



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 24, 2021

Administrator
Bayshore Residence & Rehab Ctr
1601 St Louis Avenue
Duluth, MN 55802

RE: CCN: 245227
Cycle Start Date: September 24, 2020

Dear Administrator:

On October 12, 2020, we notified you a remedy was imposed. On February 22, 2021 the Minnesota Department of Health completed a revisit to verify that your facility had achieved and maintained compliance. We have determined that your facility has achieved substantial compliance as of February 19, 2021.

As authorized by CMS the remedy of:

- Discretionary denial of payment for new Medicare and Medicaid admissions effective October 27, 2020 be discontinued as of February 19, 2021. (42 CFR 488.417 (b))

However, as we notified you in our letter of October 12, 2020, in accordance with Federal law, as specified in the Act at § 1819(f)(2)(B)(iii)(I)(b) and § 1919(f)(2)(B)(iii)(I)(b), we notified you that your facility is prohibited from conducting Nursing Aide Training and/or Competency Evaluation Programs (NATCEP) for two years from September 24, 2020. This does not apply to or affect any previously imposed NATCEP loss.

The CMS Region V Office may notify you of their determination regarding any imposed remedies.

Feel free to contact me if you have questions.

Sincerely,

A handwritten signature in black ink, appearing to read 'Joanne Simon', with a long horizontal line extending to the right.

Joanne Simon, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program
Program Assurance Unit
Health Regulation Division
Telephone: 651-201-4161 Fax: 651-215-9697
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 3, 2021

Administrator
Bayshore Residence & Rehab Ctr
1601 St Louis Avenue
Duluth, MN 55802

RE: CCN: 245227
Cycle Start Date: September 24, 2020

Dear Administrator:

On October 12, 2020, we informed you of imposed enforcement remedies.

On January 19, 2021, the Minnesota Department of Health completed a revisit and it has been determined that your facility continues to not to be in substantial compliance. The most serious deficiencies in your facility were found to be widespread deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level F), as evidenced by the electronically attached CMS-2567, whereby corrections are required.

The deficiency not corrected is as follows:

F0886 -- S/S: F -- 483.80 (h)(1)-(6) -- Covid-19 Testing-Residents & Staff

As a result of the revisit findings:

- Discretionary Denial of Payment for new Medicare and/or Medicaid Admissions, Federal regulations at 42 CFR § 488.417(a), effective October 27, 2020, will remain in effect.

This Department continues to recommend that CMS impose a civil money penalty. (42 CFR 488.430 through 488.444). You will receive a formal notice from the CMS RO only if CMS agrees with our recommendation.

The CMS Region V Office will notify your Medicare Administrative Contractor (MAC) that the denial of payment for new admissions is effective October 27, 2020. They will also notify the State Medicaid Agency that they must also deny payment for new Medicaid admissions effective October 27, 2020.

You should notify all Medicare/Medicaid residents admitted on, or after, this date of the restriction. The remedy must remain in effect until your facility has been determined to be in substantial compliance or your provider agreement is terminated. Please note that the denial of payment for new admissions includes Medicare/Medicaid beneficiaries enrolled in managed care plans. It is your obligation to inform managed care plans contracting with your facility of this denial of payment for new admissions.

An equal opportunity employer.

As we notified you in our letter of October 12, 2020, in accordance with Federal law, as specified in the Act at Section 1819(f)(2)(B)(iii)(I)(b) and 1919(f)(2)(B)(iii)(I)(b), your facility is prohibited from conducting Nursing Aide Training and/or Competency Evaluation Programs (NATCEP) for two years from September 24, 2020.

ELECTRONIC PLAN OF CORRECTION (ePOC)

Within ten (10) calendar days after your receipt of this notice, you must submit an acceptable plan of correction (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. The failure to submit an acceptable ePOC can lead to termination of your Medicare and Medicaid participation (42 CFR 488.456(b)).

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.

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:

- Optional denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417 (a));
- Per day civil money penalty (42 CFR 488.430 through 488.444).

DEPARTMENT CONTACT

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

**Teresa Ament, Unit Supervisor
Duluth District Office
Licensing and Certification Program
Health Regulation Division**

Minnesota Department of Health
Duluth Technology Village
11 East Superior Street, Suite 290
Duluth, Minnesota 55802-2007
Email: teresa.ament@state.mn.us
Phone: (218) 302-6151

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 - Health Regulation Division staff and/or the Department of Public Safety, State Fire Marshal Division staff, if your ePoC for their respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

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

If substantial compliance has been achieved, certification of your facility in the Medicare and/or Medicaid program(s) will be continued and remedies will not be imposed. Compliance is certified as of the latest correction date on the approved ePoC, unless it is determined that either correction actually occurred between the latest correction date on the ePoC and the date of the first revisit, or correction occurred sooner than the latest correction date on the ePoC.

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

We will also recommend to the CMS Region V Office and/or the Minnesota Department of Human Services that your provider agreement be terminated by March 24, 2021 (six months after the identification of noncompliance) if your facility does not achieve substantial compliance. 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.

