



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered

April 5, 2024

Licensee

Villages Of Lonsdale, LLC
1000 Birch Street Northeast
Lonsdale, MN 55046

RE: Project Number(s) SL26754016

Dear Licensee:

The Minnesota Department of Health (MDH) completed a survey on March 6, 2024, for the purpose of evaluating and assessing compliance with state licensing statutes. At the time of the survey, MDH noted violations of the laws pursuant to Minnesota Statute, Chapter 144G, Minnesota Food Code, Minnesota Rules Chapter 4626, Minnesota Statute 626.5572 and/or Minnesota Statute Chapter 260E.

STATE CORRECTION ORDERS

The enclosed State Form documents the state correction orders. MDH documents state licensing correction orders using federal software. Tag numbers are assigned to Minnesota state statutes for Assisted Living Facilities. The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute number and the corresponding text of the state statute out of compliance are listed in the "Summary Statement of Deficiencies" column. This column also includes the findings that are in violation of the state statute after the statement, "This MN Requirement is not met as evidenced by . . ."

In accordance with Minn. Stat. § 144G.31 Subd. 4, MDH may assess fines based on the level and scope of the violations; **however, no immediate fines are assessed for this survey of your facility.**

DOCUMENTATION OF ACTION TO COMPLY

In accordance with Minn. Stat. § 144G.30, Subd. 5(c), the licensee must document actions taken to comply with the correction orders within the time period outlined on the state form; however, plans of correction are not required to be submitted for approval.

The correction order documentation should include the following:

- Identify how the area(s) of noncompliance was corrected related to the resident(s)/employee(s) identified in the correction order.
- Identify how the area(s) of noncompliance was corrected for all of the provider's resident(s)/employees that may be affected by the noncompliance.
- Identify what changes to your systems and practices were made to ensure compliance with the specific statute(s).

CORRECTION ORDER RECONSIDERATION PROCESS

In accordance with Minn. Stat. § 144G.32, Subd. 2, you may challenge the correction order(s) issued, including the level and scope, and any fine assessed through the correction order reconsideration process. The request for reconsideration must be in writing and received by MDH within 15 calendar days of the correction order receipt date.

To submit a reconsideration request, please visit:

<https://forms.web.health.state.mn.us/form/HRDAppealsForm>

The MDH Health Regulation Division (HRD) values your feedback about your experience during the survey and/or investigation process. Please fill out this anonymous provider feedback questionnaire at your convenience at this link: **<https://forms.office.com/g/Bm5uQEpHV>**. Your input is important to us and will enable MDH to improve its processes and communication with providers. If you have any questions regarding the questionnaire, please contact Susan Winkelmann at susan.winkelmann@state.mn.us or call 651-201-5952.

You are encouraged to retain this document for your records. It is your responsibility to share the information contained in the letter and state form with your organization's Governing Body.

If you have any questions, please contact me.

Sincerely,

A handwritten signature in black ink, appearing to read "Jodi Johnson", with a long horizontal flourish extending to the right.

Jodi Johnson, Supervisor

State Evaluation Team

Email: jodi.johnson@state.mn.us

Telephone: 507-344-2730 Fax: 1-866-890-9290

PMB

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 26754	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____		(X3) DATE SURVEY COMPLETED 03/06/2024
NAME OF PROVIDER OR SUPPLIER VILLAGES OF LONSDALE LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1000 BIRCH STREET NORTHEAST LONSDALE, MN 55046		
(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
0 000	Initial Comments *****ATTENTION***** ASSISTED LIVING PROVIDER LICENSING CORRECTION ORDER(S) In accordance with Minnesota Statutes, section 144G.08 to 144G.95, these correction orders are issued pursuant to a survey. Determination of whether violations are corrected requires compliance with all requirements provided at the Statute number indicated below. When Minnesota Statute contains several items, failure to comply with any of the items will be considered lack of compliance. INITIAL COMMENTS: SL26754016 On March 4, 2024 through March 6, 2024, the Minnesota Department of Health conducted a full survey at the above provider, and the following correction orders are issued. At the time of the survey, there were 44 residents; 19 receiving services under the Assisted Living with Dementia Care license.		0 000	Minnesota Department of Health is documenting the State Correction Orders using federal software. Tag numbers have been assigned to Minnesota State Statutes for Assisted Living License Providers. The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state Statute number and the corresponding text of the state Statute out of compliance is listed in the "Summary Statement of Deficiencies" column. This column also includes the findings which are in violation of the state requirement after the statement, "This Minnesota requirement is not met as evidenced by." Following the surveyors' findings is 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. The letter in the left column is used for tracking purposes and reflects the scope and level issued pursuant to 144G.31 subd. 1, 2, and 3.	
0 620 SS=D	144G.42 Subd. 6 (a) / 626.557, Subd. 3 Compliance with requirements for reporting ma (a) The assisted living facility must comply with		0 620		

