator annunciator due to supply chain issues. The exit date was 02/03/2022. It is recommended that CMS approve this waiver to allow the facility time to locate an annunciator panel and have it installed by 09/09/2022.																					
17. SURVEYOR SIGNATURE <u>Karen Aldinger, Unit Supervisor</u>			Date : 04/06/2022 (L19)		18. STATE SURVEY AGENCY APPROVAL <u>Melissa Poepping, Enforcement Specialist</u>																
					Date: 04/06/2022 (L20)																

PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY

19. DETERMINATION OF ELIGIBILITY <u> </u> 1. Facility is Eligible to Participate <u> </u> 2. Facility is not Eligible (L21)		20. COMPLIANCE WITH CIVIL RIGHTS ACT:		21. 1. Statement of Financial Solvency (HCFA-2572) 2. Ownership/Control Interest Disclosure Stmt (HCFA-1513) 3. Both of the Above : <u> </u>	
22. ORIGINAL DATE OF PARTICIPATION 01/01/1987 (L24)		23. LTC AGREEMENT BEGINNING DATE (L41)		24. LTC AGREEMENT ENDING DATE (L25)	
25. LTC EXTENSION DATE: (L27)		27. ALTERNATIVE SANCTIONS A. Suspension of Admissions: (L44) B. Rescind Suspension Date: (L45)			
26. TERMINATION ACTION: (L30) <u>VOLUNTARY</u> <u>00</u> 01-Merger, Closure 02-Dissatisfaction W/ Reimbursement 03-Risk of Involuntary Termination 04-Other Reason for Withdrawal			INVOLUNTARY 05-Fail to Meet Health/Safety 06-Fail to Meet Agreement <u>OTHER</u> 07-Provider Status Change 00-Active		
28. TERMINATION DATE:		29. INTERMEDIARY/CARRIER NO. 00160 (L28)		30. REMARKS (L31)	
31. RO RECEIPT OF CMS-1539 (L32)		32. DETERMINATION OF APPROVAL DATE (L33)			
DETERMINATION APPROVAL					



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
April 6, 2022

CMS Certification Number (CCN): 245409

Administrator
Edenbrook Of Rochester
1875 19th Street Northwest
Rochester, MN 55901

Dear Administrator:

The Minnesota Department of Health assists the Centers for Medicare and Medicaid Services (CMS) by surveying skilled nursing facilities and nursing facilities to determine whether they meet the requirements for participation. To participate as a skilled nursing facility in the Medicare program or as a nursing facility in the Medicaid program, a provider must be in substantial compliance with each of the requirements established by the Secretary of Health and Human Services found in 42 CFR part 483, Subpart B.

Based upon your facility being in substantial compliance, we are recommending to CMS that your facility be recertified for participation in the Medicare and Medicaid program.

Effective March 15, 2022 the above facility is certified for:

81 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 81 skilled nursing facility beds.

We have recommended CMS approve the waivers that you requested for the following Life Safety Code Requirements: K916.

If you are not in compliance with the above requirements at the time of your next survey, you will be required to submit a Plan of Correction for these deficiency(ies) or renew your request for waiver in order to continue your participation in the Medicare and Medicaid Program.

You should advise our office of any changes in staffing, services, or organization, which might affect your certification status. If, at the time of your next survey, we find your facility to not be in substantial compliance your Medicare and/or Medicaid provider agreement may be subject to non-renewal or termination.

Please contact me if you have any questions.

Sincerely,

Edenbrook Of Rochester

April 6, 2022

Page 2

A handwritten signature in black ink, appearing to read "Melissa Poepping". The signature is fluid and cursive, with a large initial "M" and a long, sweeping underline.

Melissa Poepping, Health Program Representative Senior

Program Assurance | Licensing and Certification

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
April 6, 2022

Administrator
Edenbrook Of Rochester
1875 19th Street Northwest
Rochester, MN 55901

RE: CCN: 245409
Cycle Start Date: February 3, 2022

Dear Administrator:

On March 16, 2022, 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.

Correction of the Life Safety Code deficiency(ies) cited under K916 at the time of the February 3, 2022 standard survey, has not yet been verified. Your plan of correction for this deficiency / these deficiencies, including your request for a temporary waiver with a date of completion of September 9, 2022, has been forwarded to the Region V Office of the Centers for Medicare and Medicaid Services (CMS) for their review and determination. Failure to come into substantial compliance with this deficiency / these deficiencies by the date indicated in your plan of correction may result in the imposition of enforcement remedies.

Feel free to contact me if you have questions.

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

Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
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

April 6, 2022

Administrator
Edenbrook Of Rochester
1875 19th Street Northwest
Rochester, MN 55901

Re: Reinspection Results
Event ID: 3YDJ12

Dear Administrator:

On March 16, 2022 survey staff of the Minnesota Department of Health - Health Regulation Division completed a reinspection of your facility, to determine correction of orders found on the survey completed on February 3, 2022. At this time these correction orders were found corrected.

Please feel free to call me with any questions.

Sincerely,

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

Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64900
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
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MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

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17. SURVEYOR SIGNATURE <u>Kyla Einertson, HFE NE II</u>			Date : 03/08/2022 (L19)		18. STATE SURVEY AGENCY APPROVAL <u>Melissa Poepping, Enforcement Specialist</u> Date: 03/18/2022 (L20)											

PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY

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Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 14, 2022

Administrator
Edenbrook Of Rochester
1875 19th Street Northwest
Rochester, MN 55901

RE: CCN: 245409
Cycle Start Date: February 3, 2022

Dear Administrator:

On February 3, 2022, 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.

Edenbrook Of Rochester

February 14, 2022

Page 2

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:

Karen Aldinger, Unit Supervisor
St. Cloud A District Office
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
3333 Division Street, Suite 212
Saint Cloud, Minnesota 56301-4557
Email: karen.aldinger@state.mn.us
Office: (651) 201-3794 Mobile: (320) 249-2805

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

Edenbrook Of Rochester

February 14, 2022

Page 3

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 May 3, 2022 (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 August 3, 2022 (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.

Edenbrook Of Rochester

February 14, 2022

Page 4

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,

A handwritten signature in black ink, appearing to read "Melissa Poepping". The signature is fluid and cursive, with the first name "Melissa" and last name "Poepping" clearly distinguishable.

Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64900
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: melissa.poepping@state.mn.us

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

PRINTED: 03/18/2022
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245409	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 02/03/2022
NAME OF PROVIDER OR SUPPLIER EDENBROOK OF ROCHESTER			STREET ADDRESS, CITY, STATE, ZIP CODE 1875 19TH STREET NORTHWEST ROCHESTER, MN 55901		
(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 000	Initial Comments On 1/31/22 through 2/3/22, 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. 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. 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.	E 000			
E 015 SS=C	Subsistence Needs for Staff and Patients CFR(s): 483.73(b)(1) §403.748(b)(1), §418.113(b)(6)(iii), §441.184(b)(1), §460.84(b)(1), §482.15(b)(1), §483.73(b)(1), §483.475(b)(1), §485.625(b)(1) [(b) Policies and procedures. [Facilities] must develop and implement emergency preparedness policies and procedures, based on the emergency plan set forth in paragraph (a) of this section, risk assessment at paragraph (a)(1) of this section, and the communication plan at paragraph (c) of this section. The policies and procedures must be reviewed and updated every 2 years [annually for LTC facilities]. At a minimum, the policies and procedures must address the following: (1) The provision of subsistence needs for staff and patients whether they evacuate or shelter in	E 015		3/4/22	

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

TITLE

(X6) DATE

Electronically Signed

02/23/2022

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
CENTERS FOR MEDICARE & MEDICAID SERVICES

PRINTED: 03/18/2022
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245409	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 02/03/2022
NAME OF PROVIDER OR SUPPLIER EDENBROOK OF ROCHESTER			STREET ADDRESS, CITY, STATE, ZIP CODE 1875 19TH STREET NORTHWEST ROCHESTER, MN 55901		
(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 015	<p>Continued From page 1</p> <p>place, include, but are not limited to the following:</p> <p>(i) Food, water, medical and pharmaceutical supplies</p> <p>(ii) Alternate sources of energy to maintain the following:</p> <p>(A) Temperatures to protect patient health and safety and for the safe and sanitary storage of provisions.</p> <p>(B) Emergency lighting.</p> <p>(C) Fire detection, extinguishing, and alarm systems.</p> <p>(D) Sewage and waste disposal.</p> <p>*[For Inpatient Hospice at §418.113(b)(6)(iii):] Policies and procedures.</p> <p>(6) The following are additional requirements for hospice-operated inpatient care facilities only. The policies and procedures must address the following:</p> <p>(iii) The provision of subsistence needs for hospice employees and patients, whether they evacuate or shelter in place, include, but are not limited to the following:</p> <p>(A) Food, water, medical, and pharmaceutical supplies.</p> <p>(B) Alternate sources of energy to maintain the following:</p> <p>(1) Temperatures to protect patient health and safety and for the safe and sanitary storage of provisions.</p> <p>(2) Emergency lighting.</p> <p>(3) Fire detection, extinguishing, and alarm systems.</p> <p>(C) Sewage and waste disposal.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility failed to include in their emergency preparedness plan (EPP) how to obtain</p>	E 015	<p>On 2/21/2022 EP book was updated with information to obtain pharmaceutical supplies and how to maintain sewage and</p>		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245409	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 02/03/2022
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E 015	Continued From page 2 pharmaceutical supplies and how to maintain sewage and waste disposal during an emergency. This had the potential to affect 44 residents at the facility. Findings include: Review of the facility's EPP revealed the facility failed to address how they would obtain pharmaceutical supplies and how they would maintain sewage and waste disposal during an emergency. During interview on 2/2/22, at 10:27 a.m. the administrator verified this information.	E 015	waste disposal during emergency. NHA/Maintenance will be educated on the process Audits will be done monthly to ensure documentation is in place. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. NHA will be responsible for making sure documentation is in place		
E 030 SS=C	Names and Contact Information CFR(s): 483.73(c)(1) §403.748(c)(1), §416.54(c)(1), §418.113(c)(1), §441.184(c)(1), §460.84(c)(1), §482.15(c)(1), §483.73(c)(1), §483.475(c)(1), §484.102(c)(1), §485.68(c)(1), §485.625(c)(1), §485.727(c)(1), §485.920(c)(1), §486.360(c)(1), §491.12(c)(1), §494.62(c)(1). [(c) The [facility must develop and maintain an emergency preparedness communication plan that complies with Federal, State and local laws and must be reviewed and updated at least every 2 years [annually for LTC facilities]. The communication plan must include all of the following:] (1) Names and contact information for the following: (i) Staff. (ii) Entities providing services under arrangement. (iii) Patients' physicians	E 030		3/4/22	

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

PRINTED: 03/18/2022
FORM APPROVED
OMB NO. 0938-0391

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E 030	Continued From page 3 (iv) Other [facilities]. (v) Volunteers. *[For Hospitals at §482.15(c) and CAHs at §485.625(c)] The communication plan must include all of the following: (1) Names and contact information for the following: (i) Staff. (ii) Entities providing services under arrangement. (iii) Patients' physicians (iv) Other [hospitals and CAHs]. (v) Volunteers. *[For RNHCIs at §403.748(c):] The communication plan must include all of the following: (1) Names and contact information for the following: (i) Staff. (ii) Entities providing services under arrangement. (iii) Next of kin, guardian, or custodian. (iv) Other RNHCIs. (v) Volunteers. *[For ASCs at §416.45(c):] The communication plan must include all of the following: (1) Names and contact information for the following: (i) Staff. (ii) Entities providing services under arrangement. (iii) Patients' physicians. (iv) Volunteers. *[For Hospices at §418.113(c):] The communication plan must include all of the following: (1) Names and contact information for the	E 030			

