



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered
August 14, 2024

Administrator
Martin Luther Care Center
1401 East 100th Street
Bloomington, MN 55425

RE: CCN: 245272
Cycle Start Date: June 21, 2024

Dear Administrator:

On July 31, 2024, the Minnesota Department of Health completed a revisit to verify that your facility had achieved and maintained compliance. Based on our review, we have determined that your facility has achieved substantial compliance; therefore no remedies will be imposed.

Feel free to contact me if you have questions.

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

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



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
June 28, 2024

Administrator
Martin Luther Care Center
1401 East 100th Street
Bloomington, MN 55425

RE: CCN: 245272
Cycle Start Date: June 21, 2024

Dear Administrator:

On June 21, 2024, a survey was completed at your facility by the Minnesota Department of Health 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.

The state agency may, in lieu of an onsite revisit, determine correction and compliance by accepting

Martin Luther Care Center

June 28, 2024

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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, Regional Operations Supervisor, Federal Rapid Response
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
625 Robert Street North
P.O. Box 64975
Saint Paul, Minnesota 55164-0975
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 September 21, 2024 (three months after the identification of noncompliance), the CMS Region V Office must deny payment for new admissions as mandated by the Social Security Act (the Act) at Sections 1819(h)(2)(D) and 1919(h)(2)(C) and Federal regulations at 42 CFR Section 488.417(b).

In addition, if substantial compliance with the regulations is not verified by December 21, 2024 (six months after the identification of noncompliance) your provider agreement will be terminated. This action is mandated by the Social Security Act at Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

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

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

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

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

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

Martin Luther Care Center

June 28, 2024

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Feel free to contact me if you have questions.

Sincerely,

A handwritten signature in black ink that reads "Kamala Fiske-Downing". The signature is written in a cursive style with a loop at the end of the last name.

Kamala Fiske-Downing

Federal Enforcement | Health Regulation Division

Minnesota Department of Health

Health Regulation Division

Telephone: (651) 201-4112

Email: Kamala.Fiske-Downing@state.mn.us



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered

June 28, 2024

Administrator
Martin Luther Care Center
1401 East 100th Street
Bloomington, MN 55425

Re: Event ID: BB8C11

Dear Administrator:

The above facility survey was completed on June 21, 2024 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 that reads 'Kamala Fiske-Downing'.

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

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00227	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 06/21/2024
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NAME OF PROVIDER OR SUPPLIER MARTIN LUTHER CARE CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 1401 EAST 100TH STREET BLOOMINGTON, MN 55425
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(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETE DATE
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2 000	<p>Initial Comments</p> <p style="text-align: center;">*****ATTENTION*****</p> <p style="text-align: center;">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 6/20/24 -6/21/24, a complaint survey was conducted at your facility by surveyors from the Minnesota Department of Health (MDH). Your facility was IN compliance with the MN State Licensure</p> <p>The following complaints were reviewed during</p>	2 000		
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Minnesota Department of Health
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

Electronically Signed

TITLE

(X6) DATE

07/03/24

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00227	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 06/21/2024
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2 000	Continued From page 1 the survey: H52724723C (MN104115) 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		

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

PRINTED: 07/08/2024
FORM APPROVED
OMB NO. 0938-0391

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F 000	INITIAL COMMENTS On 6/20/24 - 6/21/24, a standard abbreviated survey was conducted at your facility. Your facility was NOT in compliance with the requirements of 42 CFR 483, Subpart B, Requirements for Long Term Care Facilities. The following complaints were reviewed. H52724723C (MN104115) with deficiencies issued at F609 and F755 The facility's plan of correction (POC) will serve as your allegation of compliance upon the Departments 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 609 SS=D	Reporting of Alleged Violations CFR(s): 483.12(b)(5)(i)(A)(B)(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		7/26/24

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 07/03/2024
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Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