Minnesota Department of Health

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

TITLE

(X6) DATE

Minnesota Department of Health

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0 620	Continued From page 1 the requirements for the reporting of maltreatment of vulnerable adults in section 626.557. The facility must establish and implement a written procedure to ensure that all cases of suspected maltreatment are reported. The requirement in Minnesota Statute section 626.557, Subd. 3 is: (a) A mandated reporter who has reason to believe that a vulnerable adult is being or has been maltreated, or who has knowledge that a vulnerable adult has sustained a physical injury which is not reasonably explained shall immediately report the information to the common entry point. If an individual is a vulnerable adult solely because the individual is admitted to a facility, a mandated reporter is not required to report suspected maltreatment of the individual that occurred prior to admission, unless: (1) the individual was admitted to the facility from another facility and the reporter has reason to believe the vulnerable adult was maltreated in the previous facility; or (2) the reporter knows or has reason to believe that the individual is a vulnerable adult as defined in section 626.5572, subdivision 21, paragraph (a), clause (4). (b) A person not required to report under the provisions of this section may voluntarily report as described above. (c) Nothing in this section requires a report of known or suspected maltreatment, if the reporter knows or has reason to know that a report has been made to the common entry point. (d) Nothing in this section shall preclude a reporter from also reporting to a law enforcement agency. (e) A mandated reporter who knows or has	0 620			

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0 620	Continued From page 2 reason to believe that an error under section 626.5572, subdivision 17, paragraph (c), clause (5), occurred must make a report under this subdivision. If the reporter or a facility, at any time believes that an investigation by a lead investigative agency will determine or should determine that the reported error was not neglect according to the criteria under section 626.5572, subdivision 17, paragraph (c), clause (5), the reporter or facility may provide to the common entry point or directly to the lead investigative agency information explaining how the event meets the criteria under section 626.5572, subdivision 17, paragraph (c), clause (5). The lead investigative agency shall consider this information when making an initial disposition of the report under subdivision 9c. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to immediately report to the Minnesota Adult Abuse Reporting Center (MAARC) suspected maltreatment for one of one resident (R4). This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or a limited number of staff are involved or the situation has occurred only occasionally). The findings include: R4's diagnoses included dementia.	0 620			

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0 620	Continued From page 3 R4's Individual Abuse Prevention Plan (IAPP) dated January 15, 2024, indicated the resident did not ambulate as was unable to ambulate safely, and utilized a EZ stand for transfers. The IAPP further indicated R4 was not susceptible to abuse from another individual, including other vulnerable adults. R6's diagnoses included Alzheimer's disease. R6's IAPP dated January 26, 2024, indicated the resident wandered into other apartments and was at risk of abusing other vulnerable adults. R6's progress note dated February 5, 2024, at 1:14 p.m. indicated staff reported during rounds R6 was in another resident's apartment, was undressed and in bed with other resident (R4). Staff had difficulty getting resident out of the bed but were able to finally redirect him. Other resident family has a camera in the apartment and notified ED (executive director) and nursing they observed event on the camera. Per family, resident used other resident's bathroom, came out with his pants off then undressed and crawled over the legs of the other resident and got into bed with her. R6's progress note dated February 12, 2024, at 10:14 a.m. indicated staff reported during shift change at approximately 6:00 a.m. another resident (R4) pushed her pendant and when staff entered apartment, resident was in bed "spooning and hugging resident". Resident was on the inside of the bed and appears resident crawled over other resident. The MAARC report submitted February 13, 2024, at 15:16 (3:16 p.m.), indicated on February 5, 2024, at approximately 9:30 p.m., R6 was	0 620			

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0 620	Continued From page 4 observed in bed with R4; staff had a difficult time getting R6 out of R4's bed. On February 12, 2024, at approximately 6:00 a.m., R4 pushed her pendant and when staff entered R4's apartment, R6 was observed hugging R4 in bed. R6 was on the inside of the bed and appeared had crawled over R4. R6's record did not include evidence a MAARC report had been submitted for the February 5, 2024, incident until February 13, 2024. On March 6, 2024, at 10:38 a.m. assisted living quality director (ALQD)-F reviewed the above MAARC report and stated the incident on February 5, 2024, should have been reported within 24 hours to MAARC. The licensee's Maltreatment Prohibition policy dated August 1, 2021, indicated: G. Investigation of the Alleged Incident a. Any reported allegations of abuse towards a vulnerable individual will be immediately assessed by the nurse in charge. b. Once the allegations/incident are initially assessed by the nurse in charge and interventions are initiated to prevent further occurrences, the alleged incident needs to be reported to the MAARC within 24 hours. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	0 620			
0 660 SS=F	144G.42 Subd. 9 Tuberculosis prevention and control (a) The facility must establish and maintain a	0 660			

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0 660	Continued From page 5 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 the CDC's Morbidity and Mortality Weekly Report. The program must include a tuberculosis infection control plan that covers all paid and unpaid employees, contractors, students, and regularly scheduled volunteers. The commissioner shall provide technical assistance regarding implementation of the guidelines. (b) The facility must maintain written evidence of compliance with this subdivision. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to maintain a tuberculosis (TB) prevention program, based on the most current guidelines issued by the Centers for Disease Control and Prevention (CDC) which included a completed facility TB risk assessment, and documentation of a completed health history and symptom screening for one of two employees (unlicensed personnel (ULP)-E). This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety) and was issued at a widespread scope (when problems are pervasive or represent a systemic failure that has affected or has the potential to affect a large portion or all of the residents). The findings include: FACILITY TB RISK ASSESSMENT	0 660			

