



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 24, 2023

Administrator
The Waterview Pines LLC
1201 8th Street South
Virginia, MN 55792

RE: CCN: 245283
Cycle Start Date: January 11, 2023

Dear Administrator:

On February 16, 2023, we notified you a remedy was imposed. On February 17, 2023 the Minnesota Department of Health completed a revisit to verify that your facility had achieved and maintained compliance. We have determined that your facility has achieved substantial compliance as of February 9, 2023.

As authorized by CMS the remedy of:

- Mandatory denial of payment for new Medicare and Medicaid admissions effective April 11, 2023 did not go into effect. (42 CFR 488.417 (b))

In our letter of January 31, 2023, in accordance with Federal law, as specified in the Act at § 1819(f)(2)(B)(iii)(I)(b) and § 1919(f)(2)(B)(iii)(I)(b), we notified you that your facility was prohibited from conducting a Nursing Aide Training and/or Competency Evaluation Program (NATCEP) for two years from April 11, 2023 due to denial of payment for new admissions. Since your facility attained substantial compliance on February 9, 2023, the original triggering remedy, denial of payment for new admissions, did not go into effect. Therefore, the NATCEP prohibition is rescinded. However, this does not apply to or affect any previously imposed NATCEP loss.

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

Feel free to contact me if you have questions.

Sincerely,

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

Joanne Simon, Compliance Analyst
Minnesota Department of Health
Health Regulation Division
Telephone: 651-201-4161
Email: joanne.simon@state.mn.us
cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

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February 24, 2023

Administrator
The Waterview Pines LLC
1201 8th Street South
Virginia, MN 55792

Re: Reinspection Results
Event ID: LGOC12

Dear Administrator:

On February 22, 2023 survey staff of the Minnesota Department of Health - Health Regulation Division completed a reinspection of your facility, to determine correction of orders found on the survey completed on January 26, 2023. At this time these correction orders were found corrected.

Please feel free to call me with any questions.

Sincerely,

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

Joanne Simon, Compliance Analyst
Minnesota Department of Health
Health Regulation Division
Telephone: 651-201-4161
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered

February 2, 2023

Administrator
The Waterview Pines Llc
1201 8th Street South
Virginia, MN 55792

RE: CCN: 245283
Cycle Start Date: January 11, 2023

Dear Administrator:

On January 31, 2023, we informed you that we may impose enforcement remedies.

On January 26, 2023, the Minnesota Department of Health completed a survey and it has been determined that your facility is not in substantial compliance. The most serious deficiencies in your facility were found to be isolated deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level D), as evidenced by the electronically attached CMS-2567, whereby corrections are required.

REMEDIES

As a result of the survey findings and in accordance with survey and certification memo 16-31-NH, this Department recommended the enforcement remedy(ies) listed below to the CMS Region V Office for imposition. The CMS Region V Office concurs and is imposing the following remedy and has authorized this Department to notify you of the imposition:

- Mandatory Denial of Payment for new Medicare and/or Medicaid Admissions, Federal regulations at 42 CFR § 488.417(a), effective Medicaid Admissions, Federal regulations at 42 CFR § 488.417(a), effective April 11, 2023

The CMS Region V Office will notify your Medicare Administrative Contractor (MAC) that the denial of payment for new admissions is effective April 11, 2023. They will also notify the State Medicaid Agency that they must also deny payment for new Medicaid admissions effective April 11, 2023.

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

This Department is also recommending that CMS impose a civil money penalty. You will receive a formal notice from the CMS RO only if CMS agrees with our recommendation.

- Civil money penalty. (42 CFR 488.430 through 488.444)

NURSE AIDE TRAINING PROHIBITION

Please note that Federal law, as specified in the Act at §§ 1819(f)(2)(B) and 1919(f)(2)(B), prohibits approval of nurse aide training and competency evaluation programs and nurse aide competency evaluation programs offered by, or in, a facility which, within the previous two years, has operated under a § 1819(b)(4)(C)(ii)(II) or § 1919(b)(4)(C)(ii) waiver (i.e., waiver of full-time registered professional nurse); has been subject to an extended or partial extended survey as a result of a finding of substandard quality of care; has been assessed a total civil money penalty of not less than \$11,292, has been subject to a denial of payment, the appointment of a temporary manager or termination; or, in the case of an emergency, has been closed and/or had its residents transferred to other facilities.

