



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered

December 23, 2021

Administrator
Robbinsdale A Villa Center
3130 Grimes Avenue North
Robbinsdale, MN 55422

RE: CCN: 245417
Cycle Start Date: December 16, 2021

Dear Administrator:

On December 16, 2021, 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 isolated deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level D), 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.

An equal opportunity employer.

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:

Annette Winters, Rapid Response Unit Supervisor
Metro 1, Golden Rule Office
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
85 East Seventh Place, Suite 220
P.O. Box 64900
Saint Paul, Minnesota 55164-0900
Email: annette.m.winters@state.mn.us
Mobile: (651) 558-7558

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

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

VERIFICATION OF SUBSTANTIAL COMPLIANCE

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

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

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

If substantial compliance with the regulations is not verified by March 16, 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 June 16, 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

Robbinsdale A Villa Center

December 23, 2021

Page 4

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,

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

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

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

PRINTED: 09/21/2022
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245417	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 12/16/2021
NAME OF PROVIDER OR SUPPLIER ROBBINSDALE A VILLA CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 3130 GRIMES AVENUE NORTH ROBBINSDALE, MN 55422		
(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
F 000	INITIAL COMMENTS On 12/15/2021 to 12/16/2021 a standard abbreviated survey was conducted at your facility. 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 complaint was found to be SUBSTANTIATED: H5417202C (MN00079188, MN00079106, and MN00079103) with deficiencies cited at F760. 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 that substantial compliance with the regulations has been attained.	F 000			
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 interview and record review the facility failed to ensure an antiviral medication was administered as prescribed for 1 of 3 residents (R1) reviewed for medications. When R1's antiviral medication for the treatment human immunodeficiency virus (HIV) was not received	F 760	The submission of this Plan of Correction is not an admission by the provider of any fact or conclusion set forth in the Statement of Deficiency. This Plan of Correction is being submitted because it is required by law. However,		1/11/22

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

TITLE

(X6) DATE

Electronically Signed

01/04/2022

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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F 760	Continued From page 1 from the pharmacy the facility failed to remedy the omission. R1's Admission Record printed 4/5/21, indicated R1's diagnoses included Human Immunodeficiency virus (HIV) disease, syncope and collapse, compartment syndrome bilateral lower extremities, renal disease, chronic obstructive pulmonary disease, asthma, major depressive disorder, chronic pain, anxiety disorder, muscle wasting and atrophy, and unsteadiness on feet. R1's medical order sheet printed 12/15/2021, listed the following pharmacy order: Juluca Tablet 50-25 milligrams (mg's), (dolutegravir-rilpivirine). Give 1 tablet by mouth one time a day for antiviral, HIV - D/C Date- 12/09/2021. Order written 12/2/2021, to be started on 12/3/2021 at 08:00 a.m. Ordering Physician was listed as MD-A Review of R1's medication administration record (MAR) for December 2021, revealed a column on the left side of the page where the order was transcribed. A grid follows to the right containing entry opportunities for several times during each date. A legend near the bottom labeled "Chart Codes" depicted numerical codes and their significance. The code "8" indicated, "8=Other/See Nurses Notes." R1's MAR for the month of December depicted the following: in the row labeled "Juluca Tablet 50-25 MG (Dolutegravir-rilpivirine). Give 1 tablet by mouth one time a day for antiviral, HIV." The MAR depicted the following: The entry for 12/3/2021, at 0800, a code "8" was entered with the initials "BNT4,"	F 760	evidencing Robbinsdale A Villa Center good faith, the facility offers the following Plan of Correction. 1.R1 no longer resides at facility, discharged against medical advice 12/7/2021. 2.All residents with medications have the potential to be affected by this practice and medication administration record has been reviewed for missed medications and follow up being completed 3.Education has been completed on Medication administration policy with licensed nursing staff. 4.Audits will be completed by the DON and/or designee on 3 residents twice weekly to ensure medications are being ordered and administered. Will be reviewed at Quality Assurance Meeting (QAPI) monthly to determine if any trends are identified and recommendations made for continued audits and monitoring needs. 5.Completion date: 01/11/2022		