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0 660	Continued From page 6 During the entrance conference on March 4, 2024, at 11:22 a.m. with licensed assisted living director (LALD)-A, clinical nurse supervisor (CNS)-B, and assisted living quality director (ALQD)-F, a request was made to review the facility TB risk assessment. On March 6, 2024, at 2:56 p.m. a TB risk assessment, undated, was provided and was incomplete. ALQD-F stated she scanned the odd pages of the current TB risk assessment and placed assessment in the shred box. ALQD-F stated she could not verify the date on the scanned TB risk assessment, and was unable to retrieve the assessment out of the shred box. HEALTH HISTORY AND SYMPTOM SCREENING ULP-E was hired April 19, 2010, to provide direct care and services to the facility's residents. ULP-E's employee record lacked evidence of TB history and symptom screening. On March 5, 2024, at 2:40 p.m. ALQD-F stated ULP-E's record lacked a TB history and symptom screening. The Minnesota Department of Health guidelines Regulations for Tuberculosis Control in Minnesota Health Care Settings dated July 2013, and based on CDC guidelines, indicated a TB infection control program should include an annual facility TB risk assessment. The guidelines also indicated an employee may begin working with patients (residents) after a negative TB history and symptom screen (no symptoms of active TB disease) and a negative IGRA (serum blood test) or TST (first step) dated within 90 days before hire. The second TST may be performed after the	0 660			

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0 660	Continued From page 7 HCW starts working with patients. Baseline TB screening should be documented in the employee's record. The licensee's Tuberculosis Prevention and Control policy, revised August 10, 2022, indicated a TB facility risk assessment will be completed at least bi-annually. This policy lacked the most current guidelines issued by the CDC. In addition, this policy indicated a provider's order will be obtained and kept on file as a standard protocol to administer a TB screening and testing of all volunteers and employees upon hire. No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 660			
0 680 SS=F	144G.42 Subd. 10 Disaster planning and emergency preparedness (a) The facility must meet the following requirements: (1) have a written emergency disaster plan that contains a plan for evacuation, addresses elements of sheltering in place, identifies temporary relocation sites, and details staff assignments in the event of a disaster or an emergency; (2) post an emergency disaster plan prominently; (3) provide building emergency exit diagrams to all residents; (4) post emergency exit diagrams on each floor; and (5) have a written policy and procedure regarding missing residents. (b) The facility must provide emergency and disaster training to all staff during the initial staff	0 680			

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0 680	Continued From page 8 orientation and annually thereafter and must make emergency and disaster training annually available to all residents. Staff who have not received emergency and disaster training are allowed to work only when trained staff are also working on site. (c) The facility must meet any additional requirements adopted in rule. This MN Requirement is not met as evidenced by: Based on interview and record review the licensee failed to have a written emergency preparedness plan (EPP) with all the required content. This had the potential to affect all residents receiving services under the assisted living license. This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at a widespread scope (when problems are pervasive or represent a systemic failure that has affected or has potential to affect a large portion or all of the residents). The findings include: The licensee's emergency preparedness plan lacked the following: - EPP training program for staff (including documentation of training provided); and - EPP testing/annual testing requirements. On March 4, 2024, at 2:40 p.m. assisted living quality director (ALQD)-F stated the above listed contents were not found in the EPP binder and emailed the maintenance director requesting	0 680			

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0 680	Continued From page 9 documentation of training dates. Nothing further was provided. The licensee's Emergency Preparedness Program policy dated March 16, 2023, indicated training on the facility Emergency Preparedness Plan will be upon hire, as well as conducted on an annual basis for all employees' volunteers, and available contract staff. Staff training on fire safety and evacuation plans will be conducted at least twice per year. No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 680			
0 950 SS=C	144G.50 Subd. 3 Designation of representative (a) Before or at the time of execution of an assisted living contract, an assisted living facility must offer the resident the opportunity to identify a designated representative in writing in the contract and must provide the following verbatim notice on a document separate from the contract: "RIGHT TO DESIGNATE A REPRESENTATIVE FOR CERTAIN PURPOSES. You have the right to name anyone as your "Designated Representative." A Designated Representative can assist you, receive certain information and notices about you, including some information related to your health care, and advocate on your behalf. A Designated Representative does not take the place of your guardian, conservator, power of attorney ("attorney-in-fact"), or health care power of attorney ("health care agent"), if applicable."	0 950			

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0 950	Continued From page 10 (b) The contract must contain a page or space for the name and contact information of the designated representative and a box the resident must initial if the resident declines to name a designated representative. Notwithstanding subdivision 1, paragraph (f), the resident has the right at any time to add, remove, or change the name and contact information of the designated representative. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to ensure the licensee provided the required notice for right to a designated representative with the required verbiage on a document separate from the contract for one of one resident (R1). This practice resulted in a level one violation (a violation that has no potential to cause more than a minimal impact on the resident and does not affect health or safety), and was issued at a widespread scope (when problems are pervasive or represent a systemic failure that has affected or has potential to affect a large portion or all of the residents). The findings include: R1's Assisted Living Contract was signed by her daughter on December 15, 2023. R1's records lacked evidence in writing of providing on a document separate from the contact verbatim notice of "RIGHT TO DESIGNATE A REPRESENTATIVE FOR CERTAIN PURPOSES. You have the right to name anyone as your "Designated	0 950			