If you have not achieved substantial compliance by April 11, 2023, the remedy of denial of payment for new admissions will go into effect and this provision will apply to your facility. Therefore, The Waterview Pines Llc will be prohibited from offering or conducting a Nurse Aide Training and/or Competency Evaluation Program (NATCEP) for two years from April 11, 2023. You will receive further information regarding this from the State agency. This prohibition is not subject to appeal. Further, this prohibition may be rescinded at a later date if your facility achieves substantial compliance prior to the effective date of denial of payment for new admissions. However, under Public Law 105-15, you may contact the State agency and request a waiver of this prohibition if certain criteria are met.

ELECTRONIC PLAN OF CORRECTION (ePOC)

Within ten (10) calendar days after your receipt of this notice, you must submit an acceptable ePOC for the deficiencies cited. An acceptable ePOC will serve as your allegation of compliance. Upon receipt of an acceptable ePOC, we will authorize a revisit to your facility to determine if substantial compliance has been achieved. The failure to submit an acceptable ePOC can lead to termination of your Medicare and Medicaid participation (42 CFR 488.456(b)).

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

- How corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- How the facility will identify other residents having the potential to be affected by the same deficient practice.
- What measures will be put into place, or systemic changes made, to ensure that the deficient practice will not recur.
- How the facility will monitor its corrective actions to ensure that the deficient practice is being corrected and will not recur.
- The date that each deficiency will be corrected.

- An electronic acknowledgement signature and date by an official facility representative.

DEPARTMENT CONTACT

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

Terri Ament, Rapid Response
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Duluth Technology Village
11 East Superior Street, Suite 290
Duluth, Minnesota 55802-2007
Email: teresa.ament@state.mn.us
Office: (218) 302-6151 Mobile: (218) 766-2720

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

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

VERIFICATION OF SUBSTANTIAL COMPLIANCE

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

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

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

We will also recommend to the CMS Region V Office and/or the Minnesota Department of Human Services that your provider agreement be terminated by July 11, 2023 (six months after the identification of noncompliance) if your facility does not achieve substantial compliance. This action is mandated by the Social Security Act at § 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR § 488.412 and § 488.456.

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

APPEAL RIGHTS

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

Steven.Delich@cms.hhs.gov

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

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

A request for a hearing should identify the specific issues, findings of fact and conclusions of law with which you disagree. It should also specify the basis for contending that the findings and conclusions are incorrect. At an appeal hearing, you may be represented by counsel at your own expense. If you have any questions regarding this matter, please contact Steven Delich, Program Representative at (312) 886-5216. Information may also be emailed to Steven.Delich@cms.hhs.gov.

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

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

Nursing Home Informal Dispute Process

The Waterview Pines Llc

February 2, 2023

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Minnesota Department of Health
Health Regulation Division
P.O. Box 64900
St. Paul, Minnesota 55164-0900

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

You must notify MDH at this website of your request for an IDR or IIDR within the 10 calendar day period allotted for submitting an acceptable electronic plan of correction. A copy of the Department's informal dispute resolution policies are posted on the MDH Information Bulletin website at: https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html

Please note that the failure to complete the informal dispute resolution process will not delay the dates specified for compliance or the imposition of remedies.

Feel free to contact me if you have questions.

Sincerely,

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

Joanne Simon, Compliance Analyst
Minnesota Department of Health
Health Regulation Division
Telephone: 651-201-4161
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File

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

PRINTED: 02/27/2023
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245283	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 01/26/2023
NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETION DATE
F 000	INITIAL COMMENTS On 1/25/23, and 1/26/23, a standard abbreviated survey was completed at your facility to conduct a complaint investigation. Your facility was found to be NOT in compliance with 42 CFR Part 483, Requirements for Long Term Care Facilities. The following complaint was found to be SUBSTANTIATED: H52837840C (MN00090347) no deficiencies were cited due to actions implemented by the facility prior to survey. However, an unrelated deficiency was cited at F609. The facility's plan of correction (POC) will serve as your allegation of compliance upon the Departments acceptance. Because you are enrolled in ePOC, your signature is not required at the bottom of the first page of the CMS-2567 form. Your electronic submission of the POC will be used as verification of compliance. Upon receipt of an acceptable electronic POC, an onsite revisit of your facility may be conducted to validate that substantial compliance with the regulations has been attained.	F 000			
F 609 SS=D	Reporting of Alleged Violations CFR(s): 483.12(b)(5)(i)(A)(B)(c)(1)(4) §483.12(c) In response to allegations of abuse, neglect, exploitation, or mistreatment, the facility must: §483.12(c)(1) Ensure that all alleged violations involving abuse, neglect, exploitation or mistreatment, including injuries of unknown source and misappropriation of resident property, are reported immediately, but not later than 2	F 609			2/1/23

