



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered

May 12, 2023

Licensee
Cedar Court
810 West Main Street
Adams, MN 55909

RE: Project Number(s) SL30231015

Dear Licensee:

The Minnesota Department of Health (MDH) completed a survey on April 26, 2023, for the purpose of evaluating and assessing compliance with state licensing statutes. At the time of the survey, the 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.

The MDH concludes the licensee is in substantial compliance. State law requires the facility must take action to correct the state correction orders and document the actions taken to comply in the facility's records. The Department reserves the right to return to the facility at any time should the Department receive a complaint or deem it necessary to ensure the health, safety, and welfare of residents in your care.

STATE CORRECTION ORDERS

The enclosed State Form documents the state correction orders. The 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 and enforcement actions 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 the MDH within 15 calendar days of the correction order receipt date.

A state correction order under Minn. Stat. § 144G.91, Subd. 8, Free from Maltreatment is associated with a maltreatment determination by the Office of Health Facility Complaints. If maltreatment is substantiated, you will receive a separate letter with the reconsideration process under Minn. Stat. § 626.557.

Please email reconsideration requests to: **Health.HRD.Appeals@state.mn.us**. Please attach this letter as part of your reconsideration request. Please clearly indicate which tag(s) you are contesting and submit information supporting your position(s).

Please address your cover letter for reconsideration requests to:

Reconsideration Unit
Health Regulation Division
Minnesota Department of Health
P.O. Box 64970
85 East Seventh Place
St. Paul, MN 55164-0970

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,



Jodi Johnson, Supervisor
State Rapid Response Team
Email: jodi.johnson@state.mn.us
Telephone: 507-344-2730 Fax: 651-215-6894

HHH

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 30231	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 04/26/2023
NAME OF PROVIDER OR SUPPLIER CEDAR COURT		STREET ADDRESS, CITY, STATE, ZIP CODE 810 WEST MAIN STREET ADAMS, MN 55909		
(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: SL#30231015 On April 24, 2023, through April 26, 2023, the Minnesota Department of Health conducted a survey at the above provider, and the following correction orders are issued. At the time of the survey, there were 40 active residents; 20 receiving services under the Assisted Living with Dementia Care license.	0 000	Minnesota Department of Health is documenting the State Licensing 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.	
0 480 SS=F	144G.41 Subd 1 (13) (i) (B) Minimum requirements (13) offer to provide or make available at least the	0 480		

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

TITLE

(X6) DATE

Minnesota Department of Health

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0 480	Continued From page 1 following services to residents: (B) food must be prepared and served according to the Minnesota Food Code, Minnesota Rules, chapter 4626; and This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure food was prepared and served according to the Minnesota Food Code. 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 the residents). The findings include: Please refer to the included document titled, Food and Beverage Establishment Inspection Report dated April 26, 2023, for the specific Minnesota Food Code deficiencies. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 480		
0 510 SS=D	144G.41 Subd. 3 Infection control program (a) All assisted living facilities must establish and maintain an infection control program that complies with accepted health care, medical, and nursing standards for infection control. (b) The facility's infection control program must be consistent with current guidelines from the national Centers for Disease Control and	0 510		

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0 510	Continued From page 2 Prevention (CDC) for infection prevention and control in long-term care facilities and, as applicable, for infection prevention and control in assisted living facilities. (c) The facility must maintain written evidence of compliance with this subdivision. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure infection control standards were followed for one of one unlicensed personnel (ULP-E) during personal cares for 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) 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: On April 25, 2023, at 7:00 a.m. ULP-E was observed providing personal cares and toileting for R3. ULP-E was observed to transfer R3 with use of an EZ stand (mechanical lift machine) to the bathroom, and place R3 on the toilet. Without donning gloves, ULP-E removed R3's pants and placed them in a plastic bag sitting on the bathroom floor, pulled off the wet incontinence brief and placed it in the trash. ULP-E grabbed a clean incontinence brief, and put on to R3's knee level, put on clean pants, touched multiple surfaces while providing cares without washing hands or using gloves. ULP-E donned gloves while administering topical creams and removed	0 510		

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0 510	Continued From page 3 after; however, she failed to wash her hands after glove use and before moving on to next tasks. On April 25, 2023, at 11:30 a.m. ULP-E was observed assisting R3 with toileting. ULP-E donned gloves to assist R3 with cleansing after a bowel movement. ULP-E then removed her gloves but failed to wash or use hand sanitizer before using the EZ stand to transfer R3 back to his wheelchair. After ULP-E transferred R3 and moved him to the hallway, she returned to the bathroom and washed her hands. On April 26, 2023, at 9:40 a.m. clinical nurse supervisor (CNS)-B stated, "we all know the requirement for proper glove use and hand hygiene" and indicated retraining would need to be completed for staff. The licensee's Procedure for Using Glove policy dated November 14, 2022, read "Gloves are to be worn whenever there may be direct contact between the caregiver's hands and blood, body fluids, secretions, feces, or a contaminated item, such as soiled linens or wound dressings." Additionally, the policy read, "Complete task, if gloves become torn or heavily soiled and additional tasks must be performed for the client, then change the gloves (washing hands before putting on new gloves before starting a new task). Always remove gloves before working with clean areas and re-wash hands. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	0 510		