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F 609	<p>Continued From page 1</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 record review the facility failed to report an allegation neglect to the State Agency (SA) within 24 hours for 1 of 1 resident (R1). R1 was given the incorrect medications at the facility, which required R1 to be sent to the hospital for bradycardia (low heart rate), pain and anxiety.</p> <p>Findings include:</p> <p>R1's care plan dated 6/12/24 indicated R1's diagnoses included pneumonia, shortness of breath, saddle embolus of the pulmonary artery with acute cor pulmonale (a large blood clot that lodges in the pulmonary artery obstructing blood flow to both lungs), chronic pain, stage 3 kidney disease, hypertension (high blood pressure), cerebral infarction (stroke), hemiplegia and hemiparesis affecting the left side (weakness), osteoarthritis, and depression.</p>	F 609	<p>All staff within the facility will be re-educated on the vulnerable adult policy, including how every staff member in the facility is a mandated reporter. All previous medication errors were reviewed and reported as necessary per federal guidelines. Facility will audit all incidents and allegation of neglect twice per week for two months. Audits will be presented to the Quality Assurance Performance Improvement (QAPI) committee to determine if additional monitoring is necessary. The correction will be monitored by the campus administrator or designee.</p>	

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F 609	<p>Continued From page 2</p> <p>R1's medication/treatment error report dated 6/15/24 at 9:15 a.m. indicated R1 was given Keppra (an anticonvulsant), metoprolol (a beta blocker to treat high blood pressure), and clopidogrel (a blood thinner). The description of the error indicated the nurse on the cart pulled out a patient's medications and did not check the right room. She entered the room and gave the medications to R1 not knowing she was in the wrong room. The provider was notified and gave orders to monitor R1's blood pressure, pulse, and respirations every 30 minutes, monitor for sedation and bleeding. R1 was to be hydrated and hold his morning Gabapentin (indicated for nerve pain, apixaban (blood thinner), methocarbamol (a muscle relaxant) and assess before the next dose.</p> <p>R1's progress note dated 6/15/24 at 1:17 p.m. indicated R1 had taken the wrong medications. His vital signs were within normal range expect his heart rate. The on-call provider was notified and R1 was sent to the hospital for close evaluation.</p> <p>R1's Hospital Emergency Department after visit summary dated 6/15/24 indicated R1 was seem for 1. drug overdose: multidrug. 2. Metoprolol organ tablet. The reason for R1's visit was he was given the wrong medications. His diagnosis was medication reaction. Labs taken were a complete blood culture (CBC) with platelets and differential and a complete metabolic panel. Imaging Tests included a 12-lead electrocardiography (EKG). R1 was given Tylenol, Lidocaine (for irregular heart rate), and Oxycodone was given twice for pain.</p> <p>R1's admission Minimum Data Set (MDS) dated</p>	F 609		

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F 609	<p>Continued From page 3</p> <p>6/18/24 indicated R1's Brief Inventory for Mental Status (BIMs) score was 15 indicating R1 was cognitively intact. R1's MDS did not indicate his diagnoses.</p> <p>Upon interview on 6/20/24 R1 stated he was given the wrong medications at the facility. He stated his stomach began to hurt and he left like he was going "throw-up." He stated he had never been "so terrified" in his life, stating he felt like his chest was going to collapse. R1 stated he did not know if the feeling in his chest was due to the incorrect medications being administered or if the medications were hurting him.</p> <p>Upon interview on 6/20/24 at 3:05 p.m. registered nurse RN-A, the unit manager stated R1 was given another resident's medications on 6/15/24 which was a Saturday. She stated she was not aware of the incident until 6/17/24 the following Monday upon reporting to work. She stated she was not certain why the incident was not reported to the SA over the weekend when R1 required hospitalization.</p> <p>Upon interview on 6/20/24 at 4:12 p.m. family member (FM)-A stated she was visiting R1 shortly after the medication error occurred. She stated the medication error caused him "great anxiety," which was still lingering as R1 does not trust the staff at the facility anymore. She stated she felt like she could not trust the facility as well since the medication error caused his heart rate to drop 46 beat per minute. She stated the medication error keeps her awake at night.</p> <p>Upon interview on 6/21/24 at 1:15 p.m. RN-B, the nursing educator stated the medication error should have been reported. She stated her</p>	F 609		