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
Electronically Signed		02/16/2023

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

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

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F 609	Continued From page 1 hours after the allegation is made, if the events that cause the allegation involve abuse or result in serious bodily injury, or not later than 24 hours if the events that cause the allegation do not involve abuse and do not result in serious bodily injury, to the administrator of the facility and to other officials (including to the State Survey Agency and adult protective services where state law provides for jurisdiction in long-term care facilities) in accordance with State law through established procedures. §483.12(c)(4) Report the results of all investigations to the administrator or his or her designated representative and to other officials in accordance with State law, including to the State Survey Agency, within 5 working days of the incident, and if the alleged violation is verified appropriate corrective action must be taken. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to ensure significant medication errors (an error of which causes the resident discomfort or jeopardizes the resident's health and safety) were reported to the State Agency (SA) for 1 of 3 residents (R1) reviewed for significant medication errors. Findings include: R1's Admission Record dated 1/26/23, indicated R1's diagnoses included diabetes, long term insulin use, aphasia (loss of ability to understand or express speech, caused by brain damage) following a stroke, and dementia. R1's quarterly Minimum Data Set dated 1/23/23, indicated R1 was cognitively intact and was	F 609	F609 Reporting of Alleged Violations Immediate Corrective Action: Resident #1 continues to remain from adverse effects d/t medication error. Corrective Action as it applies to others: The Vulnerable Adult Policy was reviewed and remains current. All medication errors have been reviewed that occurred in last 30 days to determine if medication error was reported as appropriate. Clinical leaders and Administrator will be educated on need to report appropriate medication errors to state agency in a		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245283	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 01/26/2023
NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792		
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F 609	Continued From page 2 insulin dependent. A Medication Error Reconciliation Form dated 1/20/23, indicated on 1/20/23, at 9:00 a.m. registered nurse (RN)-A gave R1 36 units (u) of Novolog insulin (a rapid-acting insulin that helps lower mealtime blood sugar spikes) and 90 u of Toujeo insulin (a long-acting insulin that helps control high blood sugar levels) of which was intended for another resident. The form indicated R1 received the wrong drug or dosage and an extra dosage was given. The outcome to R1 was to continue to monitor blood glucose levels every hour. The error was categorized as a category B error described as one that required medication discontinuation or dose modification or the need for treatment or intervention. The form indicated this was not a significant medication error with the rationale of R1's provider was notified as soon as the error was discovered. Appropriate interventions were initiated, R1 was monitored hourly and all insulins were held for the remainder of the day. The form's investigation summary indicated RN-A was still completing training. The error was made when RN-A did not ask R1 for her name. The medication error was reviewed with RN-A, and moving forward RN-A would ask who a resident is and ensure it was the correct resident prior to administering medications. The form further indicated RN-A reviewed the medication error policy and procedure. A progress note dated 1/20/23, at 9:30 a.m. indicated R1's blood glucose was 141. R1 received Novolog insulin 36 u and 90 u of Toujeo insulin that morning. The physician assistant (PA) was updated and received the following orders: 1. Check blood glucose every hour 2. Give 15 grams (gm) glucose gel now.	F 609	timely manner per facility policy. Date of Compliance: 2/1/23 Recurrence will be prevented by: All medication errors from last week will be reviewed to ensure that the medication error was reported as appropriate in a timely manner. This will occur weekly x4 weeks then monthly times 2 months. The results of the audits will be shared with the facility QAPI committee for input on the need to increase, decrease, or discontinue the audit. Corrections will be monitored by: Director of Nursing or Designee		

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

PRINTED: 02/27/2023
FORM APPROVED
OMB NO. 0938-0391

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NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETION DATE
F 609	Continued From page 3 3. Monitor for signs and symptoms of hypoglycemia. 4. If the bedtime blood glucose is less than 150, give 15 gm glucose gel. 5. If bedtime blood glucose is greater than 180, it was okay to decrease the blood glucose checks to every two to three hours. 6. Hold all insulin today. 7. Return to normal insulin usage on 1/21/23. During an interview on 1/26/23, at 8:34 a.m. the PA stated the significant medication error was a large error and should not have happened. The adverse effects to R1 could have been hypoglycemia (low blood glucose) leading to hospitalization and even death. During an interview on 1/26/23, at 10:36 a.m. the director of nursing (DON) stated the medication error was not significant and not reported to the SA. The DON did not feel the medication error was significant and did not report to the SA because the PA and physician were notified and orders were received. R1 was monitored for the rest of the day and her blood sugars were fine. The DON stated the medication error would have been reported to the SA if R1's blood sugars were dropping or if R1 was sent to the hospital. The facility's Abuse Prohibition/Vulnerable Adult policy dated 4/11/22, indicated directed suspicion of neglect, exploitation, or misappropriation of resident property must be reported to the SA not later than 2 hours if the incident resulted in serious bodily injury. If the suspected neglect, exploitation, or misappropriation of resident property did not result in serious bodily injury, the report must be made within 24 hours. The policy further included medication errors with adverse	F 609			