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0 650	Continued From page 4	0 650		
0 650 SS=D	144G.42 Subd. 8 Employee records (a) The facility must maintain current records of each paid employee, each regularly scheduled volunteer providing services, and each individual contractor providing services. The records must include the following information: (1) evidence of current professional licensure, registration, or certification if licensure, registration, or certification is required by this chapter or rules; (2) records of orientation, required annual training and infection control training, and competency evaluations; (3) current job description, including qualifications, responsibilities, and identification of staff persons providing supervision; (4) documentation of annual performance reviews that identify areas of improvement needed and training needs; (5) for individuals providing assisted living services, verification that required health screenings under subdivision 9 have taken place and the dates of those screenings; and (6) documentation of the background study as required under section 144.057. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the facility failed to ensure one of two employee records (unlicensed personnel (ULP)-E) contained the required content. 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	0 650		

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0 650	Continued From page 5 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: During the entrance conference on April 24, 2023, at 10:50 a.m. licensed assisted living director (LALD)-A stated the facility used supplemental staffing agencies to meet their staffing needs for unlicensed personnel. ULP-E started providing services to the licensee via a contracted supplemental staffing agency on January 9, 2023. On April 25, 2023, at 7:00 a.m. ULP-D was observed to hand ULP-E diclofenac 1% gel (for mild joint pain), and antifungal cream (for R3's abdominal skin folds and buttock crease area). ULP-E was observed to apply the diclofenac 1% gel to R3's left foot and ankle, and the antifungal cream to his abdominal skin folds and to his buttock crease area and continued with the remainder of R3's personal cares, applied compression stockings, and finished dressing him for the day. ULP-E's employee record lacked the required content of ULP-E's training and competency by the licensee's registered nurse (RN) for the administration of treatments such as prescribed creams and lotions. On April 26, 2023, at 10:45 a.m. clinical nurse supervisor (CNS)-B stated ULP-E was a "temp staff" (temporary or pool staff from a supplemental staffing agency) and was a certified nursing assistant (CNA) who had previous medication training by the staffing agency. The	0 650		

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0 650	Continued From page 6 licensee's practice was for "temp staff" to perform certain tasks with resident cares but not pass medications or complete treatments. A former licensee RN trained ULP-E to provide topical creams and lotions during resident cares when those were provided by the designated medication passer; however, CNS-B stated there was no documentation of ULP-E's training/competency to provide the topical creams. The licensee's Personnel Records policy dated November 14, 2022, indicated the personnel record for each person would include a record of all required training for unlicensed personnel and competency evaluations. No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-One (21) days	0 650		
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	0 680		

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0 680	Continued From page 7 missing residents. (b) The facility must provide emergency and disaster training to all staff during the initial staff 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 observation, interview, and record review, the licensee failed to practice and document testing of its emergency preparedness (EP) plan at least twice annually as required, post an emergency disaster plan prominently, and evaluate its missing person policy at least quarterly. This had the potential to affect all residents, staff, and visitors of the facility. 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 the residents). The findings include: EP PLAN TESTING The EP plan lacked evidence the facility had tested its EP plan, conducted and/or participated in a full-scale community-based exercise, or conducted a tabletop exercise to test it's plan and document the results of testing at least twice	0 680		

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0 680	Continued From page 8 annually as required. On April 26, 2023, at 8:50 a.m. licensed assisted living director (LALD)-A stated she had not conducted any EP plan testing in the past year. LALD-A indicated she was not aware of the requirement. On April 26, 2023, at 9:30 a.m. clinical nurse supervisor (CNS)-B stated she had not kept LALD-A informed of the emergency preparedness testing requirements indicating this facility was in a different coalition from the other building she oversaw. EMERGENCY DISASTER PLAN POSTED The licensee failed to prominently post the emergency disaster plan. On April 24, 2023, at 12:00 p.m. observation of the facility identified no posted facility emergency disaster plan. On April 25, 2023, at 1:25 p.m. operations manager (OM)-F stated the Emergency Disaster Plan binder was kept within the nurse's station and was not sure it was posted prominently anywhere else in the facility. MISSING RESIDENT PLAN/POLICY The licensee's Missing Resident policy dated January 12, 2023, as part of the annual EP review, was included in the EP plan. The policy lacked evidence the assisted living director and clinical registered nurse reviewed or updated the policy and documented any changes, at least quarterly as required. On April 26, 2023, at 9:30 a.m. CNS-B stated the missing resident plan was last reviewed on	0 680			

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0 680	Continued From page 9 January 12, 2023. CNS-B stated she was not aware the missing resident plan needed to be reviewed quarterly. The licensee's Emergency and Disaster Preparedness policy dated December 29, 2022, included: 7. A written policy and procedure regarding missing residents is complete in compliance with MN Rule 4659.0110 8. The facility will post the emergency disaster plan prominently 11. The facility conducts exercises to test the emergency preparedness plan at least twice per year in compliance with 42 C.F.R. 483.73(d)(2). No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 680		
0 730 SS=D	144G.43 Subd. 3 Contents of resident record Contents of a resident record include the following for each resident: (1) identifying information, including the resident's name, date of birth, address, and telephone number; (2) the name, address, and telephone number of the resident's emergency contact, legal representatives, and designated representative; (3) names, addresses, and telephone numbers of the resident's health and medical service providers, if known; (4) health information, including medical history, allergies, and when the provider is managing medications, treatments or therapies that require documentation, and other relevant health records;	0 730		

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0 730	Continued From page 10 (5) the resident's advance directives, if any; (6) copies of any health care directives, guardianships, powers of attorney, or conservatorships; (7) the facility's current and previous assessments and service plans; (8) all records of communications pertinent to the resident's services; (9) documentation of significant changes in the resident's status and actions taken in response to the needs of the resident, including reporting to the appropriate supervisor or health care professional; (10) documentation of incidents involving the resident and actions taken in response to the needs of the resident, including reporting to the appropriate supervisor or health care professional; (11) documentation that services have been provided as identified in the service plan; (12) documentation that the resident has received and reviewed the assisted living bill of rights; (13) documentation of complaints received and any resolution; (14) a discharge summary, including service termination notice and related documentation, when applicable; and (15) other documentation required under this chapter and relevant to the resident's services or status. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to ensure the resident record included a discharge summary with the required content for one of one discharged resident (R1). This practice resulted in a level two violation (a violation that did not harm a resident's health or	0 730		