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0 950	Continued From page 11 Representative." A Designated Representative can assist you, receive certain information and notices about you, including some information related to your health care, and advocate on your behalf. A Designated Representative does not take the place of your guardian, conservator, power of attorney ("attorney-in-fact"), or health care power of attorney ("health care agent"), if applicable." On March 6, 2024, at 10:57 a.m. assisted living quality director (ALQD)-F stated the contract needed to be updated with the verbatim notice and this would be for all current residents. No additional information was provided. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 950			
01060 SS=D	144G.52 Subd. 9 Emergency relocation (a) A facility may remove a resident from the facility in an emergency if necessary due to a resident's urgent medical needs or an imminent risk the resident poses to the health or safety of another facility resident or facility staff member. An emergency relocation is not a termination. (b) In the event of an emergency relocation, the facility must provide a written notice that contains, at a minimum: (1) the reason for the relocation; (2) the name and contact information for the location to which the resident has been relocated and any new service provider; (3) contact information for the Office of Ombudsman for Long-Term Care and the Office of Ombudsman for Mental Health and Developmental Disabilities;	01060			

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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
01060	Continued From page 12 (4) if known and applicable, the approximate date or range of dates within which the resident is expected to return to the facility, or a statement that a return date is not currently known; and (5) a statement that, if the facility refuses to provide housing or services after a relocation, the resident has the right to appeal under section 144G.54. The facility must provide contact information for the agency to which the resident may submit an appeal. (c) The notice required under paragraph (b) must be delivered as soon as practicable to: (1) the resident, legal representative, and designated representative; (2) for residents who receive home and community-based waiver services under chapter 256S and section 256B.49, the resident's case manager; and (3) the Office of Ombudsman for Long-Term Care if the resident has been relocated and has not returned to the facility within four days. (d) Following an emergency relocation, a facility's refusal to provide housing or services constitutes a termination and triggers the termination process in this section.currently known; and This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to provide a written notice with the required content for an emergency relocation for one of one resident (R2). This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or	01060			

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01060	Continued From page 13 a limited number of staff are involved or the situation has occurred only occasionally). The findings include: R2 began receiving assisted living services from the licensee on June 15, 2022. R2's service plan signed March 1, 2024, indicated R2 received services which included dressing, housekeeping, laundry, medication administration, blood glucose monitoring, weight monitoring, and assistance with bilateral leg pumps. R2's progress notes revealed the resident had been hospitalized on December 17, 2023, following a fall at the facility. The progress notes further revealed R2 returned to the assisted living facility on December 19, 2023. R2's medical record did not include evidence an emergency relocation notice had been provided for the above hospitalization. On March 6, 2024, at 1:57 p.m. assisted living quality director (ALQD)-F stated being unable to find evidence an emergency location notice had been provided to R2. No further information was provided. TIME PERIOD TO CORRECT: Twenty-one (21) days	01060			
01640 SS=D	144G.70 Subd. 4 (a-e) Service plan, implementation and revisions to (a) No later than 14 calendar days after the date	01640			

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01640	Continued From page 14 that services are first provided, an assisted living facility shall finalize a current written service plan. (b) The service plan and any revisions must include a signature or other authentication by the facility and by the resident documenting agreement on the services to be provided. The service plan must be revised, if needed, based on resident reassessment under subdivision 2. The facility must provide information to the resident about changes to the facility's fee for services and how to contact the Office of Ombudsman for Long-Term Care and the Office of Ombudsman for Mental Health and Developmental Disabilities. (c) The facility must implement and provide all services required by the current service plan. (d) The service plan and the revised service plan must be entered into the resident record, including notice of a change in a resident's fees when applicable. (e) Staff providing services must be informed of the current written service plan. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure a written service plan was revised to reflect the current services provided for one of two residents (R2). This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or a limited number of staff are involved or the situation has occurred only occasionally). The findings include:	01640			

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01640	Continued From page 15 R2 R2's diagnoses included cerebral infarction (stroke), diabetes, and edema (swelling due to excess fluid accumulation in the body tissues). R2's Service Plan dated March 1, 2024, included direction for staff to apply lotion to both feet and legs up to knees, then apply compression stockings and Velcro wraps to both legs every evening. R2's Service Checkoff List dated March 2024, included the above treatment; however, it was not being signed off by staff. On March 4, 2024, at 2:26 a.m. R2 was observed lying in bed with bilateral leg pumps on his legs. The pumps were plugged in and running. R2 stated he wore the leg pumps once a day for an hour and staff assisted with putting them on and taking them off. When asked if R2 wore any other kind of leg wraps he stated, "not anymore." On March 6, 2024, at 10:38 a.m. clinical nurse supervisor (CNS)-B stated R2 no longer wore the compression stockings or Velcro wraps because they were discontinued in December 2023, upon return from hospitalization. CNS-B stated R2's Service Plan should have been revised to reflect this. The licensee's Content, Development and Revision of the Service Plan policy dated August 1, 2021, indicated: J. Revision to the Service Plan: a. Each resident's Service Plan is reviewed by the Registered Nurse, as follows: i. During each regular resident monitoring visit, which occurs at least every 90	01640			