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245283	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 01/26/2023
NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792		
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F 609	Continued From page 4 effects or had the potential for adverse effects.	F 609			



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
February 2, 2023

Administrator
The Waterview Pines Llc
1201 8th Street South
Virginia, MN 55792

Re: State Nursing Home Licensing Orders
Event ID: LGOC11

Dear Administrator:

The above facility was surveyed on January 25, 2023 through January 26, 2023 for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules and Statutes. At the time of the survey, the survey team from the Minnesota Department of Health - Health Regulation Division noted one or more violations of these rules or statutes that are issued in accordance with Minn. Stat. § 144.653 and/or Minn. Stat. § 144A.10. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a civil fine for each deficiency not corrected shall be assessed in accordance with a schedule of fines promulgated by rule and/or statute of the Minnesota Department of Health.

To assist in complying with the correction order(s), a "suggested method of correction" has been added. This provision is being suggested as one method that you can follow to correct the cited deficiency. Please remember that this provision is only a suggestion and you are not required to follow it. Failure to follow the suggested method will not result in the issuance of a penalty assessment. You are reminded, however, that regardless of the method used, correction of the order within the established time frame is required. The "suggested method of correction" is for your information and assistance only.

You have agreed to participate in the electronic receipt of State licensure orders consistent with the Minnesota Department of Health Informational Bulletin 14-01, available at https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html. The State licensing orders are delineated on the Minnesota Department of Health State Form and are being delivered to you electronically. The Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Tag numbers have been assigned to Minnesota state statutes/rules for Nursing Homes.

The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule number and the corresponding text of the state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction order. This column also includes the findings that are in violation of the state statute or rule after the statement, "This MN Requirement is not met as evidenced by." Following the surveyors findings are the Suggested Method of Correction and the Time Period For Correction.

The Waterview Pines Llc

February 2, 2023

Page 2

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/RULES.

Although no plan of correction is necessary for State Statutes/Rules, please enter the word "corrected" in the box available for text. You must then indicate in the electronic State licensure process, under the heading completion date, the date your orders will be corrected prior to electronically submitting to the Minnesota Department of Health. We urge you to review these orders carefully, item by item, and if you find that any of the orders are not in accordance with your understanding at the time of the exit conference following the survey, you should immediately contact:

Terri Ament, Rapid Response
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Duluth Technology Village
11 East Superior Street, Suite 290
Duluth, Minnesota 55802-2007
Email: teresa.ament@state.mn.us
Office: (218) 302-6151 Mobile: (218) 766-2720

You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance.

Please feel free to call me with any questions.

Sincerely,



Joanne Simon, Compliance Analyst
Minnesota Department of Health
Health Regulation Division
Telephone: 651-201-4161
Email: joanne.simon@state.mn.us

cc: Licensing and Certification File

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00582	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 01/26/2023
NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792		
(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
2 000	Initial Comments *****ATTENTION***** NH LICENSING CORRECTION ORDER In accordance with Minnesota Statute, section 144A.10, this correction order has been issued pursuant to a survey. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a fine for each violation not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health. Determination of whether a violation has been corrected requires compliance with all requirements of the rule provided at the tag number and MN Rule number indicated below. When a rule contains several items, failure to comply with any of the items will be considered lack of compliance. Lack of compliance upon re-inspection with any item of multi-part rule will result in the assessment of a fine even if the item that was violated during the initial inspection was corrected. You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance. INITIAL COMMENTS: On 1/25/23, and 1/26/23, a complaint survey was conducted at your facility by surveyors from the Minnesota Department of Health (MDH). Your facility was found NOT in compliance with the MN State Licensure. The following complaint was found to be	2 000			

Minnesota Department of Health

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

Electronically Signed

TITLE

(X6) DATE

02/16/23

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00582	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 01/26/2023
NAME OF PROVIDER OR SUPPLIER THE WATERVIEW PINES LLC		STREET ADDRESS, CITY, STATE, ZIP CODE 1201 8TH STREET SOUTH VIRGINIA, MN 55792			
(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
2 000	Continued From page 1 SUBSTANTIATED: H52837840C (MN00090347) with no licensing orders were issued. However, an unrelated licensing order was issued at 626.557 Subd 3. The Minnesota Department of Health is documenting the State Licensing Correction Orders using Federal software.	2 000			
21980	MN St. Statute 626.557 Subd. 3 Reporting - Maltreatment of Vulnerable Adults Subd. 3. Timing of report. (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, 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.	21980			2/1/23