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E 030	<p>Continued From page 4 following:</p> <ul style="list-style-type: none"> (i) Hospice employees. (ii) Entities providing services under arrangement. (iii) Patients' physicians. (iv) Other hospices. <p>*[For HHAs at §484.102(c):] The communication plan must include all of the following:</p> <ul style="list-style-type: none"> (1) Names and contact information for the following: <ul style="list-style-type: none"> (i) Staff. (ii) Entities providing services under arrangement. (iii) Patients' physicians. (iv) Volunteers. <p>*[For OPOs at §486.360(c):] The communication plan must include all of the following:</p> <ul style="list-style-type: none"> (2) Names and contact information for the following: <ul style="list-style-type: none"> (i) Staff. (ii) Entities providing services under arrangement. (iii) Volunteers. (iv) Other OPOs. (v) Transplant and donor hospitals in the OPO's Donation Service Area (DSA). <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility's communication plan failed to include the required information including names and contact information for physicians. This had the potential to affect all 44 residents in the facility.</p> <p>Findings include:</p> <p>Review of the facility's EPP revealed the facility failed to list the physician and their contact numbers in the EPP.</p>	E 030	<p>On 2/21/2022 EPP book was updated with all physicians for the community and phone numbers. NHA and DON were educated on the process and how to update Audits will be done monthly to ensure documentation is in place and all correct physicians are listed. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. NHA will be responsible for making sure</p>		

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

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E 030	Continued From page 5 During interview on 2/2/22, at 11:04 a.m. the administrator acknowledged there was not a list of physicians and their contact numbers in the emergency preparedness plan.	E 030	documentation is in place		
E 031 SS=C	Emergency Officials Contact Information CFR(s): 483.73(c)(2) §403.748(c)(2), §416.54(c)(2), §418.113(c)(2), §441.184(c)(2), §460.84(c)(2), §482.15(c)(2), §483.73(c)(2), §483.475(c)(2), §484.102(c)(2), §485.68(c)(2), §485.625(c)(2), §485.727(c)(2), §485.920(c)(2), §486.360(c)(2), §491.12(c)(2), §494.62(c)(2). [(c) The [facility] must develop and maintain an emergency preparedness communication plan that complies with Federal, State and local laws and must be reviewed and updated at least every 2 years [annually for LTC facilities]. The communication plan must include all of the following: (2) Contact information for the following: (i) Federal, State, tribal, regional, and local emergency preparedness staff. (ii) Other sources of assistance. *[For LTC Facilities at §483.73(c):] (2) Contact information for the following: (i) Federal, State, tribal, regional, and local emergency preparedness staff. (ii) The State Licensing and Certification Agency. (iii) The Office of the State Long-Term Care Ombudsman. (iv) Other sources of assistance. *[For ICF/IIDs at §483.475(c):] (2) Contact information for the following:	E 031	3/4/22		

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E 031	Continued From page 6 (i) Federal, State, tribal, regional, and local emergency preparedness staff. (ii) Other sources of assistance. (iii) The State Licensing and Certification Agency. (iv) The State Protection and Advocacy Agency. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to ensure their Emergency Preparedness Plan (EPP) included contact information for Federal emergency preparedness staff, the State Licensing and Certification Agency and contact information for the Ombudsman. This had the potential to affect all 44 residents currently residing in the facility. Findings include: The facility's EPP was reviewed with the administrator. The plan-included components of a communication plan however, lacked documentation of contact information for Federal emergency preparedness staff, the State Licensing and Certification Agency and the Ombudsman. On 2/2/22, at 11:08 a.m. the administrator verified this information.	E 031	Emergency Preparedness book was updated to include contact information for federal emergency preparedness staff, the state licensing and certification agency and contact information for the Ombudsman. NHA and DON were educated regarding the process of the documentation and the effect it can have on residents. Audits will be done monthly to ensure documentation is in place for the correct contact information. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. NHA will be responsible for making sure documentation is in place.		
E 041 SS=C	Hospital CAH and LTC Emergency Power CFR(s): 483.73(e) §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.	E 041		3/4/22	

DEPARTMENT OF HEALTH AND HUMAN SERVICES
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E 041	Continued From page 7 §483.73(e), §485.625(e) (e) Emergency and standby power systems. The [LTC facility and the CAH] must implement emergency and standby power systems based on the emergency plan set forth in paragraph (a) of this section. §482.15(e)(1), §483.73(e)(1), §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. 482.15(e)(2), §483.73(e)(2), §485.625(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. 482.15(e)(3), §483.73(e)(3), §485.625(e)(3) 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. *[For hospitals at §482.15(h), LTC at §483.73(g), and CAHs §485.625(g):]	E 041			

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E 041	<p>Continued From page 8</p> <p>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 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>	E 041			

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E 041	Continued From page 9 (xi) TIA 12-4 to NFPA 101, issued October 22, 2013. (xiii) NFPA 110, Standard for Emergency and Standby Power Systems, 2010 edition, including TIAs to chapter 7, issued August 6, 2009.. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to test their emergency generator as required. This had the potential to affect all 44 residents in the facility. Findings include: As a result of the Life Safety Code survey on 2/10/22, during documentation review with the environmental service director (ESD) on 2/10/22, it was revealed that the generator had not been exercised under load for 30 minutes during November and December, 2021 and January, 2022. The ESD verified a test had not occurred.	E 041	On 02/23/2022 the emergency generator was exercised under load for 30 minutes. Monthly testing will be conducted for future months. Maintenance will be educated on the requirement for generator testing on a monthly basis. Audits for testing and inspecting the generator under load for 30 minutes will be completed monthly. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. The Director of Maintenance will be responsible for compliance.		
F 000	INITIAL COMMENTS On 1/31/21 through 2/3/22, a standard recertification survey was conducted at your facility. Complaint investigations were also conducted. Your facility was found to be NOT in compliance with the requirements of 42 CFR 483, Subpart B, Requirements for Long Term Care Facilities. The following complaints were found to be substantiated with no deficiencies cited due to actions implemented by the facility prior to survey. H5409104C/MN80488 H5409105C/MN80202 H5409109C/MN72703	F 000			

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F 000	Continued From page 10 The following complaints were found to be UNSUBSTANTIATED: H5409106C/MN79143 H5409107C/MN76266 H5409108C/MN75190 H5409110C/MN71280 H5409111C/MN70882 H5409112C/MN65322 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. Your electronic submission of the POC will be used as verification of compliance. Upon receipt of an acceptable electronic POC, an onsite revisit of your facility may be conducted to validate substantial compliance with the regulations has been attained.	F 000			
F 582 SS=D	Medicaid/Medicare Coverage/Liability Notice CFR(s): 483.10(g)(17)(18)(i)-(v) §483.10(g)(17) The facility must-- (i) Inform each Medicaid-eligible resident, in writing, at the time of admission to the nursing facility and when the resident becomes eligible for Medicaid of- (A) The items and services that are included in nursing facility services under the State plan and for which the resident may not be charged; (B) Those other items and services that the facility offers and for which the resident may be charged, and the amount of charges for those services; and (ii) Inform each Medicaid-eligible resident when	F 582		3/15/22	

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F 582	<p>Continued From page 11</p> <p>changes are made to the items and services specified in §483.10(g)(17)(i)(A) and (B) of this section.</p> <p>§483.10(g)(18) The facility must inform each resident before, or at the time of admission, and periodically during the resident's stay, of services available in the facility and of charges for those services, including any charges for services not covered under Medicare/ Medicaid or by the facility's per diem rate.</p> <p>(i) Where changes in coverage are made to items and services covered by Medicare and/or by the Medicaid State plan, the facility must provide notice to residents of the change as soon as is reasonably possible.</p> <p>(ii) Where changes are made to charges for other items and services that the facility offers, the facility must inform the resident in writing at least 60 days prior to implementation of the change.</p> <p>(iii) If a resident dies or is hospitalized or is transferred and does not return to the facility, the facility must refund to the resident, resident representative, or estate, as applicable, any deposit or charges already paid, less the facility's per diem rate, for the days the resident actually resided or reserved or retained a bed in the facility, regardless of any minimum stay or discharge notice requirements.</p> <p>(iv) The facility must refund to the resident or resident representative any and all refunds due the resident within 30 days from the resident's date of discharge from the facility.</p> <p>(v) The terms of an admission contract by or on behalf of an individual seeking admission to the facility must not conflict with the requirements of these regulations.</p> <p>This REQUIREMENT is not met as evidenced by:</p>	F 582			

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F 582	<p>Continued From page 12</p> <p>Based on interview, and document review, the facility failed to provide the required skilled nursing facility advanced beneficiary notice of non-coverage (SNFABN) (CMS 10055), notice of Medicare part A non-coverage two days in advance for 2 of 4 residents (R7, R25) whose Medicare part A coverage ended and then remained in the facility.</p> <p>Findings include:</p> <p>R7's Notice of Medicare Non-Coverage (CMS 10123-NOMNC) form dated 9/3/21, revealed R7 would be liable for skilled nursing stay beginning 9/8/21. The SNFABN form was signed and dated 9/7/21, the last covered day of Medicare-A services.</p> <p>R25's Notice of Medicare Non-Coverage (CMS 10123-NOMNC) revealed R25 would be liable for skilled nursing stay beginning 12/21/21. The progress notes indicated this was reviewed with the responsible party on 12/17/21. The SNFABN form was reviewed with the responsible party on 12/20/21, the last covered day of Medicare-A services.</p> <p>On 2/01/22, at 10:11 a.m. registered nurse (RN)-A stated the SNFABN did not require a two-day notice.</p> <p>On 02/01/22, at 2:03 p.m. the director of nursing (DON) stated the SNFABN notices should be given 2 days in advance and verified the SNFABN notices for R7 and R25 were given their last day of Medicare-A coverage.</p> <p>The facility's Beneficiary Notice Guidelines dated 2021, indicated notices were to be given in writing</p>	F 582	<p>The noted residents R7 and R25 were not financially affected nor were displaced related to the notice less than 48 hours. Both remained in facility R7 went to private pay and R25 remained on Medicaid.</p> <ol style="list-style-type: none"> 1. Review of all SNFABN in the last 30 days. 2. Education to MDS nurse on process of issuing SNFABN. 3. Audit SNFABN notices weekly times four weeks 4. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. 5. NHA/MDS or designee are responsible for compliance 		

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F 582	Continued From page 13 far enough in advance to enable resident to make an informed decision and delivered to the resident with at least two days' notice even if he/she agrees with the notice/decision.	F 582			
F 585 SS=D	Grievances CFR(s): 483.10(j)(1)-(4) §483.10(j) Grievances. §483.10(j)(1) The resident has the right to voice grievances to the facility or other agency or entity that hears grievances without discrimination or reprisal and without fear of discrimination or reprisal. Such grievances include those with respect to care and treatment which has been furnished as well as that which has not been furnished, the behavior of staff and of other residents, and other concerns regarding their LTC facility stay. §483.10(j)(2) The resident has the right to and the facility must make prompt efforts by the facility to resolve grievances the resident may have, in accordance with this paragraph. §483.10(j)(3) The facility must make information on how to file a grievance or complaint available to the resident. §483.10(j)(4) The facility must establish a grievance policy to ensure the prompt resolution of all grievances regarding the residents' rights contained in this paragraph. Upon request, the provider must give a copy of the grievance policy to the resident. The grievance policy must include: (i) Notifying resident individually or through postings in prominent locations throughout the facility of the right to file grievances orally	F 585		3/15/22	

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

PRINTED: 03/18/2022
FORM APPROVED
OMB NO. 0938-0391

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F 585	Continued From page 14 (meaning spoken) or in writing; the right to file grievances anonymously; the contact information of the grievance official with whom a grievance can be filed, that is, his or her name, business address (mailing and email) and business phone number; a reasonable expected time frame for completing the review of the grievance; the right to obtain a written decision regarding his or her grievance; and the contact information of independent entities with whom grievances may be filed, that is, the pertinent State agency, Quality Improvement Organization, State Survey Agency and State Long-Term Care Ombudsman program or protection and advocacy system; (ii) Identifying a Grievance Official who is responsible for overseeing the grievance process, receiving and tracking grievances through to their conclusions; leading any necessary investigations by the facility; maintaining the confidentiality of all information associated with grievances, for example, the identity of the resident for those grievances submitted anonymously, issuing written grievance decisions to the resident; and coordinating with state and federal agencies as necessary in light of specific allegations; (iii) As necessary, taking immediate action to prevent further potential violations of any resident right while the alleged violation is being investigated; (iv) Consistent with §483.12(c)(1), immediately reporting all alleged violations involving neglect, abuse, including injuries of unknown source, and/or misappropriation of resident property, by anyone furnishing services on behalf of the provider, to the administrator of the provider; and as required by State law; (v) Ensuring that all written grievance decisions include the date the grievance was received, a	F 585			