APPEAL RIGHTS

If you disagree with this action imposed on your facility, you or your legal representative may request a

Bayshore Residence & Rehab Ctr

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hearing before an administrative law judge of the Department of Health and Human Services, Departmental Appeals Board (DAB). Procedures governing this process are set out in 42 C.F.R. 498.40, et seq. You must file your hearing request electronically by using the Departmental Appeals Board's Electronic Filing System (DAB E-File) at <https://dab.efile.hhs.gov> no later than sixty (60) days after receiving this letter. Specific instructions on how to file electronically are attached to this notice. A copy of the hearing request shall be submitted electronically to:

Tamika.Brown@cms.hhs.gov

Requests for a hearing submitted by U.S. mail or commercial carrier are no longer accepted as of October 1, 2014, unless you do not have access to a computer or internet service. In those circumstances you may call the Civil Remedies Division to request a waiver from e-filing and provide an explanation as to why you cannot file electronically or you may mail a written request for a waiver along with your written request for a hearing. A written request for a hearing must be filed no later than sixty (60) days after receiving this letter, by mailing to the following address:

**Department of Health & Human Services
Departmental Appeals Board, MS 6132
Director, Civil Remedies Division
330 Independence Avenue, S.W.
Cohen Building – Room G-644
Washington, D.C. 20201
(202) 565-9462**

A request for a hearing should identify the specific issues, findings of fact and conclusions of law with which you disagree. It should also specify the basis for contending that the findings and conclusions are incorrect. At an appeal hearing, you may be represented by counsel at your own expense. If you have any questions regarding this matter, please contact Tamika Brown, Principal Program Representative by phone at (312) 353-1502 or by e-mail at Tamika.Brown@cms.hhs.gov.

INFORMAL DISPUTE RESOLUTION/ 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

Bayshore Residence & Rehab Ctr

February 3, 2021

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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 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.

Feel free to contact me if you have questions.

Sincerely,



Joanne Simon, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program
Program Assurance Unit
Health Regulation Division
Telephone: 651-201-4161 Fax: 651-215-9697
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
December 17, 2020

Administrator
Bayshore Residence & Rehab Ctr
1601 St Louis Avenue
Duluth, MN 55802

RE: CCN: 245227
Cycle Start Date: September 24, 2020

Dear Administrator:

On October 12, 2020, we informed you of imposed enforcement remedies.

On November 25, 2020, the Minnesota Department(s) of Health completed a survey and it has been determined that your facility continues to not to be in substantial compliance. Your facility was not in substantial compliance with the participation requirements and the conditions in your facility which constituted **immediate jeopardy** to resident health or safety. The most serious deficiencies in your facility were found to be a pattern of deficiencies that constituted immediate jeopardy (Level K), as evidenced by the electronically attached CMS-2567, whereby corrections are required.

REMOVAL OF IMMEDIATE JEOPARDY

On November 25, 2020, the situation of immediate jeopardy to potential health and safety cited at F886 was removed. However, continued non-compliance remains at the lower scope and severity of E.

As a result of the survey findings:

- Discretionary Denial of Payment for new Medicare and/or Medicaid Admissions, Federal regulations at 42 CFR § 488.417(a), effective October 27, 2020, will remain in effect.
- Directed plan of correction, Federal regulations at 42 CFR § 488.424 Please see electronically attached documents for the DPOC.

This Department continues to recommend that CMS impose a civil money penalty. (42 CFR 488.430 through 488.444).

You will receive a formal notice from the CMS RO only if CMS agrees with our recommendation.

The CMS Region V Office will notify your Medicare Administrative Contractor (MAC) that the denial of payment for new admissions is effective October 27, 2020. They will also notify the State Medicaid Agency that they must also deny payment for new Medicaid admissions effective October 27, 2020.

You should notify all Medicare/Medicaid residents admitted on, or after, this date of the restriction. The remedy must remain in effect until your facility has been determined to be in substantial compliance or your provider agreement is terminated. Please note that the denial of payment for new admissions includes Medicare/Medicaid beneficiaries enrolled in managed care plans. It is your obligation to inform managed care plans contracting with your facility of this denial of payment for new admissions.

As we notified you in our letter of October 12, 2020, in accordance with Federal law, as specified in the Act at Section 1819(f)(2)(B)(iii)(I)(b) and 1919(f)(2)(B)(iii)(I)(b), your facility is prohibited from conducting Nursing Aide Training and/or Competency Evaluation Programs (NATCEP) for two years from September 24, 2020.

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

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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:

- Optional denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417 (a));
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DEPARTMENT CONTACT

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Cohen Building – Room G-644
Washington, D.C. 20201
(202) 565-9462**

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INFORMAL DISPUTE RESOLUTION/ 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:

Bayshore Residence & Rehab Ctr
December 17, 2020
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Nursing Home Informal Dispute Process
Minnesota Department of Health
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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 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.

Feel free to contact me if you have questions.