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21980	Continued From page 2 (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 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 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 agency information explaining how the event meets the criteria under section 626.5572, subdivision 17, paragraph (c), clause (5). The lead 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 document review, the facility failed to ensure significant medication errors (an error of which causes the resident discomfort or jeopardizes the resident's health and safety) were reported to the State Agency (SA) for 1 of 3 residents (R1) reviewed for significant medication errors. Findings include: R1's Admission Record dated 1/26/23, indicated R1's diagnoses included diabetes, long term insulin use, aphasia (loss of ability to understand or express speech, caused by brain damage) following a stroke, and dementia.	21980	Corrected		

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21980	Continued From page 3 R1's quarterly Minimum Data Set dated 1/23/23, indicated R1 was cognitively intact and was insulin dependent. A Medication Error Reconciliation Form dated 1/20/23, indicated on 1/20/23, at 9:00 a.m. registered nurse (RN)-A gave R1 36 units (u) of Novolog insulin (a rapid-acting insulin that helps lower mealtime blood sugar spikes) and 90 u of Toujeo insulin (a long-acting insulin that helps control high blood sugar levels) of which was intended for another resident. The form indicated R1 received the wrong drug or dosage and an extra dosage was given. The outcome to R1 was to continue to monitor blood glucose levels every hour. The error was categorized as a category B error described as one that required medication discontinuation or dose modification or the need for treatment or intervention. The form indicated this was not a significant medication error with the rationale of R1's provider was notified as soon as the error was discovered. Appropriate interventions were initiated, R1 was monitored hourly and all insulins were held for the remainder of the day. The form's investigation summary indicated RN-A was still completing training. The error was made when RN-A did not ask R1 for her name. The medication error was reviewed with RN-A, and moving forward RN-A would ask who a resident is and ensure it was the correct resident prior to administering medications. The form further indicated RN-A reviewed the medication error policy and procedure. A progress note dated 1/20/23, at 9:30 a.m. indicated R1's blood glucose was 141. R1 received Novolog insulin 36 u and 90 u of Toujeo insulin that morning. The physician assistant (PA) was updated and received the following orders: 1. Check blood glucose every hour	21980			

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21980	Continued From page 4 2. Give 15 grams (gm) glucose gel now. 3. Monitor for signs and symptoms of hypoglycemia. 4. If the bedtime blood glucose is less than 150, give 15 gm glucose gel. 5. If bedtime blood glucose is greater than 180, it was okay to decrease the blood glucose checks to every two to three hours. 6. Hold all insulin today. 7. Return to normal insulin usage on 1/21/23. During an interview on 1/26/23, at 8:34 a.m. the PA stated the significant medication error was a large error and should not have happened. The adverse effects to R1 could have been hypoglycemia (low blood glucose) leading to hospitalization and even death. During an interview on 1/26/23, at 10:36 a.m. the director of nursing (DON) stated the medication error was not significant and not reported to the SA. The DON did not feel the medication error was significant and did not report to the SA because the PA and physician were notified and orders were received. R1 was monitored for the rest of the day and her blood sugars were fine. The DON stated the medication error would have been reported to the SA if R1's blood sugars were dropping or if R1 was sent to the hospital. The facility's Abuse Prohibition/Vulnerable Adult policy dated 4/11/22, indicated directed suspicion of neglect, exploitation, or misappropriation of resident property must be reported to the SA not later than 2 hours if the incident resulted in serious bodily injury. If the suspected neglect, exploitation, or misappropriation of resident property did not result in serious bodily injury, the report must be made within 24 hours. The policy further included medication errors with adverse	21980			

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21980	Continued From page 5 effects or had the potential for adverse effects. SUGGESTED METHOD OF CORRECTION: The administrator or designee could develop/revise policies or procedures to ensure timely reporting of all allegations of significant medication errors are within appropriate timeframes for reporting. The facility could re-educate staff to policies and procedures, and audit all complaints of alleged abuse or neglect in a measurable and specific way. The results of those audits should be taken to the Quality Assurance Performance Improvement (QAPI) committee to determine the need for further monitoring or compliance. Those audits should be ongoing and random after compliance is determined by QAPI to ensure compliance is being maintained. TIME PERIOD FOR CORRECTION: Twenty-one (21) days.	21980			