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F 585	<p>Continued From page 15</p> <p>summary statement of the resident's grievance, the steps taken to investigate the grievance, a summary of the pertinent findings or conclusions regarding the resident's concerns(s), a statement as to whether the grievance was confirmed or not confirmed, any corrective action taken or to be taken by the facility as a result of the grievance, and the date the written decision was issued;</p> <p>(vi) Taking appropriate corrective action in accordance with State law if the alleged violation of the residents' rights is confirmed by the facility or if an outside entity having jurisdiction, such as the State Survey Agency, Quality Improvement Organization, or local law enforcement agency confirms a violation for any of these residents' rights within its area of responsibility; and</p> <p>(vii) Maintaining evidence demonstrating the result of all grievances for a period of no less than 3 years from the issuance of the grievance decision.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on observation, interview, and document review, the facility failed to follow its grievance policy for 1 of 1 residents (R21) reviewed for missing personal items.</p> <p>Findings include:</p> <p>RR21's quarterly Minimum Data Set (MDS) dated 12/24/21, included cognitively intact.</p> <p>When interviewed on 1/31/22, at 12:18 p.m. R21 stated she had been missing a shirt and several other items of clothing for over two and a half weeks. R21 stated. she felt as though she had reported it to everyone in laundry services, but did not feel as though anything was being done. At that time, a housekeeper (HK)-B entered the</p>	F 585	<p>The resident has the right to voice grievances to the facility or other agency or entity that hears grievances without discrimination or reprisal and without fear of discrimination or reprisal.</p> <p>R21 was helped to find items that went missing and shown the grievance process. Resident had no mental or physical distress from this incident.</p> <p>1. Post signage around building those grievances can be written or verbal. Posting will include how to report a grievance. Who is the grievance official is Dave Molitor, Administrator, (507) 282-9449, d.molitor@edenbrookrochester.com</p> <p>2. Education to staff about the grievance process and where to find form.</p>		

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F 585	<p>Continued From page 16</p> <p>room, and R21 was observed to repeat her report of missing items to HK-B. HK-B said, "yeah, maybe someone put it in the wrong closet." R21 asked HK-B if they couldn't look in other resident closets as she was sure she had seen her gray Vikings shirt on a rack going down the hall at some time. HK-B stated, "I guess I could make a suggestion to my supervisor."</p> <p>When interviewed on 2/1/22, at 10:42 a.m. HK-A stated HK-B had told her about R21's concern of missing clothing on 1/31/22. HK-A recalled she had received an undated, unsigned note providing a description of the missing items, but it had just been laying on the desk and she was unsure for how long. HK-A stated that after HK-B told her about the missing clothing on 1/31/22, she had looked in the lost and found, and then told R21, "in a couple days I will look in the closets." HK-A stated she had not filled out any report or grievance.</p> <p>When interviewed on 2/1/22, at 10:42 a.m. the environmental services director (ESD) stated a grievance form should have been filled out and given it to the facility social worker (SW). ESD stated the proper procedure for missing items was for the person receiving the report to fill out the grievance form, hand it to the SW who would take note, then bring to ESD. The ESD would initiate a search for any missing items and complete the grievance form with the findings of their investigation, and return it to the SW. ESD stated any resident or staff could get a grievance form from the nurses station or from the SW. ESD stated there had been plenty of training to staff about the process.</p> <p>On 2/01/22, at 10:59 a.m. a grievance form</p>	F 585	<p>3. Audit- Random testing of staff on the process every week for four weeks.</p> <p>4. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring.</p> <p>5. NHA/SS or designee are responsible for compliance</p>		

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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 585	Continued From page 17 related to R21's missing items was observed laying on the SW's desk. The SW stated, "I got the grievance sheet about two minutes before you walked in the door. [The ESD] filled it out after talking to the surveyor." SW stated an expectation for staff to fill out a grievance form at the time any resident should have a concern such as missing personal items. According to a facility policy titled Grievance/Concern last revised 1/14/22: 1. Facility will make prompt efforts to resolve all grievances. 2. Facility will provide this policy and form to residents and have them readily available in common areas of the facility, areas may include facility entrance, reception desk, nurses' station, etc. 3. Facility staff will immediately report all alleged violations 8. Facility will use the information gathered in a grievance investigation to prevent further potential violations of any resident rights	F 585			
F 609 SS=D	Reporting of Alleged Violations CFR(s): 483.12(c)(1)(4) §483.12(c) In response to allegations of abuse, neglect, exploitation, or mistreatment, the facility must: §483.12(c)(1) Ensure that all alleged violations involving abuse, neglect, exploitation or mistreatment, including injuries of unknown source and misappropriation of resident property, are reported immediately, but not later than 2 hours after the allegation is made, if the events that cause the allegation involve abuse or result in serious bodily injury, or not later than 24 hours if	F 609		3/15/22	

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F 609	<p>Continued From page 18</p> <p>the events that cause the allegation do not involve abuse and do not result in serious bodily injury, to the administrator of the facility and to other officials (including to the State Survey Agency and adult protective services where state law provides for jurisdiction in long-term care facilities) in accordance with State law through established procedures.</p> <p>§483.12(c)(4) Report the results of all investigations to the administrator or his or her designated representative and to other officials in accordance with State law, including to the State Survey Agency, within 5 working days of the incident, and if the alleged violation is verified appropriate corrective action must be taken. This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility did not file a report to the State Agency for 1 of 2 residents' (R10) allegations of missing money while living in the facility.</p> <p>Findings include:</p> <p>R10's significant change Minimum Data Set (MDS) dated 1/25/22, identified them as cognitively intact.</p> <p>When interviewed on 1/31/22, at 1:30 p.m. R10 stated he had reported \$800 missing, "some time ago," and could not recall if anything had been done about his concern.</p> <p>When interviewed on 1/31/22, at 6:07 p.m. the administrator stated he had never received a report R10 was missing \$800.</p> <p>When interviewed on 2/1/22, at 11:00 a.m. the</p>	F 609	<p>R10 had possibly lost money. NHA and SS checked in with resident and he did not want help with finding the money and has no emotional or physical distress from the event. Resident was educated on the VA process and the resources that the community can give him. Resident already had lock box in place to secure money.</p> <ol style="list-style-type: none"> 1. Community will report all incidents of abuse of a resident to the state in a timely manner. 2. Review all grievances for the past month to ensure that none were reportable as VAs. 3. Educate staff about the VA process. 4. Signs around community about how to report, who to report, what to report and how to report. (Pink sheets) 5. Audit weekly to ensure that grievances have been reported if needed 		

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F 609	<p>Continued From page 19</p> <p>social worker (SW) stated she did not recall receiving a concern form related to R10, and had never heard from R10 that he was missing any money. SW stated R10 had a lockbox in his room where he could keep his personal items.</p> <p>When interviewed on 2/2/22, at 1:38 p.m. R10 stated, the \$800 had been missing for several months but he was unable to give a specific date. R10 said his sister brought the money from the bank for him because he had bills to pay and wanted to go buy some personal items. He said he kept it in an envelope on his window sill, and stated, "I never had a problem with that before." R10 reported he was out of his room at one point, and when he came back, he was looking for his money, but did not see it on the window sill. R10 said he later found the empty envelope on his bedside stand but the money was gone. R10 repeated that he had reported it to someone, but could not recall exactly the person he told. Following this, R10 said he recalled the person he had reported to told him it would be reported and investigated, but said, "I never heard anything since." At the time of interview, R10 was observed to have many personal items scattered about his room, including medical records on his window sill, and also a few dollars bills and some loose change in a clear plastic cup at his bedside.</p> <p>When interviewed on 2/2/22, at 2:36 p.m. the administrator stated he had found a grievance related to R10's missing \$800 that had been completed in August prior to the time he had been hired. Administrator stated he had not yet talked to R10 about the missing money as R10 could be difficult, and the Administrator was, "trying to find him on a good day." The administrator stated the grievance form indicated R10 did not want</p>	F 609	<p>for 4 weeks</p> <p>6. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring.</p> <p>7. NHA/DON/SS or designee are responsible for compliance</p>		

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F 609	Continued From page 20 anything done. The administrator stated R10's missing money grievance had been documented by the previous social worker, and had been minimally investigated by the administrator at that time. The administrator stated the allegation of missing \$800 should have been reported to the state agency and investigated at that time as possible theft, but had not. As of 2/2/22, no report related to R10's missing money was found on the State Agencies Nursing Home Incident Reporting site. According to a facility policy titled Vulnerable Adult Abuse and Neglect Prevention revised 11/17/20, with a policy addendum effective 6/23/21: "all residents are susceptible to maltreatment and exploitation due to their need for nursing home care." "Any nursing home employee or volunteer who becomes aware of abuse, mistreatment, neglect or misappropriation shall intervene to safeguard the resident and then immediately report to the Nursing Home Administrator or designee. The Nursing Home Administrator or designee will report abuse to the state agency per State and Federal requirements." The addendum indicated police were to be notified if a crime was suspected.	F 609			
F 657 SS=D	Care Plan Timing and Revision CFR(s): 483.21(b)(2)(i)-(iii) §483.21(b) Comprehensive Care Plans §483.21(b)(2) A comprehensive care plan must be- (i) Developed within 7 days after completion of the comprehensive assessment. (ii) Prepared by an interdisciplinary team, that includes but is not limited to--	F 657			3/15/22

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F 657	<p>Continued From page 21</p> <p>(A) The attending physician.</p> <p>(B) A registered nurse with responsibility for the resident.</p> <p>(C) A nurse aide with responsibility for the resident.</p> <p>(D) A member of food and nutrition services staff.</p> <p>(E) To the extent practicable, the participation of the resident and the resident's representative(s). An explanation must be included in a resident's medical record if the participation of the resident and their resident representative is determined not practicable for the development of the resident's care plan.</p> <p>(F) Other appropriate staff or professionals in disciplines as determined by the resident's needs or as requested by the resident.</p> <p>(iii) Reviewed and revised by the interdisciplinary team after each assessment, including both the comprehensive and quarterly review assessments.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility failed to ensure a care conference was held after a significant change in condition assessment for 1 of 3 residents (R7) reviewed for care conferences.</p> <p>Findings include:</p> <p>R7's admission record indicated he was admitted to the facility on 8/9/21 and had a diagnosis of unspecified dementia without behavioral disturbance.</p> <p>R7's significant change of condition Minimum Data Set (MDS) dated 11/19/21, indicated R7 had moderate cognitive impairment, clear speech, was understood and understands.</p>	F 657	<p>The resident R7 care conference scheduled 02/23/2022.</p> <ol style="list-style-type: none"> Care conferences were held upon admission and when a quarterly, annual or change of condition MDS assessment was completed. Educate SS on the Care Plan process Review residents to make sure Care conferences have been completed on time or have been scheduled Audit care conferences weekly for four weeks to ensure that all needed care conferences are completed. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. 		