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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 609	<p>Continued From page 4</p> <p>rationale was R1 was exhibiting a low heart rate of 46 and a change in sedation. RN-B stated, "if anything prompts them to be sent out it would be a VA" (vulnerable adult report).</p> <p>Upon interview on 6/21/24 at 1:37 p.m. the Administrator stated she chose not to report the medication error incident at the time because R1 did not have serious bodily harm.</p> <p>A facility policy titled Vulnerable Adult - Adult Prohibition Plan dated 10/6/22 indicated Mandated reporters in skilled nursing facilities ensure that all alleged violations involving abuse, neglect, exploitation or mistreatment, including injuries of unknown source and misappropriation of resident property, are reported, and a report made 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 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 in accordance with State law through established procedures. The report will be made to the Minnesota Department of Health (MDH)/OHFC. To identify and meet this VAA reporting obligation, follow these procedures: Neglect Is the failure of the facility, its employees or service providers to provide goods and services to a resident that are necessary to avoid physical harm pain, mental anguish, or emotional distress.</p>	F 609		
F 755 SS=D	<p>Pharmacy Srvcs/Procedures/Pharmacist/Records CFR(s): 483.45(a)(b)(1)-(3)</p> <p>§483.45 Pharmacy Services</p>	F 755		7/26/24

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F 755	<p>Continued From page 5</p> <p>The facility must provide routine and emergency drugs and biologicals to its residents, or obtain them under an agreement described in §483.70(g). The facility may permit unlicensed personnel to administer drugs if State law permits, but only under the general supervision of a licensed nurse.</p> <p>§483.45(a) Procedures. A facility must provide pharmaceutical services (including procedures that assure the accurate acquiring, receiving, dispensing, and administering of all drugs and biologicals) to meet the needs of each resident.</p> <p>§483.45(b) Service Consultation. The facility must employ or obtain the services of a licensed pharmacist who-</p> <p>§483.45(b)(1) Provides consultation on all aspects of the provision of pharmacy services in the facility.</p> <p>§483.45(b)(2) Establishes a system of records of receipt and disposition of all controlled drugs in sufficient detail to enable an accurate reconciliation; and</p> <p>§483.45(b)(3) Determines that drug records are in order and that an account of all controlled drugs is maintained and periodically reconciled. This REQUIREMENT is not met as evidenced by: Based on observation, interview, and record review the facility failed to provide pharmaceutical services to meet the needs for 1 of 4 residents (R4) reviewed for medication administration. R4 had an over-the-counter medication on his tray table that he had been taking for approximately two weeks, the facility failed to monitor his intake</p>	F 755	<p>All licensed nursing staff within the facility will receive education regarding the medication storage policy. They will also review the medications self-administration policy. Residents will be assessed for medications self-administration upon admission. All other previous medication</p>	

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NAME OF PROVIDER OR SUPPLIER MARTIN LUTHER CARE CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 1401 EAST 100TH STREET BLOOMINGTON, MN 55425		
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F 755	<p>Continued From page 6 of the medication. This medication had an interaction with a prescription medication R4 was taking.</p> <p>Finding include:</p> <p>R4's self-administration of medication assessment dated 6/8/24 indicated:</p> <ul style="list-style-type: none"> -Required assistance for storing medications in a secure location. -Required assistance for opening and closing medication containers. -Could count not accurately tell time to know when medications need to be taken. -R4 did understand that skipping a medication dose is a refusal and staff will be notified when refusal has occurred. -Required assistance administering eye drops/ointments, topical lotions, ear drops, suppositories, subcutaneous injections, nasal sprays, and oral medications. -Required assistance naming his medications and their prescribed use. -Fully capable of reading the labels for medications. -Required assistance to identify common side effects of medications. -Required assistance with time of medications, dosage, proper amount, and documentation of self-administration of medications. -Required assistance with being able to identify when needing a prn (as needed) medication. <p>R4 was not approved for self-administration of medications or to keep medications at bedside.</p> <p>R4's care plan dated 6/12/24 did not indicate R4 was able to self-administer any medications.</p> <p>R4's admission Minimum Data Set (MDS) dated</p>	F 755	<p>storage errors have been reviewed and reported per federal guidelines. Facility will audit resident rooms for inappropriate medication storage and self-administration assessments three times a week for four weeks. Audits will be presented to the Quality Assurance Performance Improvement (QAPI) committee to determine if additional monitoring is necessary. The correction will be monitored by the director of nursing or designee.</p>	