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0 730	Continued From page 11 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 was admitted to the facility on February 1, 2021, with a diagnosis of obesity ,and discharged to a skilled nursing facility following hospitalization on February 20, 2023. R1's service plan signed by the resident on January 10, 2023, noted services included medication administration, assistance with dressing/undressing, bathing, standby help with walking long distances, light housekeeping, and laundry. R1's record lacked a discharge summary to include: - a summary of the resident's stay including diagnoses, course of illnesses, allergies, treatments and therapies, and pertinent lab, radiology, and consultation results; - a final summary of the resident's status from the latest assessment or review including baseline and current mental, behavioral, and functional status; and -a postdischarge plan to identify any arrangements that have been made for the resident's follow-up care, and any postdischarge medical and nonmedical services the resident will need. On April 24, 2023, at 3:00 p.m. clinical nurse supervisor (CNS)-B stated the nurse that	0 730		

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0 730	Continued From page 12 completed the discharge summary was new and did not complete all required components of the discharge summary form. The licensee's Discharge Summary policy dated reviewed November 28, 2022, indicated: 4. The discharge summary will include, if applicable, a. summary of the resident's stay, including i. diagnosis ii. courses of illnesses, iii. allergies iv. treatments v. therapies vi. pertinent lab results vii. pertinent radiology results viii. pertinent consultation results ix. final summary of the resident's status. The summary will be based upon the latest assessment or review and, if applicable, will include resident status including baseline and current mental, behavioral, and functional status. d. The post discharge plan will include: i. Resident's planned residence ii. Arrangements made for follow-up care iii. Needed post discharge medical and nonmedical services No further information was provided. TIME PERIOD FOR CORRECTIONS: Twenty-one (21) days	0 730			
0 780 SS=F	144G.45 Subd. 2 (a) (1) Fire protection and physical environment (a) Each assisted living facility must comply with the State Fire Code in Minnesota Rules, chapter 7511, and:	0 780			

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0 780	Continued From page 13 (1) for dwellings or sleeping units, as defined in the State Fire Code: (i) provide smoke alarms in each room used for sleeping purposes; (ii) provide smoke alarms outside each separate sleeping area in the immediate vicinity of bedrooms; (iii) provide smoke alarms on each story within a dwelling unit, including basements, but not including crawl spaces and unoccupied attics; (iv) where more than one smoke alarm is required within an individual dwelling unit or sleeping unit, interconnect all smoke alarms so that actuation of one alarm causes all alarms in the individual dwelling unit or sleeping unit to operate; and (v) ensure the power supply for existing smoke alarms complies with the State Fire Code, except that newly introduced smoke alarms in existing buildings may be battery operated; This MN Requirement is not met as evidenced by: Based on observation and interview, the licensee failed to provide smoke alarms that complied with fire protection requirements. This deficient condition had the potential to affect all staff, residents, and visitors. 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).	0 780		

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0 780	Continued From page 14 Findings include: On April 27, 2023, between 9:30 a.m. and 10:45 a.m., survey staff toured the facility with the director of maintenance (DM)-H, the licensed assisted living director (LALD)-A, and operations manager (OM)-F. During the facility tour, survey staff observed resident sleeping rooms in several of the units did not have the required smoke alarms. This deficient condition was verified by DM-H, LALD-A, and OM-F accompanying on the facility tour. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	0 780		
01370 SS=D	144G.61 Subd. 2 (a) Training and evaluation of unlicensed personn (a) Training and competency evaluations for all unlicensed personnel must include the following: (1) documentation requirements for all services provided; (2) reports of changes in the resident's condition to the supervisor designated by the facility; (3) basic infection control, including blood-borne pathogens; (4) maintenance of a clean and safe environment; (5) appropriate and safe techniques in personal hygiene and grooming, including: (i) hair care and bathing; (ii) care of teeth, gums, and oral prosthetic devices; (iii) care and use of hearing aids; and (iv) dressing and assisting with toileting;	01370		

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01370	Continued From page 15 (6) training on the prevention of falls; (7) standby assistance techniques and how to perform them; (8) medication, exercise, and treatment reminders; (9) basic nutrition, meal preparation, food safety, and assistance with eating; (10) preparation of modified diets as ordered by a licensed health professional; (11) communication skills that include preserving the dignity of the resident and showing respect for the resident and the resident's preferences, cultural background, and family; (12) awareness of confidentiality and privacy; (13) understanding appropriate boundaries between staff and residents and the resident's family; (14) procedures to use in handling various emergency situations; and (15) awareness of commonly used health technology equipment and assistive devices. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to ensure training and competency evaluations were completed as required prior to providing direct care for one of two employees (unlicensed personnel (ULP)-G). 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).	01370		