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F 657	Continued From page 22 During an interview on 1/31/22, at 1:59 p.m. R7 stated he was not aware of any care conferences and was not sure what this writer was talking about when asked about care conferences. R7's medical record revealed R7 had one care conference since admission held on 9/1/21. On 2/01/22, at 11:11 a.m. social services (SS) stated care conferences should be held on admission, quarterly, annual or change of condition basis. SS verified the only documentation in R7's medical record for a care conference held was on 9/1/21. SS stated R7 should have had another care conference by this time. SS stated R7 had a change of condition MDS on 9/15/21 and 11/19/21 and verified no care conferences were held for these change of condition MDS assessments. SS stated R7's next scheduled care conference was scheduled 2/23/22. On 2/02/22, at 1:56 p.m. the director of nursing (DON) stated her expectation was care conferences were held upon admission and when a quarterly, annual or change of condition MDS assessment was completed. Care Plan/Conference policy dated 1/14/22, included, "Every effort will be made to schedule care plan meetings at the best time of the day for the resident, resident's representative and family."	F 657	6. SS or designee are responsible for compliance		
F 684 SS=D	Quality of Care CFR(s): 483.25 § 483.25 Quality of care	F 684		3/15/22	

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F 684	<p>Continued From page 23</p> <p>Quality of care is a fundamental principle that applies to all treatment and care provided to facility residents. Based on the comprehensive assessment of a resident, the facility must ensure that residents receive treatment and care in accordance with professional standards of practice, the comprehensive person-centered care plan, and the residents' choices. This REQUIREMENT is not met as evidenced by:</p> <p>Based on observation, interview, and document review, the facility failed to monitor the condition of 1 of 1 residents (R4) with lymphedema and failed to follow-up on orders provided for R4 to receive edema care.</p> <p>Findings include:</p> <p>R4's 5 day perspective payment system (PPS) Minimum Data Set (MDS) dated 1/17/21, included, cognitively intact, required extensive assistance for dressing, bathing and grooming and limited assistance of 1 person for wheel chair mobility, and had limited range of motion of left upper extremity. R4's diagnoses included, a malignant neoplasm of connective and soft tissue of left upper limb, including the shoulder. Edema was not marked on the MDS.</p> <p>During observation and interview on 1/31/22, at 2:43 p.m. R4 was observed to have significant swelling of her left upper arm, with the area appearing to be nearly twice the size of her right arm. R4 had an ace wrap on her lower left arm that had slid down and was wrinkled, bunched and sloppy. R4 stated she wrapped the arm herself, but said a nurse would wrap it if she requested this. R4 also stated that therapy used to wrap it for her, but she was no longer in</p>	F 684	<p>Quality of Care</p> <p>The resident R4 with lymphedema was evaluated during the survey and continues to be treated for lymphedema by therapy. There was no negative impact on the resident due to the delay of the order being placed.</p> <ol style="list-style-type: none"> 1. Review the last 30 days of orders for therapy to validate they have been entered and addressed 2. Education on order processing and review to nursing staff, Health Unit Coordinator, and therapy 3. Audit therapy orders weekly to ensure accuracy of process times four weeks 4. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. 5. DON or designee are responsible for compliance 		

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F 684	<p>Continued From page 24</p> <p>therapy and had to do the care herself. R4 stated she has had lymphedema for many years, but that her upper arm was now larger than it had ever been.</p> <p>During observation on 2/1/22, at 11:13 a.m. R4 was in her room and stated her arm had not been wrapped yet today.</p> <p>R4's current physician orders for February 2022 did not show an order to monitor edema or to wrap her left upper extremity.</p> <p>R4's care plan did not contain information regarding her lymphedema or any directions for care or monitoring of this condition.</p> <p>A hand written form in R4's medical record indicated she had been seen 1/6/22 for left lymphedema consultation, and ordered recommendations were written for physical or occupation therapy (PT or OT) to wrap R4's left upper extremity fingers, hand and arm with short stretch wrap daily and leave in place for 23 hours per day and then rewrap.</p> <p>A hospital discharge summary dated 1/19/22 indicated R4 had lymphedema for over twenty years.</p> <p>A daily skilled charting note dated 1/20/22 indicated R4 had edema of her left upper extremity.</p> <p>When interviewed on 2/1/22, at 11:38 a.m. occupational therapist (OT)-A stated she was not lymphedema certified but had wrapped R4's arm a few times. OT-A did not recall receiving an order for PT or OT to wrap R4's arm, and said</p>	F 684			

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

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F 684	<p>Continued From page 25</p> <p>she was also unaware that no orders had been provided to the nursing staff to provide edema care. OT-A stated that even if R4 wrapped her arm herself, the facility was responsible to make sure the wrap was in place and correctly applied.</p> <p>When interviewed on 2/3/22, at 9:46 a.m. licensed practical nurse (LPN)-C stated she was aware that one of R4's arms was, "very swollen," but she did not provide any treatment for the edema. LPN-C said she thought perhaps PT wrapped the arm, but she did not know. LPN-C was not able to describe any nursing interventions a nurse could take independently to provide care for edematous extremities and said she had been taught to go by orders, and since there were never any treatments orders for R4's edema she had not done anything. She did not recall R4 asking her to wrap her arm.</p> <p>On 2/03/22, at 11:25 a.m. the director of nursing (DON) stated R4's edema should be on her diagnosis list, and should have been listed on her care plan. DON stated nursing staff should be monitoring edema and note if it is a chronic condition, and monitor for changes, updating the medical provider as needed. DON stated edema monitoring and any orders to wrap an extremity should be listed on the medication or treatment administration order sheets so a nurse would be reminded to monitor the resident's condition. DON also said a nurse noting a resident with edema, but not seeing any orders for care should clarify this with the medical provider. DON confirmed that R4 had not had edema listed on her care plan, and also confirmed the order for lymphedema wraps from 1/6/22, had not been followed up on.</p>	F 684			

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F 684	Continued From page 26 A policy specific to edema care and monitoring was not provided, but a facility policy titled Resident Assessment and Examination dated 8/1/15 indicated residents should be assessed for abnormalities in health states in order that the care team could implement interventions to address the concern. The policy indicated a body systems review, which included edema, was to be performed and any abnormalities reported to the physician.	F 684			
F 759 SS=D	Free of Medication Error Rts 5 Prcnt or More CFR(s): 483.45(f)(1) §483.45(f) Medication Errors. The facility must ensure that its- §483.45(f)(1) Medication error rates are not 5 percent or greater; This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review, the facility failed to ensure a medication error rate of less than 5%, medication errors were noted for 3 out of 26 medications observed, affecting 2 residents (R44 and R35) for a medication error rate of 12%. Findings include: R44's significant change Minimum Data Set (MDS) dated 1/14/22, included, cognitively intact, a diagnosis of diabetes and received insulin injections. R35's quarterly MDS dated 1/25/22, included, severe cognitive impairment, diagnosis of diabetes and received insulin injections.	F 759	The residents noted in findings R 35 and R 44 blood sugars were monitored post error with no noted adverse effects. 1. Education immediately started on insulin pen administration for nursing staff and continues 2. Audit hands on observation of the insulin administration technique RN-C and LPN-B weekly times 4 weeks, random audits for other nurses <input type="checkbox"/> weekly times four weeks 3. All findings will be brought to the facility <input type="checkbox"/> s next QAPI meeting for review and recommendation for continuance of monitoring. 4. DON or designee are responsible for compliance	3/15/22	

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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 759	<p>Continued From page 27</p> <p>On 2/1/22, at 9:15 a.m. a registered nurses (RN)-C was observed to prepare two insulin flex-pens for R44's ordered breakfast insulin (a flex pen contains the vial of insulin inside the pen, and has a mechanism where the dose to be administered is set on a dial at the top of the pen, and only that amount can then be injected). R44 was to receive 7 units of Aspart insulin (a rapid acting insulin) twice daily with meals, and 32 units of Tresiba insulin (a long acting insulin) once each morning according to his medication administration record (MAR) for February 2020. RN-C performed hand hygiene, cleansed the top of the flex-pen with an alcohol wipe and attached a needle. RN-C then set the dial for each insulin to match the ordered dose in the MAR. RN-C failed to prime the needle (setting the flex-pen dial to 2 units and injecting that into the needle prior to setting the dose). RN-C ascertained R44's morning blood sugar and proceeded to R44's room. RN-C applied gloves, cleansed a spot on R44's abdomen where she injection the Aspart, and held the needle in place for 10 seconds and then gave the Tesiba, also on the right side of the abdomen in the same manner. RN-C stated she had received training in how to properly use a flex-pen when she was hired.</p> <p>On 2/2/22, 8:24 a.m. a licensed practical nurse (LPN)-B was observed in preparation of R35's Aspart insulin dose, 26 units to be injected with her morning meal. LPN-B performed hand hygiene, retrieved R35's insulin flex pen from the medication cart, wiped the top and dialed the dose to match the physician ordered dose listed in the MAR. LPN-B then carried the flex-pen and a needle to R35's room where she attached the needle to the flex-pen. LPN-B then observed R35's abdomen to find a spot that was not</p>	F 759			

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F 759	<p>Continued From page 28</p> <p>bruised, cleansed the area and gently inserted the flex-pen needle into the flesh, and injected the dose. After about 2 seconds, and not more than 3 seconds she removed the needle from the abdomen, carried it to the medication cart to dispose of the needle, and performed hand hygiene. LPN-B stated she had received training on how to use an insulin flex-pen "a long time ago," but did not recall learning that the needle should be primed before setting the insulin dose. LPN-B was aware that the needle was supposed to remain in the tissue for "awhile" after injection, but said, "I usually just hold it until, I don't know." LPN-B stated she thought she had left the needle in place for five seconds.</p> <p>When interviewed on 2/2/22, at 8:40 a.m. the director of nursing (DON) stated the correct procedure when using an insulin flex-pen is to place the needle on the pen and to prime it with two units of insulin. After priming the pen, the nurse should check the order for the correct dose and set the dial to match the order. DON stated the needle should be inserted straight into the subcutaneous tissue before injecting the dose. After injection the needle should be removed and discarded and the pen re-capped.</p> <p>An undated facility procedure titled, Insulin Administration with Pen Device, included, "remove the pen cap, wipe the pen tip with an alcohol wipe, remove the protective seal from a new needle, screw the needle in place, dial a dose of 2 units to prime the pen, hold the pen with needle pointing straight up and tap lightly so the bubbles will rise to the top. Press the injection button all the way in and check to see that the insulin comes out of the needle (if no insulin comes out, repeat the test. If insulin still does not</p>	F 759			

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F 759	Continued From page 29 come out, get a new needle.) Check the order for the correct dose. Select the correct dose and dial until the number shows in the window." When administering insulin, the procedure included, "keep the pen straight and insert the needle into the skin. Using your thumb, press the injection button all the way down, when the number in the window returns to "0," slowly count to 10 before removing the needle."	F 759			
F 760 SS=D	Residents are Free of Significant Med Errors CFR(s): 483.45(f)(2) The facility must ensure that its- §483.45(f)(2) Residents are free of any significant medication errors. This REQUIREMENT is not met as evidenced by: Based on observation, interview and record review, improper injection technique was used for 2 of 3 residents (R44 and R35) who received insulin. R44 received 2 doses, and R35 received one, and the improper injection technique had the potential to result in the wrong dose of insulin being injected in all three cases. Findings include: R44's significant change Minimum Data Set (MDS) dated 1/14/22, included, cognitively intact, with diagnosis including diabetes and received insulin injections daily. R35's quarterly MDS dated 1/25/22, included severe cognitive impairment with diagnosis of diabetes and received insulin injections daily. On 2/1/22, 9:15 a.m. registered nurses (RN)-C was observed to prepare two insulin flex-pens for	F 760	The residents noted in findings R 35 and R 44 blood sugars were monitored post error with no noted adverse effects. 1. Education immediately started on insulin pen administration for nursing staff and continues 2. Audit hands on observation of the insulin administration technique RN-C and LPN-B weekly times 4 weeks, random audits for other nurses <input type="checkbox"/> weekly times four weeks 3. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. 4. DON or designee are responsible for compliance	3/15/22	