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245272	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 06/21/2024
NAME OF PROVIDER OR SUPPLIER MARTIN LUTHER CARE CENTER		STREET ADDRESS, CITY, STATE, ZIP CODE 1401 EAST 100TH STREET BLOOMINGTON, MN 55425		
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F 755	<p>Continued From page 7</p> <p>6/14/24 indicated R1 had a Brief Inventory for Mental Status (BIMs) score of 15 indicating R1 was cognitively intact. R1's diagnoses were Rhabdomyolysis (breakdown of muscle tissue that releases damaging protein into the blood), unspecified fall, unspecified symptoms and signs of cognitive functions and awareness, abnormalities with gait and mobility, and Type 2 Diabetes Mellitus.</p> <p>Upon observation and medication review on 6/21/24 at 8:57 a.m. R4's medication Rosuvastatin 20 mg (a statin to treat high cholesterol) card indicated to not take aluminum, magnesium, or antacids within two hours of the medication. Registered nurse (RN)-C administered all R4's morning medications including the Rosuvastatin. On R4's table was a bottle of over-the-counter Roloids. RN-C asked R4 if the facility knew he had the Roloids. R4 stated that the facility was aware. RN-C did not question R4 when he had taken the Roloids last or how often he takes them. RN-C did not remove the Roloids from R4's room.</p> <p>Upon interview on 6/21/24 at 9:10 a.m. RN-C stated it was fine for R4 to have the Roloids in his room since the facility knew about them. He stated they should not cause any adverse effects.</p> <p>Upon interview on 6/21/24 at 9:14 a.m. R4 stated his family member (FM)-B brought in the bottle of Roloids after he had been admitted. R4 stated he FM-B her to bring him some antacids because when he had daily heartburn, he would not have wait for staff to give him a medication due to long wait times. R4 stated he did not think he was on any other medications for heartburn. R4 stated the Rolaid had been on his tray table for a few</p>	F 755		

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F 755	<p>Continued From page 8 weeks.</p> <p>Upon interview on 6/21/24 at 10:34 a.m. FM-B stated she brought R4 a bottle of antacids due to his frequent heartburn and it takes the staff so long to answer his call light, she thought it was a good idea that he keeps his own bottle. She stated she brought the medications in about 6/12/24 or 6/13/24.</p> <p>Upon interview on 6/21/24 at 11:38 a.m. the pharmacist stated that R4 having the Roloids at his bedside was a concern because it did interact with R4's Rosuvastatin and the facility was not monitoring the use so the pharmacy could not address how much and how often R4 was taking the medication. R4 stated their pharmacy consultant would be doing a full assessment as soon as possible to assure R4 is taking the correct medication for his heartburn.</p> <p>Upon observation and interview RN-A the nurse manager stated she was unaware that R4 had the Roloids in the room. She stated the bottle should have been removed and the provider and pharmacy should have been notified. She stated she was not aware of any contraindications of the Roloids with any other medications R4 was taking.</p> <p>Upon interview on 6/21/24 at 1:37 p.m. the Administrator stated if a resident is not allowed to self-administer their own medications, then the nurse staff would remove the medication from the room. "If the nurse sees it, we don't rifle through their things."</p> <p>A facility policy titled Medication Administration - General Guidelines dated 3/10/23 indicated</p>	F 755		

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F 755	<p>Continued From page 9</p> <p>Residents are not allowed to self-administer any medication unless specifically authorized to do so by their attending physician/nurse practitioner and then only in accordance with the procedure for Self-Administration of Medications. Medications in a resident's room must be secured or maintained in the medication's cart/room. A physician's order is to be obtained when a resident is involved with self-administration of his/her medications in any way:</p> <p>If a resident keeps his/her medications at the bedside, an order similar to the following is to be obtained: "resident self-administer medications- medications kept at bedside" (this would include prescription medications and over-the-counter medications). The Resident is instructed to report use of PRN.</p>	F 755		