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01370	Continued From page 16 The findings include: ULP-G had a hire date of October 24, 2022, to provide services under the licensee's assisted living with dementia care. ULP-G was not a registered nursing assistant. ULP-G's record lacked evidence of the following competencies had been completed prior to providing direct cares: - appropriate and safe techniques in personal hygiene and grooming, including: -care of teeth, gums, and oral prosthetic devices; -care and use of hearing aids; and -dressing On April 26, 2023, at 11:20 a.m. clinical nurse supervisor (CNS)-B verified the above topics were lacking from ULP-G's employee file. CNS-B stated a previously employed registered nurse (RN) had completed competencies with ULP-G, but the above required areas were not signed off as completed on ULP-G's skills validation form. The licensee's Assisted Living Orientation-ULP Staff policy dated reviewed November 29, 2022, included: In addition to training all staff receive, ULPs who are not a registered nursing assistant will receive additional training on the following topics with a written or oral competency test AND a skill demonstration: c. care of teeth, gums (sic), and oral prosthetic devices d. care and use of hearing aids e. dressing No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-one	01370		

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01370	Continued From page 17 (21) days	01370		
01540 SS=D	144G.64 (a) TRAINING IN DEMENTIA CARE REQUIRED (3) for assisted living facilities with dementia care, direct-care employees must have completed at least eight hours of initial training on topics specified under paragraph (b) within 80 working hours of the employment start date. Until this initial training is complete, an employee must not provide direct care unless there is another employee on site who has completed the initial eight hours of training on topics related to dementia care and who can act as a resource and assist if issues arise. A trainer of the requirements under paragraph (b) or a supervisor meeting the requirements in clause (1) must be available for consultation with the new employee until the training requirement is complete. Direct-care employees must have at least two hours of training on topics related to dementia for each 12 months of employment thereafter; This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to ensure one of two employees (unlicensed personnel (ULP)-G) completed the required amount of dementia care training in the required time frame. 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	01540		

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01540	Continued From page 18 a limited number of staff are involved or the situation has occurred only occasionally). The findings include: ULP-G had a hire date of October 24, 2022, to provide direct care services under the licensee's assisted living with dementia care license. ULP-G's employee record contained evidence of four hours of training on dementia care topics, instead of the required eight hours. On April 26, 2023, at 11:20 a.m. clinical nurse supervisor (CNS)-B stated ULP-G had worked approximately 160 hours since her start date. CNS-B verified ULP-G lacked the required eight hours of dementia training within 80 working hours as required. The licensee's Assisted Living and Assisted Living with Memory Care Dementia Training policy dated reviewed November 29, 2022, noted direct-care employees would: 1. complete a minimum of eight hours of initial training on dementia care topics; and 2. Initial training will be completed within 80 working hours of the employment start date. No further information was provided. TIME PERIOD FOR CORRECTION: Twenty-one (21) days	01540			
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 that services are first provided, an assisted living	01640			

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01640	Continued From page 19 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 one of two resident's (R3) service plans were revised to reflect the current services provided. In addition, the licensee failed to ensure the service plan included a signature or other authentication by the resident and the facility to document agreement on the services provided for one of two 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	01640		

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01640	Continued From page 20 a limited number of staff are involved or the situation has occurred only occasionally). The findings include: R3's diagnoses included congestive heart failure (impairment of the heart's blood pumping function), atrial fibrillation (an irregular heartbeat), diabetes, and obstructive sleep apnea (a condition when a person obstructs their airway during sleep causing an interruption of the breathing pattern). R3 was also legally blind and had a right prosthetic eye. On April 25, 2023, at 11:30 a.m. the surveyor observed an oxygen concentrator (a machine/device which concentrates the air in the room to supply an oxygen-enriched product to a person) in R3's room with green oxygen tubing connected to a CPAP (continuous positive airway pressure)/Bi-PAP (two level positive airway pressure) machine which sat on R3's bedside table. The surveyor also observed a document labeled "Instruction for CPAP or Bi-PAP" taped to the wall above the machine. The instructions read: -Add DISTILLED water to humidifying reservoir. To fill line and replace reservoir -Unclip head strap from mask on one side and gather head strap in one hand. -Assist resident in properly placing mask over nose, mouth and forehead. (may have a separate gel piece that goes over nose -Slide head strap over top of head and under ears -Reclip head strap to mask. Make sure strap is tightened equally on both sides if it needs readjusting -Turn machine and oxygen on. -Check for a proper seal around the mask making sure there are no air leaks (if there is a leak	01640		

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01640	Continued From page 21 recheck placement of mask or tightness of straps) The instructions lacked the indication of what flow rate the concentrator should be set to. On April 25, 2023, at 11:35 a.m. R3 stated he used his CPAP/Bi-PAP machine each night and had oxygen "at about six liters, I think". R3 stated the staff assist him with all use of the CPAP/Bi-PAP machine and oxygen. R3's Task sheet dated April 2023, indicated CPAP assistance at bedtime; however, it lacked direction or an indication of oxygen use/flow rate. Additionally, R3's task sheet indicated twice daily blood sugar checks. R3's provider notes/orders dated April 19, 2023, included an order for oxygen at a flow rate of two liters per minute connected to R3's CPAP/Bi-PAP machine during sleep. R3's unsigned Service Plan dated March 14, 2023, indicated R3 received services to medication management, TED (thrombo-embolic deterrent) stockings (used to prevent blood clots and reduce lower extremity swelling), CPAP or Bi-PAP, blood sugar checks twice daily, toileting/bathing/dressing assistance and the use of an EZ stand (mechanical lift system) for all transfers. The service plan failed to identify the treatment of oxygen as connected to his CPAP/Bi-PAP machine. R3's Service Plan was revised on March 14, 2023, to include twice daily blood sugar check (previously was once weekly) and a daily insulin injection (a new order March 7, 2023). The licensee failed to ensure the Service Plan was signed or authenticated by the licensee or the	01640		