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F 760	<p>Continued From page 30</p> <p>R44's ordered breakfast insulin (a flex pen contains the vial of insulin inside the pen, and has a mechanism where the dose to be administered is set on a dial at the top of the pen, and only that amount can then be injected). R44 was to receive 7 units of Aspart insulin (a rapid acting insulin) twice daily with meals, and 32 units of Tresiba insulin (a long acting insulin) once each morning. RN-C performed hand hygiene, cleansed the top of the flex-pen with an alcohol wipe and attached a needle. RN-C then set the dial for each insulin to match the ordered dose in the medication administration record (MAR). RN-C failed to prime the needle (setting the flex-pen dial to 2 units and injecting that into the needle prior to setting the dose). RN-C ascertained R44's morning blood sugar and proceeded to R44's room. RN-C applied gloves, cleansed a spot on R44's abdomen where she injection the Aspart, and held the needle in place for 10 seconds and then gave the Tesiba, also on the right side of the abdomen in the same manner. RN-C stated she had received training in how to properly use a flex-pen when she was hired.</p> <p>On 2/2/22, 8:24 a.m. a licensed practical nurse (LPN)-B was observed in preparation of R35's Aspart insulin dose, 26 units to be injected with her morning meal. LPN-B performed hand hygiene, retrieved R35's insulin flex pen from the medication cart, wiped the top and dialed the dose to match the physician ordered dose listed in the MAR. LPN-B then carried the flex-pen and a needle to R35's room where she attached the needle to the flex-pen. LPN-B then observed R35's abdomen to find a spot that was not bruised, cleansed the area and gently inserted the flex-pen needle into the flesh, and injected the dose. After about 2 seconds, and not more than 3</p>	F 760			

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F 760	<p>Continued From page 31</p> <p>seconds she removed the needle from the abdomen, carried it to the medication cart to dispose of the needle, and performed hand hygiene. LPN-B stated she had received training on how to use an insulin flex-pen "a long time ago," but did not recall learning that the needle should be primed before setting the insulin dose. LPN-B was aware that the needle was supposed to remain in the tissue for "awhile" after injection, but said, "I usually just hold it until, I don't know." LPN-B stated she thought she had left the needle in place for five seconds.</p> <p>At 2/2/22, at 8:40 a.m. the director of nursing (DON) stated the correct procedure when using an insulin flex-pen is to place the needle on the pen and to prime it with two units of insulin. After priming the pen, the nurse should check the order for the correct dose and set the dial to match the order. DON stated the needle should be inserted straight into the subcutaneous tissue before injecting the dose. After injection the needle should be removed and discarded and the pen re-capped.</p> <p>When interviewed on 2/03/22, at 1:52 p.m. the DON stated using the wrong technique when giving insulin, especially if the needle of an insulin pen is not primed correctly, could result in the resident not receiving their ordered dose of medication. DON stated it could be of a significant concern if the blood sugar was running high after receiving the dose.</p> <p>According to a facility procedure titled Insulin Administration with Pen Device, no date, the proper procedure included "remove the pen cap, wipe the pen tip with an alcohol wipe, remove the protective seal from a new needle, screw the</p>	F 760			

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F 760	Continued From page 32 needle in place, dial a dose of 2 units to prime the pen, hold the pen with needle pointing straight up and tap lightly so the bubbles will rise to the top. Press the injection button all the way in and check to see that the insulin comes out of the needle (if no insulin comes out, repeat the test. If insulin still does not come out, get a new needle.)Check the order for the correct dose. Select the correct dose and dial until the number shows in the window." When administering insulin, the procedure included, "keep the pen straight and insert the needle into the skin. Using your thumb, press the injection button all the way down, when the number in the window returns to "0," slowly count to 10 before removing the needle."	F 760			
F 886 SS=D	COVID-19 Testing-Residents & Staff CFR(s): 483.80 (h)(1)-(6) §483.80 (h) COVID-19 Testing. The LTC facility must test residents and facility staff, including individuals providing services under arrangement and volunteers, for COVID-19. At a minimum, for all residents and facility staff, including individuals providing services under arrangement and volunteers, the LTC facility must: §483.80 (h)((1) Conduct testing based on parameters set forth by the Secretary, including but not limited to: (i) Testing frequency; (ii) The identification of any individual specified in this paragraph diagnosed with COVID-19 in the facility; (iii) The identification of any individual specified in this paragraph with symptoms consistent with COVID-19 or with known or suspected exposure to COVID-19;	F 886		3/15/22	

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F 886	<p>Continued From page 33</p> <p>(iv) The criteria for conducting testing of asymptomatic individuals specified in this paragraph, such as the positivity rate of COVID-19 in a county;</p> <p>(v) The response time for test results; and</p> <p>(vi) Other factors specified by the Secretary that help identify and prevent the transmission of COVID-19.</p> <p>§483.80 (h)((2) Conduct testing in a manner that is consistent with current standards of practice for conducting COVID-19 tests;</p> <p>§483.80 (h)((3) For each instance of testing:</p> <p>(i) Document that testing was completed and the results of each staff test; and</p> <p>(ii) Document in the resident records that testing was offered, completed (as appropriate to the resident's testing status), and the results of each test.</p> <p>§483.80 (h)((4) Upon the identification of an individual specified in this paragraph with symptoms consistent with COVID-19, or who tests positive for COVID-19, take actions to prevent the transmission of COVID-19.</p> <p>§483.80 (h)((5) Have procedures for addressing residents and staff, including individuals providing services under arrangement and volunteers, who refuse testing or are unable to be tested.</p> <p>§483.80 (h)((6) When necessary, such as in emergencies due to testing supply shortages, contact state and local health departments to assist in testing efforts, such as obtaining testing supplies or</p>	F 886			

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F 886	<p>Continued From page 34</p> <p>processing test results. This REQUIREMENT is not met as evidenced by: Based on observation, interview, and document review, the facility failed to ensure appropriate Personal Protective Equipment (PPE) was used by staff while collecting specimens for COVID-19 testing. The failure to appropriately collect and process COVID specimens affected 1 of 1 staff (ESD)that completed testing on 1/31/22.</p> <p>Findings include:</p> <p>During observation on 1/31/22, at 1:55 p.m. the front desk receptionist (R)-A was performing a COVID test. The environmental services director (ESD) handed the specimen collection card to the Receptionist after swabbing her nares and placing the swab in the testing card. R-A retrieved the testing card with her bare hands, placed the card on her desk, then wrote the name of the employee and time of the test on the card. Further observation revealed the eye goggles were located on the Receptionist's head and she was wearing prescription eyeglasses and a surgical face mask during the specimen collection.</p> <p>When interviewed on 1/31/22, at 5:01 p.m. the administrator stated, R-A did not don a gown, gloves, and eye protection prior to processing the COVID-19 testing card. The administrator stated R-A had been trained to don gloves, gown, and eye protection during the process. The administrator stated that the training and competency completed for R-A could not be found as the former infection preventionist (IP) was no longer employed at the facility.</p>	F 886	<p>COVID-19 Testing-Residents & Staff</p> <ol style="list-style-type: none"> 1. Staff (R)-A trained to test the community policies on infection control and COVID 19 testing. 2. Education of staff on COVID 19 and infection control related to PPE during testing 3. Review of staff testing procedures via competency for antigen testing 4. Audit testing weekly for 4 weeks and ensure proper testing procedures to include antigen testing and proper PPE use. 5. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. 6. NHA/DON or designee are responsible for compliance 		

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F 886	<p>Continued From page 35</p> <p>When interviewed on 2/1/22, at 10:17 a.m. R-A stated she had been trained by the former director of nursing (DON), and she was not aware that she should wear gloves, gown and eye protection during the specimen collection process. R-A stated it was important to wear the proper PPE to protect herself, staff, and residents from spreading the virus.</p> <p>When interviewed on 2/2/22, at 1:32 p.m. the DON, who also served as the facility's IP, stated she expected staff to follow the procedure for testing and don the proper PPE during the testing process. The DON stated, additional competency training was started on 1/31/22 with the R-A, the DON stated it was important to wear the proper PPE and sanitize the testing area to prevent the spread and transmission of COVID-19 to staff and residents in the facility.</p> <p>Review of the facility's undated document titled Competency for: Use of Binax-Now COVID-19 AG Card Rapid Antigen Testing, provided by the facility, included, "Specimen collection don appropriate PPE prior to collecting specimen."</p>	F 886			

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

PRINTED: 03/08/2022
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245409	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED 02/08/2022
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K 000	<p>INITIAL COMMENTS</p> <p>FIRE SAFETY</p> <p>An annual Life Safety recertification survey was conducted by the Minnesota Department of Public Safety, State Fire Marshal Division on 02/08/2022. At the time of this survey, Edenbrook of Rochester 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			

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

TITLE

(X6) DATE

Electronically Signed

02/24/2022

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 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>Edenbrook of Rochester is a 1-story building with a partial basement. The building was constructed at two different times. The original building was constructed in 1964 and was determined to be of Type II(111) construction. In 1974 an addition was constructed to the East wing that was determined to be of Type II(111) construction. The facility has a fire alarm system with smoke detection in the corridors and spaces open to the corridors, which is monitored for automatic fire department notification and the building is fully sprinkler</p>	K 000			

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K 000	Continued From page 2 protected. Because the original building and the 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 facility has a capacity of 81 beds and had a census of 43 at the time of the survey.	K 000			
K 211 SS=F	The requirement at 42 CFR, Subpart 483.70(a) is NOT MET as evidenced by: Means of Egress - General CFR(s): NFPA 101 Means of Egress - General Aisles, passageways, corridors, exit discharges, exit locations, and accesses are in accordance with Chapter 7, and the means of egress is continuously maintained free of all obstructions to full use in case of emergency, unless modified by 18/19.2.2 through 18/19.2.11. 18.2.1, 19.2.1, 7.1.10.1 This REQUIREMENT is not met as evidenced by: Based on observation and staff interview, the facility failed to maintain the means of egress per NFPA 101 (2012 edition), Life Safety Code, sections 7.1.3.2.3, 7.1.10.1, and 7.2.1.4.3.1. These deficient findings could have a widespread impact on the residents within the facility. Findings include: 1. On 02/08/2022 between 10:30 AM to 12:30 PM, observation revealed that the north and east fire door exits were impeded with snow and ice build-up.	K 211	<ul style="list-style-type: none"> ¿ Snow and ice build-up were removed from North and East fire door exits on 2/8/2022. ¿ Items being temporarily stored in the North stairwell were removed and placed in an alternative storage area. ¿ Maintenance and interdisciplinary team members were educated on the need for all facility egress exits to be clear of obstruction (including snow and ice build-up), and that stairwells will be free of equipment. ¿ Audits for clear fire exits and 	3/4/22	

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K 211	Continued From page 3 2. On 02/08/2022, between 10:30 AM to 12:30 PM, observation revealed that the north exit stairwell had items stored within the stairwell. These items included several wheelchairs, walkers, and mattresses. An interview with the Maintenance Director verified these findings at the time of discovery.	K 211	stairwells will be completed weekly for four weeks, and monthly thereafter. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. The Director of Maintenance will be responsible for compliance.		
K 324 SS=D	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 * cooking facilities in smoke compartments with 30 or fewer patients comply with conditions under 18.3.2.5.4, 19.3.2.5.4. 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. 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	K 324		3/4/22	

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K 324	Continued From page 4 This REQUIREMENT is not met as evidenced by: Based on observation and staff interview, the facility failed to maintain the commercial cooking fire suppression system per NFPA 101 (2012 edition), Life Safety Code, sections 19.3.2.5.1 and 9.2.3, and NFPA 96 (2011 edition), Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, section 10.5.1. This deficient finding could have a isolated impact on the residents within the facility. Findings include: On 02/08/2022 between 10:30AM to 12:30 PM, observation revealed access to the manual pull station for the Ansul fire suppression system in the Kitchen was blocked with a serving table and food carts. An interview with the Maintenance Director verified these findings at the time of discovery.	K 324	<ul style="list-style-type: none"> ¿ Pull-station for the Ansul fire suppression system in the kitchen was cleared of obstruction on 2/08/2022. ¿ All maintenance and kitchen staff have been educated that the pull-station for the fire suppression system must remain free of obstruction. ¿ Audits for keeping the pull-station clear of obstruction will be completed weekly for four weeks, and monthly thereafter. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. ¿ The Director of Maintenance will be responsible for compliance. 		
K 345 SS=F	Fire Alarm System - Testing and Maintenance CFR(s): NFPA 101 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 test and	K 345	<ul style="list-style-type: none"> ¿ Facility was able to find documentation from our fire test vendor, 	3/4/22	