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01640	Continued From page 16 days ii. Whenever changes are needed to the services to be provided because of a change in the resident's condition, after receipt of new or revised orders for the resident's health provider if these changes will require a change in the services the staff provide, following an incident if new services ar identified and accepted, or following the resident's return from a hospital or nursing home if new services are identified and accepted. No further information was provided. TIME PERIOD TO CORRECT- Twenty-one (21) days	01640			
01750 SS=D	144G.71 Subd. 7 Delegation of medication administration When administration of medications is delegated to unlicensed personnel, the assisted living facility must ensure that the registered nurse has: (1) instructed the unlicensed personnel in the proper methods to administer the medications, and the unlicensed personnel has demonstrated the ability to competently follow the procedures; (2) specified, in writing, specific instructions for each resident and documented those instructions in the resident's records; and (3) communicated with the unlicensed personnel about the individual needs of the resident. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure nursing staff administered prescribed insulin per manufacturer's instruction for one of two	01750			

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01750	Continued From page 17 residents (R3). This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or a limited number of staff are involved or the situation has occurred only occasionally). The findings include: R1's diagnoses included type 2 diabetes. R1's Service Plan dated April 17, 2023, indicated insulin administration was assigned to unlicensed personnel (ULP). R1's physician orders dated February 13, 2024, included: Basaglar Kwikpen 100 unit/milliliter - inject 44 units subcutaneously (under the skin) every morning and Humalog Kwikpen 100 unit/milliliter - inject 4 units subcutaneously every morning. On March 5, 2024, at 9:42 a.m. ULP-E was observed administering insulin to R1. ULP-E prepped the Basaglar Kwikpen with a needle, prepped upper right arm with an alcohol pad. ULP-E primed the Kwikpen by dialing pen to two units, wasted, dialed pen to 44 units and injected Basaglar insulin 44 units into mid-right deltoid (muscle). ULP-E prepped the Humalog Kwikpen with a needle, prepped upper left arm with an alcohol pad. ULP-E primed the Kwikpen by dialing pen to two units, wasted, dialed pen to 4 units and injected Humalog insulin 4 units into mid-left deltoid.	01750			

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01750	Continued From page 18 ULP-E's employee file contained a subcutaneous injection competency dated April 5, 2022, signed by previous facility clinical nurse supervisor (CNS). The competency directed subcutaneous injections are to be administered in upper arm, abdomen, or thigh (fatty area). On March 5, 2024, at 9:52 a.m. ULP-E stated she was trained by the previous CNS, to give the insulin in the deltoid. On March 5, 2024, at 9:56 a.m. CNS-B stated subcutaneous insulin given in arm should be in the subcutaneous area (fatty area), not the deltoid (intramuscularly). CNS-B stated she would complete follow-up training with staff on insulin administration. The manufacturer's instructions for Basaglar Kwikpen - insulin glargine injection solution dated November 2022, indicated Basaglar is injected under the skin (subcutaneously) of the stomach area, buttocks, upper legs or upper arms. The manufacturer's instructions for Humalog Kwikpen - injection solution dated March 31, 2020, indicated Humalog is injected under the skin (subcutaneously) of your stomach area, buttocks, upper legs or upper arms. The licensee's Administration of Medications, Treatment and Therapy by Unlicensed Personnel policy, reviewed/amended, February 15, 2022, indicated medications, treatment and therapy always need to be administered according to the "6 Rights": 1. Right person 2. Right medication, treatment or therapy 3. Right time	01750			

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01750	Continued From page 19 4. Right route (by mouth, eye drops, to the skin, etc.) 5. Right dose (how many milligrams, drops, etc.) 6. Right chart/record to document that the medication, treatment and therapy was taken. No further information provided. TIME PERIOD FOR CORRECTION: Seven (7) days	01750			
01890 SS=D	144G.71 Subd. 20 Prescription drugs A prescription drug, prior to being set up for immediate or later administration, must be kept in the original container in which it was dispensed by the pharmacy bearing the original prescription label with legible information including the expiration or beyond-use date of a time-dated drug. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure medications were maintained bearing the original prescription label for one of four residents (R2) observed during medication administration. This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or a limited number of staff are involved or the situation has occurred only occasionally).	01890			

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01890	Continued From page 20 The findings include: On March 5, 2024, at 8:01 a.m. unlicensed personnel (ULP)-D was observed administering insulin to R2. ULP-D brought a plastic box into R2's apartment containing an opened insulin glargine pen 100 unit/milliliter (ml). The insulin pen lacked an original prescription label with information regarding the directions for use, medication name, medication dosage, resident's name, and the pharmacy in which it had been issued. ULP-D stated R2's insulin pens come in a bag from the pharmacy; the prescription label is on the outside of the bag. ULP-D further stated the original label is not kept with the insulin; it is given to the nurse so she can re-order the medication. On March 5, 2024, at 1:38 p.m. clinical nurse supervisor (CNS)-B stated R2's medications were supplied by the VA (Veteran's Administration) and the insulin pens were not individually labeled though did have a prescription label on the outside of the package they came in. CNS-B further stated not having a label available for the ULP to view for the insulin pen when setting up/administering the medication, and should have. The licensee's Storage of Medications policy dated January 6, 2021, indicated: B. Proper Storage of Medications b. Until the medication is set up for immediate or later administration by a nurse, a legend drug must be kept in its original container bearing the original prescription label with legible information stating the prescription number, name of drug, strength and quantity of drug, expiration date of time-dated drug, directions for use,	01890			