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01640	Continued From page 22 resident at the time of revision. On April 25, 2023, at 1:30 p.m. operations manager (OM)-F confirmed R3's service plan dated March 14, 2023, lacked a signature by R3. On April 26, 2023, at 9:30 a.m. clinical nurse supervisor (CNS)-B stated she was not aware of the oxygen being used with R3's CPAP/Bi-PAP machine. CNS-B confirmed the treatment of oxygen was not on R3's service plan. The licensee's Contents of Service Plans policy dated November 28, 2022, indicated all Assisted Living residents have an up-to date Service Plan identifying services to be provided based on the assessment by the registered nurse (RN) and/or other licensed health professionals. Service plans and any revisions to Service Plans would have a signature or other authentication by the facility and by the resident. No further information was provided. TIME PERIOD FOR CORRECTION: 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	01750		

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01750	Continued From page 23 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 insulin administration via a prefilled insulin pen was administered according to manufacturer instructions, for one of one resident (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) 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: R3's diagnoses included congestive heart failure (impairment of the heart's blood pumping function), atrial fibrillation (an irregular heartbeat), diabetes (the body's inability to adequately manage blood sugar), and obstructive sleep apnea (a condition when a person obstructs their airway during sleep causing an interruption of the breathing pattern). R3 was also legally blind and had a right prosthetic eye. R3's unsigned Service Plan dated March 14, 2023, indicated R3 received services to include medication management, daily insulin injections, and blood sugar checks twice daily. R3's provider's orders dated April 18, 2023, indicated Lantus Solostar insulin 11 units	01750		

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01750	Continued From page 24 subcutaneously (injected at a shallow depth in the skin) daily. On April 25, 2023, at 7:30 a.m. unlicensed personnel (ULP)-D was observed at the medication cart setting up R3's morning medications. ULP-D removed R3's Lantus insulin pen cartridge from the medication cart and immediately attached a pen needle to the end of the cartridge. ULP-D primed the insulin pen with the designated two units of insulin and dialed the insulin pen to the prescribed 11 units of insulin. ULP-D was then observed to take the insulin pen to R3, told R3 what she was doing, pulled up his t-shirt, and administered/injected the insulin into his left upper abdomen and held the insulin pen in place for 10 seconds as per protocol. ULP-D then removed the insulin pen from the abdomen, repositioned his t-shirt, returned to the medication cart, disconnected the needle end, and returned the insulin cartridge to his compartment in the medication cart. ULP-D failed to wipe the insulin pen cartridge tip prior to placing the needle on it and failed to properly cleanse (alcohol wipe) R3's skin prior to the insulin injection. On April 26, 2023, at 9:19 a.m. clinical nurse supervisor (CNS)-B stated she expected the staff would follow the proper procedure and wipe the tip of the insulin cartridge prior to applying a new needle and then properly cleanse the resident's skin prior to inserting a needle and insulin. The licensee's undated, Insulin Administration-Skills Validation indicated "select injection site and swab with alcohol. Allow area to dry prior to injection. The Insulin Administration-Skills Validation did not address the need to wipe the insulin tip.	01750		

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01750	Continued From page 25 Lantus Solostar manufacturer's instructions dated 2022, instructed users to wipe the tip of the insulin cartridge with an alcohol wipe prior to applying the needle. Additionally, the instructions indicated to cleanse the injection site with an alcohol wipe prior to insulin injection. No further information was 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 time sensitive medications were dated when opened for one of two residents (R4) and failed to discard a medication when it had exceeded its recommended period of safe use time once opened for one of one resident (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	01890		

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01890	Continued From page 26 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: On April 25, 2023, at 9:55 a.m. the surveyor observed the medication cart for the memory care residents with unlicensed personnel (ULP)-D. R4 R4's latanoprost (treats increased pressure inside the eye) 0.005% eye drops lacked a date to indicate when staff opened it, or when it would expire. ULP-D confirmed the findings at this time and verified eye drops should be dated when opened and when they would expire. On April 25, 2023, at 9:20 a.m. CNS-B stated all eye drops should be dated when opened. The latanoprost manufacturer's instructions package insert revised August 2011, noted: -Once a bottle is opened for use, it may be stored at room temperature up to six weeks. R3 The licensee failed to discard R3's eye drops when they had exceeded the recommended period of time (28 days) of safe use, once opened. On April 25, 2023, at 7:30 a.m. the surveyor observed the medication cart with ULP-D. ULP-D pulled out R3's 1.4% artificial tears eye drops from the medication cart as needed for his 8:00	01890		

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01890	Continued From page 27 a.m. medication administration time. The surveyor observed the eye drop bottle included a date opened label written as "March 23" and included additional print that read do not use after 28 days, once opened. ULP-D stated she had just reviewed the medication cart the day before and had not recognized the date had then exceeded 28 days since opened. ULP-D indicated the date "March 23" written on the label was recognized as March 23, 2023. ULP-D confirmed the eye drops had now been opened for 33 days (five days past the recommended use date). On April 26, 2023, at 9:21 a.m. CNS-B stated she was made aware of R3's eye drops and the bottle in use was beyond the 28-day time period for safe use and stated she had completed a medication cart audit the week before and did not recognize the due date for reorder was approaching and failed to order a new bottle. The manufacturer's package inserts dated July 15, 2009, indicated "You must throw away the bottle 28 days after you first opened it, even if there are still some drops left." The licensee's Storage of Medications policy dated November 23, 2022, indicated medications would be handled and stored per acceptable standards. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	01890		