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K 345	Continued From page 5 inspect smoke detectors throughout the facility per NFPA 101 (2012 edition), Life Safety Code, sections 9.6.1.3 and 9.6.1.5 and NFPA 72 (2010), National Fire Alarm and Signaling Code, sections 14.4.5.3 through 14.4.5.3.4. This deficient finding could have a widespread impact on the residents within the facility. Findings include: On 02/08/2022 between 10:30 AM to 12:30 PM, during a review of the available documentation it was revealed that documentation could not be located showing that a smoke detector sensitivity test had occurred within the required time period. Last sensitivity test was dated 09/14/2018. An interview with the Maintenance Director verified this finding at the time of discovery.	K 345	Custom Alarm, for an annual sensitivity test conducted on 9/1/2021. These results have been added into the facility fire safety binder. ¿ The Maintenance Director has been educated on the importance of obtaining sensitivity test results from our contracted vendor and keeping the information available in the facility's fire and safety binder. ¿ Audits for smoke detector sensitivity testing will be completed on a quarterly basis to ensure annual testing occurs and results are documented in the facility's fire and safety binder. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. ¿ The Director of Maintenance will be responsible for compliance		
K 353 SS=D	Sprinkler System - Maintenance and Testing CFR(s): NFPA 101 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. a) Date sprinkler system last checked _____ b) Who provided system test _____ c) Water system supply source _____	K 353		3/4/22	

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K 353	Continued From page 6 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 observation and staff interview, the facility failed maintain the fire sprinkler system per NFPA 101 (2012 edition), Life Safety Code, section 9.7.5 and NFPA 25 (2011 edition), Standard for the Inspection, Testing, and Maintenance of Water-Based Fire Protection Systems, section 5.2.2.2. This deficient finding could have an isolated impact on the residents within the facility. Findings include: On 02/08/2022 between 10.30 AM to 12:30 PM, observation revealed cabling engaging the sprinkler system in the Boiler Room. The cabling was zip-tied to sprinkler system piping. An interview with the Maintenance Director verified this finding at the time of discovery.	K 353	<ul style="list-style-type: none"> ¿ Cabling that was zip-tied to the sprinkler system piping was removed on 2/25/2022 ¿ Maintenance staff were educated on the need for the sprinkler system piping to remain clear and unobstructed by and cabling or other obtrusion. ¿ Audits for ensuring that the sprinkler system remains clear of obstruction or obtrusion will be completed weekly for four weeks, and monthly thereafter. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. ¿ The Director of Maintenance will be responsible for compliance. 		
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	K 712		3/4/22	

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K 712	Continued From page 7 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 19.7.1.6. These deficient finding could have a widespread impact on the residents within the facility. Findings include: 1. On 02/08/2022 between 10:30 AM to 12:30 PM, during a review of the available fire drill reports, it was revealed that there was no first quarter, second shift fire drill conducted. 2. On 02/08/2022 between 10:30 AM to 12:30 PM, during a review of the available fire drill reports, it was revealed that there were no second and third quarter, third shift fire drills conducted. An interview with the Maintenance Director verified this finding at the time of discovery.	K 712	<ul style="list-style-type: none"> ¿ On 02/23/2022 a Q1 fire drill was completed on the PM shift. ¿ A schedule for fire drills has been constructed for the remainder of the year, which includes conducting drills quarterly on each shift. ¿ Maintenance and interdisciplinary team members were educated on the need for quarterly fire drills to be completed on each shift. ¿ Audits for ensuring fire drills are completed quarterly on each shift will be completed monthly. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. ¿ The Director of Maintenance will be responsible for compliance. 		
K 761 SS=F	Maintenance, Inspection & Testing - Doors CFR(s): NFPA 101 Maintenance, Inspection & Testing - Doors Fire doors assemblies are inspected and tested annually in accordance with NFPA 80, Standard for Fire Doors and Other Opening Protectives. Non-rated doors, including corridor doors to patient rooms and smoke barrier doors, are routinely inspected as part of the facility maintenance program. Individuals performing the door inspections and	K 761		3/4/22	

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K 761	Continued From page 8 testing possess knowledge, training or experience that demonstrates ability. Written records of inspection and testing are maintained and are available for review. 19.7.6, 8.3.3.1 (LSC) 5.2, 5.2.3 (2010 NFPA 80) This REQUIREMENT is not met as evidenced by: Based on a review of available documentation and staff interview, the facility failed to test and inspect fire door assemblies per NFPA 101 (2012 edition), Life Safety Code, section 8.3.3.1, and NFPA 80 (2010 edition), Standard for Fire Doors and Other Opening Protectives, section 5.2.1. This deficient finding could have a widespread impact on the residents within the facility. Findings include: On 02/08/2022 between 10:30 AM to 12:30 PM, a review of the available documentation revealed that no evidence could be provided to show that an annual fire door inspection and testing had occurred. An interview with the Maintenance Director verified this finding at the time of discovery.	K 761	<ul style="list-style-type: none"> ¿ On 02/23/2022 an annual inspection of all fire doors was completed, and all doors were fully operational. ¿ Maintenance staff were educated on the requirement for all fire doors to be inspected and tested annually. ¿ Audits for fire door inspections will be completed monthly to ensure all doors are in compliance. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring. ¿ The Director of Maintenance will be responsible for compliance. 		
K 916 SS=F	Electrical Systems - Essential Electric System CFR(s): NFPA 101 Electrical Systems - Essential Electric System Alarm Annunciator A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer	K 916		3/15/22	

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K 916	Continued From page 9 system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) This REQUIREMENT is not met as evidenced by: Based on observation and staff interview, the facility failed to install an essential electric system alarm annunciator per NFPA 99 (2012 edition), Health Care Facilities Code, section 6.4.1.1.17. This deficient finding could have a widespread impact on the residents within the facility. Findings include: On 02/08/2022 between 10:30 AM to 12:30 PM, observation revealed that a remote electric system alarm annunciator could not be located within the facility. An interview with the Maintenance Director verified this finding at the time of discovery.	K 916	Waiver being requested for this work. Waiver is attached in ePOC system. Completion date is not a set date refer to waiver.		
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	K 918		3/4/22	

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K 918	<p>Continued From page 10</p> <p>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 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 and inspect the emergency generator per NFPA 99 (2012 edition), Health Care Facilities Code, section 6.4.4.1.1.3 and NFPA 110 (2010 edition), Standard for Emergency and Standby Power Systems, section 8.4.1. This deficient finding could have a widespread impact on the residents within the facility.</p> <p>Findings include:</p> <p>On 02/08/2022 between 10:30 AM to 12:30 PM, a review of the available documentation revealed that the generator had not been exercised under load for 30 minutes during the months of November and December, 2021 and January,</p>	K 918	<p>¿ On 02/23/2022 the emergency generator was exercised under load for 30 minutes. Monthly testing will be conducted for future months.</p> <p>¿ Maintenance will be educated on the requirement for generator testing on a monthly basis.</p> <p>¿ Audits for testing and inspecting the generator under load for 30 minutes will be completed monthly. All findings will be brought to the facility's next QAPI meeting for review and recommendation for continuance of monitoring.</p> <p>¿ The Director of Maintenance will be responsible for compliance.</p>		

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

PRINTED: 03/08/2022
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245409	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED 02/08/2022
NAME OF PROVIDER OR SUPPLIER EDENBROOK OF ROCHESTER			STREET ADDRESS, CITY, STATE, ZIP CODE 1875 19TH STREET NORTHWEST ROCHESTER, MN 55901		
(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 918	Continued From page 11 2022. An interview with the Maintenance Director verified these findings at the time of discovery.	K 918			



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 14, 2022

Administrator
Edenbrook Of Rochester
1875 19th Street Northwest
Rochester, MN 55901

Re: State Nursing Home Licensing Orders
Event ID: 3YDJ11

Dear Administrator:

The above facility was surveyed on January 31, 2022 through February 3, 2022 for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules and Statutes. At the time of the survey, the survey team from the Minnesota Department of Health - Health Regulation Division noted one or more violations of these rules or statutes that are issued in accordance with Minn. Stat. § 144.653 and/or Minn. Stat. § 144A.10. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a civil fine for each deficiency not corrected shall be assessed in accordance with a schedule of fines promulgated by rule and/or statute of the Minnesota Department of Health.

To assist in complying with the correction order(s), a "suggested method of correction" has been added. This provision is being suggested as one method that you can follow to correct the cited deficiency. Please remember that this provision is only a suggestion and you are not required to follow it. Failure to follow the suggested method will not result in the issuance of a penalty assessment. You are reminded, however, that regardless of the method used, correction of the order within the established time frame is required. The "suggested method of correction" is for your information and assistance only.

You have agreed to participate in the electronic receipt of State licensure orders consistent with the Minnesota Department of Health Informational Bulletin 14-01, available at https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html. The State licensing orders are delineated on the Minnesota Department of Health State Form and are being delivered to you electronically. The Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Tag numbers have been assigned to Minnesota state statutes/rules for Nursing Homes.

The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule number and the corresponding text of the state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction order. This column also includes the findings that are in violation of the state statute or rule after the statement, "This MN Requirement is not met as evidenced by." Following the surveyors findings are

Edenbrook Of Rochester

February 14, 2022

Page 2

the Suggested Method of Correction and the Time Period For Correction.

PLEASE DISREGARD THE HEADING OF THE FOURTH COLUMN WHICH STATES, "PROVIDER'S PLAN OF CORRECTION." THIS APPLIES TO FEDERAL DEFICIENCIES ONLY. THIS WILL APPEAR ON EACH PAGE.

THERE IS NO REQUIREMENT TO SUBMIT A PLAN OF CORRECTION FOR VIOLATIONS OF MINNESOTA STATE STATUTES/RULES.

Although no plan of correction is necessary for State Statutes/Rules, please enter the word "corrected" in the box available for text. You must then indicate in the electronic State licensure process, under the heading completion date, the date your orders will be corrected prior to electronically submitting to the Minnesota Department of Health. We urge you to review these orders carefully, item by item, and if you find that any of the orders are not in accordance with your understanding at the time of the exit conference following the survey, you should immediately contact:

Karen Aldinger, Unit Supervisor
St. Cloud A District Office
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
3333 Division Street, Suite 212
Saint Cloud, Minnesota 56301-4557
Email: karen.aldinger@state.mn.us
Office: (651) 201-3794 Mobile: (320) 249-2805

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.

Please note it is your responsibility to share the information contained in this letter and the results of this visit with the President of your facility's Governing Body.

Please feel free to call me with any questions.



Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64900
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: melissa.poepping@state.mn.us

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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NAME OF PROVIDER OR SUPPLIER EDENBROOK OF ROCHESTER	STREET ADDRESS, CITY, STATE, ZIP CODE 1875 19TH STREET NORTHWEST ROCHESTER, MN 55901
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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 1/3/22 through 2/3/22, a licensing survey was conducted at your facility by surveyors from the Minnesota Department of Health (MDH). Your facility was found NOT in compliance with the MN State Licensure and the following correction orders are issued. Please indicate in your electronic plan of correction you have reviewed</p>	2 000		

Minnesota Department of Health
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

Electronically Signed

TITLE

(X6) DATE
02/23/22

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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2 000	<p>Continued From page 1</p> <p>these orders and identify the date when they will be completed. In addition, complaint investigations were completed.</p> <p>The following complaints were found to be substantiated with no deficiencies cited due to actions implemented by the facility prior to survey. H5409104C/MN80488 H5409105C/MN80202 H5409109C/MN72703</p> <p>The following complaints were found to be UNSUBSTANTIATED: H5409106C/MN79143 H5409107C/MN76266 H5409108C/MN75190 H5409110C/MN71280 H5409111C/MN70882 H5409112C/MN65322</p> <p>Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Tag numbers have been assigned to Minnesota state statutes/rules for Nursing Homes. The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction order. This column also includes the findings which are in violation of the state statute after the statement, "This Rule is not met as evidence by." Following the surveyors findings are the Suggested Method of Correction and Time period for Correction.</p> <p>You have agreed to participate in the electronic receipt of State licensure orders consistent with the Minnesota Department of Health Informational Bulletin</p>	2 000		

Minnesota Department of Health

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2 000	Continued From page 2 https://www.health.state.mn.us/facilities/regulation/infobulletins/ib14_1.html The State licensing orders are delineated on the attached Minnesota Department of Health orders being submitted to you electronically. Although no plan of correction is necessary for State Statutes/Rules, please enter the word "corrected" in the box available for text. You must then indicate in the electronic State licensure process, under the heading completion date, the date your orders will be corrected prior to electronically submitting to the Minnesota Department of Health. PLEASE DISREGARD THE HEADING OF THE FOURTH COLUMN WHICH STATES, "PROVIDER'S PLAN OF CORRECTION." THIS APPLIES TO FEDERAL DEFICIENCIES ONLY. THIS WILL APPEAR ON EACH PAGE. THERE IS NO REQUIREMENT TO SUBMIT A PLAN OF CORRECTION FOR VIOLATIONS OF MINNESOTA STATE STATUTES/RULES.	2 000		
2 830	MN Rule 4658.0520 Subp. 1 Adequate and Proper Nursing Care; General Subpart 1. Care in general. A resident must receive nursing care and treatment, personal and custodial care, and supervision based on individual needs and preferences as identified in the comprehensive resident assessment and plan of care as described in parts 4658.0400 and 4658.0405. A nursing home resident must be out of bed as much as possible unless there is a written order from the attending physician that the resident must remain in bed or the resident prefers to remain in bed.	2 830		3/15/22

Minnesota Department of Health

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2 830	<p>Continued From page 3</p> <p>This MN Requirement is not met as evidenced by: Based on observation, interview, and document review, the facility failed to monitor the condition of 1 of 1 residents (R4) with lymphedema and failed to follow-up on orders provided for R4 to receive edema care.</p> <p>Findings include:</p> <p>R4's 5 day perspective payment system (PPS) Minimum Data Set (MDS) dated 1/17/21, included, cognitively intact, required extensive assistance for dressing, bathing and grooming and limited assistance of 1 person for wheel chair mobility, and had limited range of motion of left upper extremity. R4's diagnoses included, a malignant neoplasm of connective and soft tissue of left upper limb, including the shoulder. Edema was not marked on the MDS.</p> <p>During observation and interview on 1/31/22, at 2:43 p.m. R4 was observed to have significant swelling of her left upper arm, with the area appearing to be nearly twice the size of her right arm. R4 had an ace wrap on her lower left arm that had slid down and was wrinkled, bunched and sloppy. R4 stated she wrapped the arm herself, but said a nurse would wrap it if she requested this. R4 also stated that therapy used to wrap it for her, but she was no longer in therapy and had to do the care herself. R4 stated she has had lymphedema for many years, but that her upper arm was now larger than it had ever been.</p> <p>During observation on 2/1/22, at 11:13 a.m. R4 was in her room and stated her arm had not been wrapped yet today.</p>	2 830	Acknowledged	

Minnesota Department of Health

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2 830	<p>Continued From page 4</p> <p>R4's physician orders for February 2022 did not show an order to monitor edema or to wrap her left upper extremity.</p> <p>R4's care plan did not contain information regarding her lymphedema or any directions for care or monitoring of this condition.</p> <p>A hand written form in R4's medical record indicated she had been seen 1/6/22, for left lymphedema consultation, and ordered recommendations were written for physical or occupation therapy (PT or OT) to wrap R4's left upper extremity fingers, hand and arm with short stretch wrap daily and leave in place for 23 hours per day and then rewrap.</p> <p>A hospital discharge summary dated 1/19/22, indicated R4 had lymphedema for over twenty years.</p> <p>A daily skilled charting note dated 1/20/22, indicated R4 had edema of her left upper extremity.</p> <p>When interviewed on 2/1/22, at 11:38 a.m. occupational therapist (OT)-A stated she was not lymphedema certified but had wrapped R4's arm a few times. OT-A did not recall receiving an order for PT or OT to wrap R4's arm, and said she was also unaware that no orders had been provided to the nursing staff to provide edema care. OT-A stated that even if R4 wrapped her arm herself, the facility was responsible to make sure the wrap was in place and correctly applied.</p> <p>When interviewed on 2/3/22, at 9:46 a.m. licensed practical nurse (LPN)-C stated she was aware that one of R4's arms was, "very swollen,"</p>	2 830		

Minnesota Department of Health

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2 830	<p>Continued From page 5</p> <p>but she did not provide any treatment for the edema. LPN-C said she thought perhaps PT wrapped the arm, but she did not know. LPN-C was not able to describe any nursing interventions a nurse could take independently to provide care for edematous extremities and said she had been taught to go by orders, and since there were never any treatments orders for R4's edema she had not done anything. She did not recall R4 asking her to wrap her arm.</p> <p>On 2/3/22, at 11:25 a.m. the director of nursing (DON) stated R4's edema should be on her diagnosis list, and should have been listed on her care plan. DON stated nursing staff should be monitoring edema and note if it is a chronic condition, and monitor for changes, updating the medical provider as needed. DON stated edema monitoring and any orders to wrap an extremity should be listed on the medication or treatment administration order sheets so a nurse would be reminded to monitor the resident's condition. DON also said a nurse noting a resident with edema, but not seeing any orders for care should clarify this with the medical provider. DON confirmed that R4 had not had edema listed on her care plan, and also confirmed the order for lymphedema wraps from 1/6/22 had not been followed up on.</p> <p>A policy specific to edema care and monitoring was not provided, but a facility policy titled Resident Assessment and Examination dated 8/1/15 indicated residents should be assessed for abnormalities in health states in order that the care team could implement interventions to address the concern. The policy indicated a body systems review, which included edema, was to be performed and any abnormalities reported to the physician.</p>	2 830		

Minnesota Department of Health

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2 830	Continued From page 6 SUGGESTED METHOD OF CORRECTION: The director of nursing (DON) or designee could audit orders for residents with fluid imbalance or lymphedema issues and ensure they are complete. DON or designee could ensure all nursing staff receive education on fluid balance issues, edema monitoring and documentation. Audits could be done to ensure compliance of proper nursing interventions, edema monitoring, and proper reporting. TIME PERIOD FOR CORRECTION: Twenty one (21) days.	2 830		
21426	MN St. Statute 144A.04 Subd. 3 Tuberculosis Prevention And Control (a) A nursing home provider must establish and maintain a comprehensive tuberculosis infection control program according to the most current tuberculosis infection control guidelines issued by the United States Centers for Disease Control and Prevention (CDC), Division of Tuberculosis Elimination, as published in CDC's Morbidity and Mortality Weekly Report (MMWR). This program must include a tuberculosis infection control plan that covers all paid and unpaid employees, contractors, students, residents, and volunteers. The Department of Health shall provide technical assistance regarding implementation of the guidelines. (b) Written compliance with this subdivision must be maintained by the nursing home.	21426		3/15/22

Minnesota Department of Health

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21426	<p>Continued From page 7</p> <p>This MN Requirement is not met as evidenced by: Based on interview and document review, the facility failed to follow their tuberculosis prevention plan and standards of practice when 4 of 6 residents (R4, R22, R39, and R45) were not properly screened for tuberculosis (TB) upon admission to the facility, and 6 of 6 employees (NA-C, NA-D, DP-A, DP-B, RN-B and EM-A) were not properly screened for TB when hired. This had the potential to affect all 44 residents and staff in the facility.</p> <p>Findings include:</p> <p>The facility policy titled Tuberculosis Surveillance and Control, last revised 5/25/21, indicated the following guidelines were to be followed for residents:</p> <ol style="list-style-type: none"> 1. All residents new to long-term care or assisted living facility who do not have documentation of a previous skin test reaction >10 mm or a history of adequate treatment of tuberculosis infection or disease, previous documented negative interferon gamma release assay (IGR) blood test within past 90 days, or tuberculin skin test (TST) within past 90 days shall have the initial test of a Mantoux PPD [purified protein derivative] two -step skin test to rule out tuberculosis within 72 hours of admission. 2. If the initial result is 0-9 mm, the second test, which can be given after admission, should be given at least on week and no more than three weeks after the first test. <p>In addition, the policy indicated, "it is important to also perform an evaluation to determine if signs</p>	21426	Acknowledged	

Minnesota Department of Health

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21426	<p>Continued From page 8</p> <p>or symptoms of tuberculosis (unexplained weight loss, fever, persistent cough) are present. Once tuberculosis disease is ruled out, it is important to record the results of the skin test in millimeters (mm), in a prominent place ..."</p> <p>The policy guidelines for employees were:</p> <ol style="list-style-type: none"> 1. Initial Examination. Provide a tuberculin skin test (Mantoux, 5 tuberculin units (TU) of purified protein derivative (PPD) to all employees during pre-employment procedures, unless a previous reaction >10 mm is documented. If the initial skin test result is 0-9 mm, a second test should be given at least one week and no more than three weeks after the first test. 4. Repeat Chest X-ray. After the initial evaluation of persons with skin test reaction >10 mm, routine repeated chest x-rays are not recommended. They are not a substitute for infection treatment nor vigilance for signs and symptoms of tuberculin disease. An annual sign and symptom review should be documented in their record. <p>R4 had a TB symptom screening completed upon admission, 10/19/21, and the medical record showed an initial "step 1" TST (tuberculin skin test) was performed at that time. The results were recorded two days later as negative, "0 mm"; however, no second step TST was recorded as having been done.</p> <p>R22 had a TB symptom screening completed upon admission, 12/20/21, and the medical record showed an initial, "step 1" TST was performed at that time. The results were recorded as negative two days later without any corresponding measurement in millimeters, but no second step TST was recorded as having been done.</p>	21426		