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01890	Continued From page 21 resident's name, prescriber's name, date of issue and the name and address of the licensed pharmacy that issued the medications. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days.	01890			
01950 SS=D	144G.72 Subd. 4 Administration of treatments and therapy Ordered or prescribed treatments or therapies must be administered by a nurse, physician, or other licensed health professional authorized to perform the treatment or therapy, or may be delegated or assigned to unlicensed personnel by the licensed health professional according to the appropriate practice standards for delegation or assignment. When administration of a treatment or therapy is delegated or assigned to unlicensed personnel, the facility must ensure that the registered nurse or authorized licensed health professional has: (1) instructed the unlicensed personnel in the proper methods with respect to each resident and the unlicensed personnel has demonstrated the ability to competently follow the procedures; (2) specified, in writing, specific instructions for each resident and documented those instructions in the resident's record; and This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to ensure unlicensed personnel (ULP) notified the registered nurse (RN) of blood glucose exceeding parameters for one of one resident (R2).	01950			

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01950	Continued From page 22 This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death), and was issued at an isolated scope (when one or a limited number of residents are affected or one or a limited number of staff are involved or the situation has occurred only occasionally). The findings include: R2's diagnoses included diabetes. R2's Service Plan dated March 1, 2024, included blood sugar monitoring. The service plan directed staff to check resident blood sugar in the morning and record. Hold insulin if blood sugar less than 100 or greater then 500. If below 100, give orange juice, cookies, peanut butter sandwich and recheck in 30 minutes to one hour, then notify nursing. If greater than 500, have resident drink lots of water and propel self in wheelchair, recheck in 30 minutes to one hour and then notify nursing. Staff will alert nurse if blood sugar is not in desired range set by physician. A review of R2's blood sugar values revealed a blood sugar of 527 on March 3, 2024. R2's progress notes did not include evidence the nurse had been contacted related to R2's elevated blood sugar. On March 6, 2024, at 12:02 p.m. assisted living quality director (ALQD)-F stated R2's elevated blood sugar on March 3, 2024, had not been reported to the nurse and should have been.	01950			

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01950	Continued From page 23 No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	01950			
02320 SS=E	144G.91 Subd. 4 (b) Appropriate care and services (b) Residents have the right to receive health care and other assisted living services with continuity from people who are properly trained and competent to perform their duties and in sufficient numbers to adequately provide the services agreed to in the assisted living contract and the service plan. This MN Requirement is not met as evidenced by: The licensee failed to ensure medications were administered according to policy and accepted standards of practice for two of four residents (R2, R5) observed during medication administration. This practice resulted in a level two violation (a violation that did not harm a resident's health or safety but had the potential to have harmed a resident's health or safety, but was not likely to cause serious injury, impairment, or death) and was issued at a pattern scope (when more than a limited number of residents are affected, more than a limited number of staff are involved, or the situation has occurred repeatedly; but is not found to be pervasive). The findings include: R2 R2's diagnoses included diabetes.	02320			

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02320	Continued From page 24 R2's Service Plan dated March 1, 2024, included medication administration. On March 5, 2024, at 8:01 a.m. unlicensed personnel (ULP)-D was observed entering R2's apartment to administer medications. ULP-D brought a plastic box into R2's apartment containing supplies to test R2's blood glucose level and an opened insulin glargine pen 100 unit/milliliter (ml). ULP-D also brought a medication cup containing R2's oral medications that had previously been set-up. ULP-D proceeded to measure R2's blood sugar, administered his oral medications, then prepared R2's insulin pen for administration and dialed the pen to 15 units. The insulin pen lacked an original prescription label with information regarding the directions for use, medication name, medication dosage, resident's name, and the pharmacy in which it had been issued. The surveyor asked ULP-D how she knew what dosage to administer without a prescription label or physician order in front of her. ULP-D stated when she set up R2's oral medications she had viewed his insulin order and "remembered" it. On March 5, 2024, at 1:38 p.m. clinical nurse supervisor (CNS)-B stated the medication cart for the assisted living residents was currently kept in the locked memory care unit. ULP were to set up the medications from the cart in the memory care unit then were expected to bring their tablet with them to compare with the electronic medication administration record (eMAR). CNS-B stated being aware R2's insulin pen did not have a prescription label and would have expected ULP-D to review the physician order on the tablet prior to dialing up the dose and administering to R2.	02320			

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NAME OF PROVIDER OR SUPPLIER VILLAGES OF LONSDALE LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1000 BIRCH STREET NORTHEAST LONSDALE, MN 55046		
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02320	Continued From page 25 R5 R5's diagnoses included spinal stenosis. R5's Service Plan dated January 17, 2024, included medication administration. On March 5, 2024, at 10:04 a.m. ULP-D was observed setting up and administering R5's medications. The medications included hydrocortisone lotion 1%, apply to back area twice daily, and diclofenac sodium topical gel 1%, apply two grams topically to neck pain every morning. ULP-D donned gloves then applied the hydrocortisone lotion to R5's back. When finished, ULP-D stated there was still some lotion on her gloves and asked R5 if she wanted her to rub the remainder of the medication onto R5's hands. R5 consented and ULP-D then rubbed what was left of the hydrocortisone lotion onto R5's hands. ULP-D then doffed her gloves and donned new gloves. She then applied R5's diclofenac sodium 1% topical gel onto the palm of her hand without measuring. ULP-D applied the gel to R5's neck then asked the resident if she wanted her to also apply the gel to R5's lower back. R5 consented and ULP-D rubbed the remainder of the gel onto R5's lower back. At 10:27 a.m. following the observation, the surveyor asked ULP-D how she measured R5's diclofenac topical gel to ensure she was getting the prescribed dosage. ULP-D indicted the approximate size of a nickel to the surveyor. When asked if she'd been trained to use a measurement device ULP-D indicated "no." The surveyor and ULP-D then inspected the bag R5's diclofenac gel was stored in and observed a plastic measuring device for the gel. ULP-D stated she had not been shown how to use the measuring device.	02320			