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F 760	Continued From page 2 the entry for 12/4/2021, at 0800, a code of "8" was entered with the initials "SYP0," the entry for 12/5/2021, at 0800, a code of "8" was entered with the initials "Onwu," the entry for 12/6/2021, at 0800, a code of "8" was entered with the initials "BNT4," and the entry for 12/7/2021, at 0800 a code of "8" was entered with the initials " BNT4." R1's Progress Notes copied on 12/6/2021, depicted the following: On 12/3/2021, at 07:48 Author LPN-A entered, "Juluca Tablet 50-25 MG Give 1 tablet by mouth one time a day for antiviral, HIV not available." On 12/4/2021, there was no entry describing pertaining to Juluca. On 12/5/2021, at 09:34 Author LPN-B entered, ""Juluca Tablet 50-25 MG Give 1 tablet by mouth one time a day for antiviral, HIV. Medication not available to give, on order. Resident asked for medication and concern of not having his HIV med." On 12/6/2021, at 07:12 Author LPN-A entered, "Juluca Tablet 50-25 MG Give 1 tablet by mouth one time a day for antiviral, HIV. Med not available, on order." On 12/7/2021, at 07:55 Author LPN-A entered, "Juluca Tablet 50-25 MG Give 1 tablet by mouth one time a day for antiviral, HIV. Awaiting for med to arrive." On 12/16/21, at 12:22 p.m. the DON was interviewed. DON stated no one at the facility reported R1's prescription for Juluca was not received from the pharmacy. Stated was notified of the expense of the medication and authorized the expense by contacting the pharmacy and sending an authorization form. Stated received no indication the medication was unavailable to the	F 760			

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F 760	Continued From page 3 pharmacy and would not be dispensed to the facility. DON stated had spoken to registered nurse pharmacy liaison (RNPL-D) earlier on 12/16/21. RNPL-D reported to the DON the pharmacy received the order for R1's Juluca, and the authorization for expense form, but the medication was not available, was on back order, and never dispensed to the facility. DON stated RNPL-D should have contacted facility stating the prescription could not be filled, but the facility was never notified. DON stated RNPL-D admitted pharmacy personnel did not, and should have, notified facility Juluca was not available. On 12/16/2021, at 1:37 p.m. NP-B stated she was not notified that R1 was not receiving Juluca as prescribed. Stated this drug was an anti-viral medication used to maintain his HIV status. Stated R1 was not compromised from his HIV at the current time. Stated it would be very difficult to prove the absence of this medication caused R1 any harm. R1 was at home after leaving facility against medical advice. Stated the medical group was no longer primary physicians for R1. On 12/16/21, at 2:08 p.m. MD-C stated had reviewed the medical record for R1. Stated that the Juluca was started to take the place of Biktarvy. Stated R1 was the anti-viral properties were established with Biktarvy. This medication was stopped because of the potential damage to R1's already compromised renal system. Stated that missing the Juluca doses would not cause any harm to the resident as R1 was already stable on the Biktarvy. The facility policy, "IIA2: Medication Administration-General Guidelines," Dated April 2018, Section D, number 6 directed, "If a dose of	F 760			

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F 760	Continued From page 4 regularly scheduled medication is withheld, refused, not available, or given at a time other than the scheduled time {e.g., the resident is not in the facility at scheduled dose time, or a starter dose of antibiotic is needed), the space provided on the front of the MAR for that dosage administration is (initialed and circled). If electronic MAR is used, documentation of the unadministered dose is done as instructed by the procedures for use of the eMAR system. An explanatory note is entered on the reverse side of the record. If [XX consecutive doses] of a vital medication are withheld, refused, or not available the physician is notified. Nursing documents the notification and physician response." The facility policy, "1B1: Non-Controlled Medication Order Documentation," Dated August 2019, Section D directed, "The prescriber is contacted by nursing for direction when delivery of a medication will be delayed or the medication is not or will not be available."	F 760			



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
December 23, 2021

Administrator
Robbinsdale A Villa Center
3130 Grimes Avenue North
Robbinsdale, MN 55422

Re: Event ID: TF5G11

Dear Administrator:

The above facility survey was completed on December 16, 2021 for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules. At the time of the survey, the survey team from the Minnesota Department of Health - Health Regulation Division noted no violations of these rules promulgated under Minnesota Stat. section 144.653 and/or Minnesota Stat. Section 144A.10.

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

Please feel free to call me with any questions.

Sincerely,

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

Joanne Simon, Enforcement Specialist
Minnesota Department of Health
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

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00122	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED C 12/16/2021
NAME OF PROVIDER OR SUPPLIER ROBBINSDALE A VILLA CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 3130 GRIMES AVENUE NORTH ROBBINSDALE, MN 55422		
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2 000	Initial Comments *****ATTENTION***** NH LICENSING CORRECTION ORDER 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. 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. 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. INITIAL COMMENTS: On 12/15/21 to 12/16/21 a complaint survey was conducted at your facility by surveyors from the Minnesota Department of Health (MDH). Your facility was found IN compliance with the MN State Licensure. The following complaint was found to be	2 000		

Minnesota Department of Health

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

TITLE

(X6) DATE

Electronically Signed

01/04/22

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00122	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED C 12/16/2021
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2 000	Continued From page 1 SUBSTANTIATED: H5417202C (MN00079188, MN00079106, and MN00079103), however NO licensing orders were issued. Minnesota Department of Health is documenting the State Licensing Correction Orders using Federal software. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of state form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.	2 000			