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01910	Continued From page 28	01910		
01910 SS=D	144G.71 Subd. 22 Disposition of medications (a) Any current medications being managed by the assisted living facility must be provided to the resident when the resident's service plan ends or medication management services are no longer part of the service plan. Medications for a resident who is deceased or that have been discontinued or have expired may be provided for disposal. (b) The facility shall dispose of any medications remaining with the facility that are discontinued or expired or upon the termination of the service contract or the resident's death according to state and federal regulations for disposition of medications and controlled substances. (c) Upon disposition, the facility must document in the resident's record the disposition of the medication including the medication's name, strength, prescription number as applicable, quantity, to whom the medications were given, date of disposition, and names of staff and other individuals involved in the disposition. This MN Requirement is not met as evidenced by: Based on interview and record review, the licensee failed to document in the resident's record the disposition of the medication including the medication strength, for one of one discharged resident (R1). 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	01910		

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01910	Continued From page 29 situation has occurred only occasionally). The findings include: R1 was discharged to a skilled nursing facility on February 20, 2022. R1's record included an untitled form that had columns to identify the date, medication, RX# (medical prescription number), quantity, location of disposal, nurse signature and witness signature. The document did not include a column for the medication strength. - The following medications were listed without a strength: vitamin D3 (supplement), duloxetine (antidepressant), levothyroxine (thyroid hormone), senna plus (constipation), Nystop powder (antifungal), and ferrous sulfate (iron deficiency). On April 24, 2023, at 3:50 p.m. clinical nurse supervisor (CNS)-B stated though the strength was identified for most of the medications listed on the form, the strength of medications as listed above was missing on the disposition log for R1. CNS-B indicated she may need to alter the form to include a column for strength to ensure it is not missed. The licensee's Disposition or Disposal of Medication policy dated reviewed November 23, 2022, noted documentation of the destruction, listing the date, quantity, name of drug, prescription number, signature of person destroying the drugs and signature of witness to the destruction must be recorded and maintained in the resident's record for two years. The policy lacked information to include the medication strength. No further information was provided.	01910		

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01910	Continued From page 30	01910		
01940 SS=D	144G.72 Subd. 3 Individualized treatment or therapy management For each resident receiving management of ordered or prescribed treatments or therapy services, the assisted living facility must prepare and include in the service plan a written statement of the treatment or therapy services that will be provided to the resident. The facility must also develop and maintain a current individualized treatment and therapy management record for each resident which must contain at least the following: (1) a statement of the type of services that will be provided; (2) documentation of specific resident instructions relating to the treatments or therapy administration; (3) identification of treatment or therapy tasks that will be delegated to unlicensed personnel; (4) procedures for notifying a registered nurse or appropriate licensed health professional when a problem arises with treatments or therapy services; and (5) any resident-specific requirements relating to documentation of treatment and therapy received, verification that all treatment and therapy was administered as prescribed, and monitoring of treatment or therapy to prevent possible complications or adverse reactions. The treatment or therapy management record must be current and updated when there are any changes. This MN Requirement is not met as evidenced	01940		

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01940	Continued From page 31 by: Based on observation, interview, and record review, the licensee failed to develop and implement a treatment or therapy management plan to include all required content for one of two 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: During the entrance conference on April 24, 2023, at 10:50 a.m. licensed assisted living director (LALD)-A stated the licensee provided treatment management services to residents, including oxygen. R3's diagnoses included congestive heart failure (impairment of the heart's blood pumping function) and obstructive sleep apnea (a condition when a person obstructs their airway during sleep causing an interruption of the breathing pattern). R3's unsigned Service Plan dated March 14, 2023, indicated R3 received services to include assistance with the application and removal of a CPAP (continuous positive airway pressure) or Bi-PAP (two level positive airway pressure), every night at bedtime. The service plan failed to identify the treatment of oxygen as connected to his CPAP/Bi-PAP machine.	01940		

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01940	Continued From page 32 On April 25, 2023, at 11:30 a.m. the surveyor observed an oxygen concentrator (a machine/device which concentrates the air in the room to supply an oxygen-enriched product to a person) in R3's room with green oxygen tubing connected to a CPAP machine which sat on R3's bedside table. The surveyor also observed a document labeled "Instruction for CPAP or Bi-PAP" taped to the wall above the machine. The instructions read: -Add DISTILLED water to humidifying reservoir. To fill line and replace reservoir -Unclip head strap from mask on one side and gather head strap in one hand. -Assist resident in properly placing mask over nose, mouth, and forehead. (may have a separate gel piece that goes over nose -Slide head strap over top of head and under ears -Reclip head strap to mask. Make sure strap is tightened equally on both sides if it needs readjusting -Turn machine and oxygen on. -Check for a proper seal around the mask making sure there are no air leaks (if there is a leak recheck placement of mask or tightness of straps) The instructions lacked the indication of what flow rate the concentrator should be set to. On April 25, 2023, at 11:35 a.m. R3 stated he used his CPAP/Bi-PAP machine each night and had oxygen "at about six liters, I think". R3 stated the staff assist him with all use of the CPAP/Bi-PAP machine and oxygen. R3's provider notes/orders dated April 19, 2023, included an order for oxygen at a flow rate of two liters per minute connected to R3's CPAP/Bi-PAP machine during sleep.	01940		