Sincerely,



Joanne Simon, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program
Program Assurance Unit
Health Regulation Division
Telephone: 651-201-4161 Fax: 651-215-9697
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File

DIRECTED PLAN OF CORRECTION

A Directed Plan of Correction (DPOC) is imposed in accordance with 42 CFR § 488.424. Your facility must include the following in their POC for the deficient practice cited at F880:

ACTIVE SCREENING

- Address how corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- Address how the facility will identify other residents having the potential to be affected by the same deficient practice.

POLICIES/PROCEDURES/SYSTEM CHANGES:

- The facility's Quality Assurance and Performance Improvement Committee must conduct a root cause analysis (RCA) to identify the problem(s) that resulted in this deficiency and develop intervention or corrective action plan to prevent recurrence.

The Infection Preventionist and Director of Nursing, shall complete the following:

- Develop and implement procedures, policies, and forms regarding active screening for temperature and signs and symptoms of COVID-19, in accordance with CDC guidelines to be conducted at the point of entry for every person who enters the facility. The procedures and policy must restrict entrance to anyone who does not meet the criteria as outlined by the CDC. This procedure must include actively measuring and recording staff temperature and assessment of shortness of breath, new or changed cough, and sore throat. The results must be documented. The MDH COVID-19 Toolkit <https://www.health.state.mn.us/diseases/coronavirus/hcp/ltctoolkit.pdf> has examples of forms to utilize for staff screening.

TRAINING/EDUCATION:

As part of a corrective action plan, the facility must provide training for Infection Preventionist and all other staff who enter the facility, as well as staff responsible for the screening. The training must cover the need for active screening. The CDC has training videos available for COVID-19 which may be utilized, Training for Healthcare Professionals; <https://www.cdc.gov/coronavirus/2019-ncov/hcp/training.html> and the MDH COVID-19 Toolkit may be utilized.

- Include documentation of the completed training with a timeline for completion.
- The training may be provided by the Director of Nursing, Infection Preventionist, or Medical Director with an attestation statement of completion.

CDC RESOURCES:

Infection Control Guidance: <https://www.cdc.gov/coronavirus/2019-ncov/hcp/infection-control.html>

CDC: Isolation Precautions Guideline:

<https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html>

CDC: Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings (2007): <https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html>

CDC: Personal Protective Equipment: <https://www.cdc.gov/niosh/ppe/>

Healthcare Infection Prevention and Control FAQs for COVID-19:

https://www.cdc.gov/coronavirus/2019-ncov/hcp/faq.html?CDC_AA_refVal=https%3A%2F%2Fwww.cdc.gov%2Fcoronavirus%2F2019-ncov%2Fhcp%2Finfection-control-faq.html

MDH RESOURCES:

Personal Protective Equipment (PPE) for Infection Control:

<https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/ppe/index.html>

MDH Contingency Standards of Care for COVID-19: Personal Protective Equipment for Congregate Care Settings (PDF): <https://www.health.state.mn.us/communities/ep/surge/crisis/ppegrid.pdf>

Interim Guidance on Facemasks as a Source Control Measure (PDF):

<https://www.health.state.mn.us/diseases/coronavirus/hcp/maskssource.pdf>

Interim Guidance on Alternative Facemasks (PDF):

<https://www.health.state.mn.us/diseases/coronavirus/hcp/masksalt.pdf>

Aerosol-Generating Procedures and Patients with Suspected or Confirmed COVID-19

(PDF): <https://www.health.state.mn.us/diseases/coronavirus/hcp/aerosol.pdf>

Droplet Precautions:

<https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html>

Airborne Precautions:

<https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html>

MONITORING/AUDITING:

- The Director of Nursing, the Infection Preventionist, and other facility leadership will conduct audits on all shifts, four times a week for one week, twice weekly for one week and biweekly thereafter, until 100% compliance is achieved to ensure active screening is being completed at the point of entry for all persons who enter the facility.

The Director of Nursing, Infection Preventionist or designee will review the results of audits and monitoring with the Quality Assurance Program Improvement (QAPI) program.

In accordance with 42 CFR § 488.402(f), the DPOC remedy is effective 15 calendar days from the date of the enforcement letter. The DPOC may be completed before or after that date. A revisit will not be approved prior to receipt of documentation confirming the DPOC was completed. To successfully complete the DPOC, the facility must provide all of the following documentation identified in the chart below.

Documentation must be uploaded as attachments through ePOC to ensure you have completed this remedy.

Imposition of this DPOC does not replace the requirement that the facility must submit a complete POC for all cited deficiencies (including F880) within 10 days after receipt of the Form CMS 2567.

Item	Checklist: Documents Required for Successful Completion of the Directed Plan
1	Documentation of the RCA and intervention or corrective action plan based on the results with signatures of the QAPI Committee members.
2	Documentation that the interventions or corrective action plan that resulted from the RCA was fully implemented
3	Content of the training provided to staff, including a syllabus, outline, or agenda, as well as any other materials used or provided to staff for the training
4	Names and positions of all staff that attended and took the trainings
5	Staff training sign-in sheets
6	Summary of staff training post-test results, to include facility actions in response to any failed post-tests
7	Documentation of efforts to monitor and track progress of the interventions or corrective action plan

In order to speed up our review, identify all submitted documents with the number in the “Item” column.