Minnesota Department of Health

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21426	<p>Continued From page 9</p> <p>R39 had a TB symptom screening completed upon admission, 5/14/21, and R45's medical record showed a step 1 TST was performed on the same day, but no results were documented. A step 2 TST was performed 14 days later, and documented as negative and 0 mm, two days later.</p> <p>R45 had a TB symptom screening completed upon admission, 12/22/21, but R45's medical record did not show a step 1 TST was performed within 72 hours of admission. A single TB TST was performed 14 days after admission and marked as "step 2." R45's medication administration record (MAR) indicated the results were read two days later and were documented as "negative" without any corresponding measurement in millimeters.</p> <p>A nursing assistant (NA)-C provided the facility a copy of a chest X-ray TB screening done 19 months (5/4/20) prior to being hired by the facility, 12/13/21; however, the facility failed to screen NA-C at the time of hire for any signs or symptoms of tuberculosis.</p> <p>NA-D completed a TB screening tool upon hire, 10/27/21, and a step 1 TST test was given on the same day plus recorded as negative with 0 mm induration two days later. A second step TST was not documented as having been performed.</p> <p>A dietary personnel (DP)-A, completed a TB screening tool upon hire, 12/6/21, and a step 1 TST was given on the same day. No results of the test were recorded, and no step 2 TST was documented as having been performed.</p> <p>DP-B completed a TB screening tool upon hire, 11/23/21, and a step 1 TST test was given on the</p>	21426		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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NAME OF PROVIDER OR SUPPLIER EDENBROOK OF ROCHESTER	STREET ADDRESS, CITY, STATE, ZIP CODE 1875 19TH STREET NORTHWEST ROCHESTER, MN 55901
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21426	<p>Continued From page 10</p> <p>same day plus recorded as negative with 0.0 mm induration two days later. A second step TST was not documented as having been performed.</p> <p>A registered nurse (RN)-B, completed a TB screening tool upon hire, 1/5/22, and a step 1 TST was given on the same day. No results of the test were recorded, and no step 2 TST was documented as having been performed.</p> <p>On 2/3/22, at approximately 11:30 a.m., an employee (EM)-A, stated the facility must have lost his TB screening from when he was hired, 10/11/21. EM-A presented a TB screening form that was completed, but dated as 2/4/22, and the form showed EM-A had just received a step 1 TST on 2/3/22 at 10:58 a.m.</p> <p>During an interview 2/3/22, 2:26 p.m. the director of nursing (DON) stated she was new to the facility, but she expected to be the one in charge of the TB program. DON stated she had been reaching out to nurses who had documented results for residents as only "negative" to see if they could recall the mm of induration. DON stated it was her expectation that TB be given as written, and results recorded to include the mm induration as indicated in the MAR. DON stated employees should be screened and tested upon hire.</p> <p>SUGGESTED METHOD OF CORRECTION:</p> <p>The director of nursing (DON) or designee could audit resident and employee files for incomplete TB testing and ensure the testing gets done. DON or designee could provide training to all staff related to TB, TB testing and documentation and follow the training with further audits to ensure</p>	21426		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21426	Continued From page 11 compliance with facility plan. TIME PERIOD FOR CORRECTION: Twenty one (21) days.	21426		
21545	MN Rule 4658.1320 A.B.C Medication Errors A nursing home must ensure that: A. Its medication error rate is less than five percent as described in the Interpretive Guidelines for Code of Federal Regulations, title 42, section 483.25 (m), found in Appendix P of the State Operations Manual, Guidance to Surveyors for Long-Term Care Facilities, which is incorporated by reference in part 4658.1315. For purposes of this part, a medication error means: (1) a discrepancy between what was prescribed and what medications are actually administered to residents in the nursing home; or (2) the administration of expired medications. B. It is free of any significant medication error. A significant medication error is: (1) an error which causes the resident discomfort or jeopardizes the resident's health or safety; or (2) medication from a category that usually requires the medication in the resident's blood to be titrated to a specific blood level and a single medication error could alter that level and precipitate a reoccurrence of symptoms or toxicity. All medications are administered as prescribed. An incident report or medication error report must be filed for any medication error that occurs. Any significant medication errors or resident reactions must be reported to the physician or the physician's designee and the resident or the resident's legal guardian or designated representative and an explanation	21545		3/15/22

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21545	<p>Continued From page 12</p> <p>must be made in the resident's clinical record.</p> <p>C. All medications are administered as prescribed. An incident report or medication error report must be filed for any medication error that occurs. Any significant medication errors or resident reactions must be reported to the physician or the physician's designee and the resident or the resident's legal guardian or designated representative and an explanation must be made in the resident's clinical record.</p> <p>This MN Requirement is not met as evidenced by: Based on observation, interview and record review, facility failed to ensure medications were passed without an error rate of 5% or less when improper injection technique was used for 2 of 3 residents (R44, R35) who received insulin. R44 received 2 doses, and R35 received one, and the improper injection technique had the potential to result in the wrong dose of insulin being injected in all three cases.</p> <p>Findings include:</p> <p>According to a Minimum Data Set (MDS) significant change assessment dated 1/14/22, R44 scored 15/15 on a cognitive assessment indicating she was cognitively intact. The MDS indicated R44 had a diagnosis of Diabetes Mellitus and received insulin injections.</p> <p>According to an MDS quarterly review dated 1/25/22, R35 scored 3/15 on a cognitive assessment indicating severe cognitive impairments. The MDS indicated R35 had a diagnosis of Diabetes Mellitus and received insulin injections.</p>	21545	Acknowledged	

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21545	<p>Continued From page 13</p> <p>On 2/1/22, 9:15 a.m. a registered nurses (RN)-C was observed to prepare two insulin "flex-pens" for R44's ordered breakfast insulin (a flex pen contains the vial of insulin inside the pen, and has a mechanism where the dose to be administered is set on a dial at the top of the pen, and only that amount can then be injected). R44 was to receive 7 units of Aspart insulin (a rapid acting insulin) twice daily with meals, and 32 units of Tresiba insulin (a long acting insulin) once each morning. RN-C performed hand hygiene, cleansed the top of the flex-pen with an alcohol wipe and attached a needle. RN-C then set the dial for each insulin to match the ordered dose in the medication administration record (MAR). RN-C failed to prime the needle (setting the flex-pen dial to 2 units and injecting that into the needle prior to setting the dose). RN-C ascertained R44's morning blood sugar and proceeded to R44's room. RN-C applied gloves, cleansed a spot on R44's abdomen where she injection the Aspart, and held the needle in place for 10 seconds and then gave the Tesiba, also on the right side of the abdomen in the same manner. RN-C stated she had received training in how to properly use a flex-pen when she was hired.</p> <p>On 2/2/22, 8:24 a.m. a licensed practical nurse (LPN)-B was observed in preparation of R35's Aspart insulin dose, 26 units to be injected with her morning meal. LPN-B performed hand hygiene, retrieved R35's insulin flex pen from the medication cart, wiped the top and dialed the dose to match the physician ordered dose listed in the MAR. LPN-B then carried the flex-pen and a needle to R35's room where she attached the needle to the flex-pen. LPN-B then observed R35's abdomen to find a spot that was not bruised, cleansed the area and gently inserted</p>	21545		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21545	<p>Continued From page 14</p> <p>the flex-pen needle into the flesh, and injected the dose. After about 2 seconds, and not more than 3 seconds she removed the needle from the abdomen, carried it to the medication cart to dispose of the needle, and performed hand hygiene. LPN-B stated she had received training on how to use an insulin flex-pen "a long time ago," but did not recall learning that the needle should be primed before setting the insulin dose. LPN-B was aware that the needle was supposed to remain in the tissue for "awhile" after injection, but said, "I usually just hold it until ...I don't know." LPN-B stated she thought she had left the needle in place for five seconds.</p> <p>At 2/2/22, 8:40 a.m. the director of nursing (DON) stated the correct procedure when using an insulin flex-pen is to place the needle on the pen and to prime it with two units of insulin. After priming the pen, the nurse should check the order for the correct dose and set the dial to match the order. DON stated the needle should be inserted straight into the subcutaneous tissue before injecting the dose. After injection the needle should be removed and discarded and the pen re-capped.</p> <p>According to a facility procedure titled Insulin Administration with Pen Device, not dated, the proper procedure included, "remove the pen cap, wipe the pen tip with an alcohol wipe, remove the protective seal from a new needle, screw the needle in place, dial a dose of 2 units to prime the pen, hold the pen with needle pointing straight up and tap lightly so the bubbles will rise to the top. Press the injection button all the way in and check to see that the insulin comes out of the needle (if no insulin comes out, repeat the test. If insulin still does not come out, get a new needle.) Check the order for the correct dose. Select the correct dose</p>	21545		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21545	Continued From page 15 and dial until the number shows in the window." When administering insulin, the procedure included, "Keep the pen straight and insert the needle into the skin. Using your thumb, press the injection button all the way down, when the number in the window returns to '0,' slowly count to 10 before removing the needle." SUGGESTED METHOD OF CORRECTION: The Director of Nursing (DON) or designee could review and educate nursing staff on the correct use of insulin pens. DON or designee could audit administration of insulin by nursing staff to ensure procedures are implemented as written. TIME PERIOD FOR CORRECTION: Twenty one (21) days.	21545		
21880	MN St. Statute 144.651 Subd. 20 Patients & Residents of HC Fac.Bill of Rights Subd. 20. Grievances. Patients and residents shall be encouraged and assisted, throughout their stay in a facility or their course of treatment, to understand and exercise their rights as patients, residents, and citizens. Patients and residents may voice grievances and recommend changes in policies and services to facility staff and others of their choice, free from restraint, interference, coercion, discrimination, or reprisal, including threat of discharge. Notice of the grievance procedure of the facility or program, as well as addresses and telephone numbers for the Office of Health Facility Complaints and the area nursing home ombudsman pursuant to the Older Americans Act, section 307(a)(12) shall be posted in a conspicuous place.	21880		3/15/22

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21880	<p>Continued From page 16</p> <p>Every acute care inpatient facility, every residential program as defined in section 253C.01, every nonacute care facility, and every facility employing more than two people that provides outpatient mental health services shall have a written internal grievance procedure that, at a minimum, sets forth the process to be followed; specifies time limits, including time limits for facility response; provides for the patient or resident to have the assistance of an advocate; requires a written response to written grievances; and provides for a timely decision by an impartial decision maker if the grievance is not otherwise resolved. Compliance by hospitals, residential programs as defined in section 253C.01 which are hospital-based primary treatment programs, and outpatient surgery centers with section 144.691 and compliance by health maintenance organizations with section 62D.11 is deemed to be compliance with the requirement for a written internal grievance procedure.</p> <p>This MN Requirement is not met as evidenced by: Based on observation, interview and document review, the facility failed to follow its grievance policy for 1 of 1 residents (R21) reviewed for missing personal items.</p> <p>Findings include:</p> <p>R21's quarterly Minimum Data Set (MDS) dated 12/24/21, included cognitively intact.</p> <p>When interviewed on 1/31/22, at 12:18 p.m. R21 stated she had been missing a shirt and several</p>	21880	Acknowledged	
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Minnesota Department of Health

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21880	<p>Continued From page 17</p> <p>other items of clothing for over two and a half weeks. R21 stated, she felt as though she had reported it to everyone in laundry services, but did not feel as though anything was being done. At that time, a housekeeper (HK)-B entered the room, and R21 was observed to repeat her report of missing items to HK-B. HK-B said, "yeah, maybe someone put it in the wrong closet." R21 asked HK-B if they couldn't look in other resident closets as she was sure she had seen her gray Vikings shirt on a rack going down the hall at some time. HK-B stated, "I guess I could make a suggestion to my supervisor."</p> <p>When interviewed on 2/1/22, at 10:42 a.m. HK-A stated HK-B had told her about R21's concern of missing clothing on 1/31/22. HK-A recalled she had received an undated, unsigned note providing a description of the missing items, but it had just been laying on the desk and she was unsure for how long. HK-A stated that after HK-B told her about the missing clothing on 1/31/22, she had looked in the lost and found, and then told R21, "in a couple days I will look in the closets." HK-A stated she had not filled out any report or grievance.</p> <p>When interviewed on 2/1/22, at 10:42 a.m. the environmental services director (ESD) stated a grievance form should have been filled out and given it to the facility social worker (SW). ESD stated the proper procedure for missing items was for the person receiving the report to fill out the grievance form, hand it to the SW who would take note, then bring to ESD. The ESD would initiate a search for any missing items and complete the grievance form with the findings of their investigation, and return it to the SW. ESD stated any resident or staff could get a grievance form from the nurses station or from the SW.</p>	21880		

Minnesota Department of Health

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21880	<p>Continued From page 18</p> <p>ESD stated there had been plenty of training to staff about the process.</p> <p>On 2/01/22, 10:59 a.m. a grievance form related to R21's missing items was observed laying on the SW's desk. The SW stated, "I got the grievance sheet about two minutes before you walked in the door. [The ESD] filled it out after talking to the surveyor." SW stated an expectation for staff to fill out a grievance form at the time any resident should have a concern such as missing personal items.</p> <p>According to a facility policy titled Grievance/Concern last revised 1/14/22:</p> <ol style="list-style-type: none"> 1. Facility will make prompt efforts to resolve all grievances. 2. Facility will provide this policy and form to residents and have them readily available in common areas of the facility, areas may include facility entrance, reception desk, nurses' station, etc. 3. Facility staff will immediately report all alleged violations 8. Facility will use the information gathered in a grievance investigation to prevent further potential violations of any resident rights. <p>SUGGESTED METHOD OF CORRECTION:</p> <p>Facility administrator or social service designee could ensure all staff receive training in the facility grievance procedure, and the procedure could be reviewed at resident council meetings. Additionally, a reminder regarding how the forms are readily available could be sent out to residents and/or family. Administrator or social service designee could initiate audits of grievances to ensure that the policy and procedure have been fully implemented.</p>	21880		

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00916	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 02/03/2022
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21880	Continued From page 19 TIME PERIOD FOR CORRECTION: Twenty one (21) days.	21880		