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01940	Continued From page 33 R3's Task sheet dated April 2023, indicated CPAP assistance at bedtime; however, it lacked direction or an indication of oxygen use/flow rate. R3's Level of Care Form V6 (90-day assessment) dated March 14, 2023, in section M identified as "Respiratory Equipment, included the treatment of CPAP or Bi-PAP, however the licensee failed to identify the use of oxygen or oxygen equipment. R3's 90-day assessment also contained an Individualized Treatment and Therapy Management Plan; however, the plan lacked the following required content for the use of oxygen: (1) a statement of the type of services that will be provided; (2) documentation of specific resident instructions relating to the treatments or therapy administration; (3) identification of treatment or therapy tasks that will be delegated to unlicensed personnel; (4) procedures for notifying a registered nurse or appropriate licensed health professional when a problem arises with treatments or therapy services; and (5) any resident-specific requirements relating to documentation of treatment and therapy received, verification that all treatment and therapy was administered as prescribed, and monitoring of treatment or therapy to prevent possible complications or adverse reactions. The treatment or therapy management record must be current and updated when there are any changes. On April 26, 2023, at 9:30 a.m. clinical nurse supervisor (CNS)-B stated R3's treatment order was new on April 5, 2023, and she was not aware R3 had oxygen with his CPAP machine. The licensee's Treatment Management Plan	01940		

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01940	Continued From page 34 policy dated November 23, 2022, indicated a registered nurse (RN) would develop a treatment management plan for each resident receiving treatment management services. The treatment management plan would include: 1. [Licensee name] staff will provide treatment/therapy management services only after a face-to-face assessment has been performed by our Registered Nurse. 2. The RN will assess the resident on pre-admission, admission, at 14-day, every 90-day and anytime there is a change in conditions. 3. The RN will discuss with the resident what treatment/therapy they are receiving the reason for the treatment/therapy. 4. The RN will obtain orders for treatment/therapies for services provided. 5. The treatment or therapy services will be recorded on the service plan/task. 6. The RN will identify what treatment and therapies will be delegated to the unlicensed staff and train them on the treatment and therapy prior to having them perform the treatment/therapy. 7. The nurse will document any specific instruction related to the treatment and therapy administration on the record for the staff to view. 8. The nurse will have a procedure for when to notify the nurse and a procedure of notifying a registered nurse or appropriate licensed health professional when a problem arises with treatments or therapy services. 9. The RN will assess the resident every 90 days or as needed based on the treatment/therapy plan that is in place. 10. The RN will update the provider on treatment or therapy status changes as needed or based on orders. 11. Treatment and therapy orders will be renewed yearly or when there is a change in	01940		

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01940	Continued From page 35 treatment/therapy needed. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	01940		
01960 SS=D	144G.72 Subd. 5 Documentation of administration of treatments Each treatment or therapy administered by an assisted living facility must be in the resident record. The documentation must include the signature and title of the person who administered the treatment or therapy and must include the date and time of administration. When treatment or therapies are not administered as ordered or prescribed, the provider must document the reason why it was not administered and any follow-up procedures that were provided to meet the resident's needs. This MN Requirement is not met as evidenced by: Based on observation, interview, and record review, the licensee failed to ensure treatment or therapies were administered as prescribed 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).	01960		

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01960	Continued From page 36 The findings include: R2's diagnoses included dementia (memory loss), atherosclerosis of native arteries of right leg (narrowing of the leg arteries), non-pressure chronic ulcer (an open sore or wound that develops on the skin caused by poor blood circulation or injury) of right lower leg. R2's service plan signed by R2's responsible party on January 13, 2023, indicated R2 received dressing changes to the right heel. R2's prescriber orders dated April 19, 2023, included an order to clean right heel wound with wound cleanser (a solution used to remove foreign materials and microorganisms from wounds), pat dry, apply Iodosorb (anti-infective ointment that promotes healing of skin ulcers and wounds) topically, cover with foam border dressing, change every 3 days, and as needed. R2's April 2023, medication administration record (MAR) indicated R2 received wound care to right heel every three days and as needed when soiled. Remove old dressing, wash hands and put on new gloves. Wash wound out with wound cleanser and gauze pad. Apply a pea size amount of Iodosorb ointment directly into wound bed. Cover ulcer site with Mepilex (foam boarder dressing) 2 x 2 dressing and tape as needed. On April 25, 2023, at 8:56 a.m. unlicensed personnel (ULP)-D was observed to change R2's right heel dressing. ULP-D sanitized hands and applied gloves. ULP-D removed R2's right sock and dressing. Drainage was present on R2's sock. ULP-D removed their gloves and sanitized their hands, and stated she needed to notify the	01960		

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01960	Continued From page 37 registered nurse (RN) due to the increase in drainage. RN-C came into the room and assessed R2's heel wound, took measurements, asked R2 and ULP-D questions, and instructed ULP-D she could continue with dressing change. ULP-D then obtained R2's wound care supplies contained in a plastic tub on upper shelf of his closet, sanitized their hands and applied clean gloves. ULP-D took a container of acetic acid (vinegar water) 0.25% and saturated a clean 4x4 gauze pad with the solution. ULP-D then put the saturated gauze pad to R2's open wound and gently scrubbed the wound bed while squeezing the acetic acid from gauze pad onto the wound. After this, ULP-D grabbed the wound cleanser and sprayed the wound area while using a clean gauze pad to "pat dry" the area. ULP-D removed their gloves, sanitized their hands, and applied a clean pair of gloves. ULP-D applied a pea size amount of Iodosorb to a sterile Q-tip and spread onto the wound bed. Following this, ULP-D took a foam dressing, covered the wound area, and taped it into place. ULP-D removed their gloves, sanitized their hands, and put R2's wound care supplies away. Immediately following the wound care procedure, ULP-D stated she has been using the acetic acid to R2's wound "for a long time". ULP-D then acknowledged there was no current order for the acetic acid indicating she had not followed the ordered treatment. ULP-D further stated, "he's had a lot of treatment changes". On April 25, 2023, at 2:12 p.m. RN-C stated she would expect wound care to be completed as ordered. RN-C further stated she would remove the acetic acid from R2's wound care container indicating it should have been removed when the order changed.	01960		