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 26754	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____		(X3) DATE SURVEY COMPLETED 03/06/2024
NAME OF PROVIDER OR SUPPLIER VILLAGES OF LONSDALE LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1000 BIRCH STREET NORTHEAST LONSDALE, MN 55046		
(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
02320	Continued From page 26 On March 5, 2024, at 1:38 p.m. CNS-B stated topical medications should only be administered to the area of the body indicated in the physician order. CNS-B further stated expecting ULP to utilize the measuring device that comes with the topical medication when a specific dose was ordered. The licensee's Administration of Medications, Treatment and Therapy by Unlicensed Personnel policy dated February 15, 2022, indicated: 2. Medications, treatment and therapy always need to be administered according to the "6 Rights" a. Right person b. Right medication, treatment or therapy c. Right time d. Right route e. Right dose f. Right chart/record to document that the medication, treatment and therapy was taken. The licensee's Medication Errors policy dated August 1, 2021, indicated: staff will report any medication errors promptly to the nurse in charge. The nurse will immediately investigate any medication error and will implement and document corrective actions to reduce the risk of similar medication error in the future. No further information provided. TIME PERIOD FOR CORRECTION: Seven (7) days	02320			

Type: Full
Date: 03/04/24
Time: 13:50:00
Report: 1033241028

Food and Beverage Establishment Inspection Report

Page 1

Location:

The Villages of Lonsdale
1000 Birch Street NE
Lonsdale, MN55046
Rice County, 66

Establishment Info:

ID #: 0023814
Risk: Medium
Announced Inspection: No

License Categories:

Expires on: / /

Operator:

Villages of Lonsdale, LLC

Phone #: 5077443453
ID #: 52222

The violations listed in this report include any previously issued orders and deficiencies identified during this inspection. Compliance dates are shown for each item.

No NEW orders were issued during this inspection.

Surface and Equipment Sanitizers

Hot Water: = at 163 Degrees Fahrenheit
Location: Dish Machine
Violation Issued: No

Food and Equipment Temperatures

Process/Item: Cold Holding
Temperature: 38 Degrees Fahrenheit - Location: Egg Salad-Two Door Cooler
Violation Issued: No

Process/Item: Cold Holding
Temperature: 38 Degrees Fahrenheit - Location: Sour Cream-Two Door Cooler
Violation Issued: No

Process/Item: Cold Holding
Temperature: 35 Degrees Fahrenheit - Location: Single Door Cooler
Violation Issued: No

Process/Item: Cold Holding
Temperature: 35 Degrees Fahrenheit - Location: Ground Beef-Walk In Cooler
Violation Issued: No

Process/Item: Cold Holding
Temperature: 0> Degrees Fahrenheit - Location: Walk In Freezer
Violation Issued: No

Process/Item: Cold Holding
Temperature: 38 Degrees Fahrenheit - Location: Event Room Cooler
Violation Issued: No

Type: Full
Date: 03/04/24
Time: 13:50:00
Report: 1033241028
The Villages of Lonsdale

Food and Beverage Establishment
Inspection Report

Total Orders	In This Report	Priority 1	Priority 2	Priority 3
		0	0	0

NOTE: Plans and specifications must be submitted for review and approval prior to new construction, remodeling or alterations.

I acknowledge receipt of the inspection report number 1033241028 of 03/04/24.

Certified Food Protection ManagerStephanie Ann Moe

Certification Number: FM107666 Expires: 08/09/24

Inspection report reviewed with person in charge and emailed.

Signed:Stephanie Ann Moe

Signed:Isaiah Armendariz
Environmental Health Specialist
Mankato District Office
507-344-2743
isaiah.armendariz@state.mn.us

Report #: 1033241028

m1

DEPARTMENT OF HEALTH

Address

1000 Birch Street NE

City/State

Lonsdale, MN

Zip Code

55046

Telephone

5077443453

License/Permit #

0023814

Permit Holder

Villages of Lonsdale, LLC

Purpose of Inspection

Full

Est Type

Risk Category

M

No. of RF/PHI Categories Out

0

No. of Repeat RF/PHI Categories Out

0

Legal Authority MN Rules Chapter 4626

Date

03/04/24

Time In

13:50:00

Time Out

FOODBORNE ILLNESS RISK FACTORS AND PUBLIC HEALTH INTERVENTIONS

Circle designated compliance status (IN, OUT, N/O, N/A) for each numbered item

Mark "X" in appropriate box for COS and/or R

IN=in compliance

OUT= not in compliance

N/O= not observed

N/A= not applicable

COS=corrected on-site during inspection

R= repeat violation

Compliance Status

COS

R

Surpervision

1

IN

OUTPIC knowledgeable; duties & oversight

2

IN

OUTN/ACertified food protection manager, duties

Employee Health

3

IN

OUTMgmt/Staff;knowledge,responsibilities&reporting

4

IN

OUTProper use of reporting, restriction & exclusion

5

IN

OUTProcedures for responding to vomiting & diarrheal events

Good Hygenic Practices

6

IN

OUT

N/O

Proper eating, tasting, drinking, or tobacco use

7

IN

OUT

N/O

No discharge from eyes, nose, & mouth

Preventing Contamination by Hands

8

IN

OUTN/OHands clean & properly washed

9

IN

OUTN/AN/ONo bare hand contact with RTE foods or pre-approved alternate pprocedure properly followed