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01960	Continued From page 38 The licensee's Administration of Medication, Treatment and Therapy by Unlicensed Personnel policy dated reviewed November 23, 2022, included: 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 (by mouth, eye drops, to the skin, etc.) e. right dose (how many milligrams, drops, etc.) f. right chart/record to document that the medication, treatment, and therapy was taken. No further information was provided. TIME PERIOD FOR CORRECTION: Seven (7) days	01960		

Type: Full
Date: 04/26/23
Time: 10:15:52
Report: 8044231136

Food and Beverage Establishment Inspection Report

Page 1

Location:

Cedar Court
810 West Main Street
Adams, MN55909
Mower County, 50

Establishment Info:

ID #: 0038229
Risk:
Announced Inspection: No

License Categories:

Expires on: / /

Operator:

Phone #: 5075823263
ID #:

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

The following orders were issued during this inspection.

3-800 Highly Susceptible Populations

3-801.11B ** Priority 1 **

MN Rule 4626.0447B Discontinue using unpasteurized eggs or egg products in the preparation of Caesar salad, hollandaise or Bearnaise sauce, mayonnaise, meringue, eggnog, ice cream, and egg-fortified beverages when serving a highly susceptible population.

Unpasteurized eggs used for undercooked fried eggs.

Comply By: 04/26/23

5-200B Plumbing: cross connections

5-203.14A ** Priority 1 **

MN Rule 4626.1085A Water used under pressure in equipment in food and beverage establishments must be drained to a sanitary sewer through an air gap. Examples: refrigeration cooling water, water softener, and drained steam jacketed kettles.

Air gap for water softener discharge hose below flood rim.

Comply By: 05/10/23

5-200B Plumbing: cross connections

5-203.14C ** Priority 1 **

MN Rule 4626.1085A Provide a reduced pressure zone (RPZ) backflow preventer to protect water distribution lines subject to back pressure, such as the water supply serving a boiler.

No backflow preventer for boiler.

Comply By: 04/26/24

Type: Full
Date: 04/26/23
Time: 10:15:52
Report: 8044231136
Cedar Court

Food and Beverage Establishment Inspection Report

Page 2

2-100 Supervision

2-102.12DMN

MN Rule 4626.0033D Post the certified food protection manager certificate.

State of Minnesota certificate not posted.

Comply By: 04/26/23

3-300C Protection from Contamination: equipment/utensils, consumers

3-304.14B

MN Rule 4626.0285B Wiping cloths used for wiping counters and other equipment surfaces must be held in an approved sanitizing solution and laundered daily.

Quaternary ammonium in wiping bucket measured at 100 ppm.

Comply By: 04/26/23

4-500 Equipment Maintenance and Operation

4-501.11AB

MN Rule 4626.0735AB All equipment and components must be in good repair and maintained and adjusted in accordance with manufacturer's specifications.

Walk in freezer leaking condensate from condenser.

Comply By: 05/26/23

4-900 Protecting Clean Items

4-903.12A

MN Rule 4626.0960A Discontinue storage of food, clean equipment, linens, utensils, or single-service and single-use articles in locker rooms; toilet rooms; garbage rooms; mechanical rooms; under unshielded sewer lines; under leaking water lines or sources of condensation or moisture; under open stairwells; or under other sources of contamination.

No splash guard for handwashing sink next to prep table.

Discontinue storing or prepping food next to prep sink until splash guard is installed.'

Condensate dripping onto bags of food in walk in freezer.

Comply By: 04/26/23

Surface and Equipment Sanitizers

Quaternary Ammonia: = 100 ppm at Degrees Fahrenheit

Location: Bucket

Violation Issued: Yes

Hot Water: = at 167.9 Degrees Fahrenheit

Location: Dishwasher

Violation Issued: No

Type: Full
Date: 04/26/23
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Food and Beverage Establishment Inspection Report

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Food and Equipment Temperatures

Process/Item: Cooking
Temperature: 208 Degrees Fahrenheit - Location: Rice
Violation Issued: No

Process/Item: Cooking
Temperature: 160.0 Degrees Fahrenheit - Location: Pork chop
Violation Issued: No

Process/Item: Cold Holding
Temperature: 39.0 Degrees Fahrenheit - Location: Milk in upright
Violation Issued: No

Process/Item: Cold Holding
Temperature: 42.0 Degrees Fahrenheit - Location: Upright cooler
Violation Issued: No

Process/Item: Cold Holding
Temperature: 39.0 Degrees Fahrenheit - Location: WIC
Violation Issued: No

Process/Item: Cold Holding
Temperature: 41.0 Degrees Fahrenheit - Location: Tuna salad in WIC
Violation Issued: No

Process/Item: Cold Holding
Temperature: 41.0 Degrees Fahrenheit - Location: Turkey in WIC
Violation Issued: No

Total Orders	In This Report	Priority 1	Priority 2	Priority 3
		3	0	4

HRD inspection conducted with Susan Kalis, Deb Jacobson and Rob Davis.

Inspection report reviewed on site with

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

I acknowledge receipt of the Minnesota Department of Health inspection report number 8044231136 of 04/26/23.

Certified Food Protection Manager Rhonda J. Musel

Certification Number: FM19642 Expires: 06/25/23

Inspection report reviewed with person in charge and emailed.

Signed: 
Inspector signed for Rhonda

Signed: 
Michael DeMars, RS
Public Health Sanitarian III
Rochester District Office
507-206-4715
michael.demars@state.mn.us