10

IN

OUTAdequate handwashing sinks supplied/accessible

Approved Source

11

IN

OUTFood obtained from approved source

12

IN

OUTN/AN/OFood received at proper temperature

13

IN

OUTFood in good condition, safe, & unadulterated

14

IN

OUT

N/A

N/OREquired records available; shellstock tags, parasite destruction

Protection from Contamination

15

IN

OUTN/AN/OFood separated and protected

16

IN

OUTN/AFood contact surfaces: cleaned & sanitized

17

IN

OUTProper disposition of returned, previously served, reconditioned, & unsafe food

Compliance Status

COS

R

Time/Temperature Control for Safety

18

IN

OUT

N/A

N/O

Proper cooking time & temperature

19

IN

OUT

N/A

N/O

Proper reheating procedures for hot holding

20

IN

OUT

N/A

N/O

Proper cooling time & temperature

21

IN

OUT

N/A

N/O

Proper hot holding temperatures

22

IN

OUT

N/A

Proper cold holding temperatures

23

IN

OUT

N/A

N/O

Proper date marking & disposition

24

IN

OUT

N/A

N/O

Time as a public health control: procedures & records

Consumer Advisory

25

IN

OUT

N/A

Consumer advisory provided for raw/undercooked food

Highly Susceptible Populations

26

IN

OUT

N/A

Pasteurized foods used; prohibited foods not offered

Food and Color Additives and Toxic Substances

27

IN

OUT

N/A

Food additives: approved & properly used

28

IN

OUT

Toxic substances properly identified, stored, & used

Conformance with Approved Procedures

29

IN

OUT

N/A

Compliance with variance/specialized process/HACCP

Risk factors (RF) are improper practices or proceeedures identified as the most prevalent contributing factors of foodborne illness or injury. Public Health Interventions (PHI) are control measures to prevent foodborne illness or injury.

GOOD RETAIL PRACTICES

Good Retail Practices are preventative measures to control the addition of pathogens, chemicals, and physical objects into foods.

Mark "X" in box if numbered item is not in compliance

Mark "X" in appropriate box for COS and/or R

COS=corrected on-site during inspection

R= repeat violation

COS

R

Safe Food and Water

30

IN

OUT

N/A

Pasteurized eggs used where required

31

IN

OUT

N/A

Water & ice obtained from an approved source

32

IN

OUT

N/A

Variance obtained for specialized processing methods

Food Temperature Control

33

IN

OUT

N/A

Proper cooling methods used; adequate equipment for temperature control

34

IN

OUT

N/A

N/O

Plant food properly cooked for hot holding

35

IN

OUT

N/A

N/O

Approved thawing methods used

36

IN

OUT

N/A

Thermometers provided & accurate

Food Identification

37

IN

OUT

N/A

Food properly labled; original container

Prevention of Food Contamination

38

IN

OUT

N/A

Insects, rodents, & animals not present

39

IN

OUT

N/A

Contamination prevented during food prep, storage & display

40

IN

OUT

N/A

Personal cleanliness

41

IN

OUT

N/A

Wiping cloths: properly used & stored

42

IN

OUT

N/A

Washing fruits & vegetables

Compliance Status

COS

R

Proper Use of Utensils

43

IN

OUT

N/A

In-use utensils: properly stored

44

IN

OUT

N/A

Utensils, equipment & linens: properly stored, dried, & handled

45

IN

OUT

N/A

Single-use/single service articles: properly stored & used

46

IN

OUT

N/A

Gloves used properly

Utensil Equipment and Vending

47

IN

OUT

N/A

Food & non-food contact surfaces cleanable, properly designed, constructed, & used

48

IN

OUT

N/A

Warewashing facilities: installed, maintained, & used; test strips

49

IN

OUT

N/A

Non-food contact surfaces clean

Physical Facilities

50

IN

OUT

N/A

Hot & cold water available; adequate pressure

51

IN

OUT

N/A

Plumbing installed; proper backflow devices

52

IN

OUT

N/A

Sewage & waste water properly disposed

53

IN

OUT

N/A

Toilet facilities: properly constructed, supplied, & cleaned

54

IN

OUT

N/A

Garbage & refuse properly disposed; facilities maintained

55

IN

OUT

N/A

Physical facilities installed, maintained, & clean

56

IN

OUT

N/A

Adequate ventilation & lighting; designated areas used

57

IN

OUT

N/A

Compliance with MCIAA

58

IN

OUT

N/A

Compliance with licensing & plan review

Food Recalls:

Person in Charge (Signature)

Date: 03/06/24

Inspector (Signature)