



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
November 3, 2020

CMS Certification Number (CCN): 245286

Administrator
Pierz Villa Inc
119 Faust Street Southeast
Pierz, MN 56364

Dear Administrator:

The Minnesota Department of Health assists the Centers for Medicare and Medicaid Services (CMS) by surveying skilled nursing facilities and nursing facilities to determine whether they meet the requirements for participation. To participate as a skilled nursing facility in the Medicare program or as a nursing facility in the Medicaid program, a provider must be in substantial compliance with each of the requirements established by the Secretary of Health and Human Services found in 42 CFR part 483, Subpart B.

Based upon your facility being in substantial compliance, we are recommending to CMS that your facility be recertified for participation in the Medicare and Medicaid program.

Effective October 15, 2020 the above facility is certified for:

50 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 50 skilled nursing facility beds.

You should advise our office of any changes in staffing, services, or organization, which might affect your certification status. If, at the time of your next survey, we find your facility to not be in substantial compliance your Medicare and/or Medicaid provider agreement may be subject to non-renewal or termination.

Please contact me if you have any questions.

Sincerely,

A handwritten signature in black ink, appearing to read 'Douglas Larson'.

Douglas Larson, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program
Program Assurance Unit
Health Regulation Division
Telephone: 651-201-4118 Fax: 651-215-9697
Email: doug.larson@state.mn.us

cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

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November 3, 2020

Administrator
Pierz Villa Inc
119 Faust Street Southeast
Pierz, MN 56364

RE: CCN: 245286
Cycle Start Date: August 18, 2020

Dear Administrator:

On September 14, 2020, we notified you a remedy was imposed. On October 23, 2020 the Minnesota Department(s) 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 October 15, 2020.

As authorized by CMS the remedy of:

- Mandatory denial of payment for new Medicare and Medicaid admissions effective November 20, 2020 did not go into effect. (42 CFR 488.417 (b))

In our letter of September 14, 2020, 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 November 20, 2020 due to denial of payment for new admissions. Since your facility attained substantial compliance on October 15, 2020, 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,

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Douglas Larson, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program

Pierz Villa Inc

November 3, 2020

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Program Assurance Unit

Health Regulation Division

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

ID: EQ9S

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

Facility ID: 00384

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245286		3. NAME AND ADDRESS OF FACILITY (L3) PIERZ VILLA INC (L4) 119 FAUST STREET SOUTHEAST (L5) PIERZ, MN (L6) 56364		4. TYPE OF ACTION: <u>7</u> (L8) 1. Initial 2. Recertification 3. Termination 4. CHOW 5. Validation 6. Complaint 7. On-Site Visit 9. Other 8. Full Survey After Complaint	
2.STATE VENDOR OR MEDICAID NO. (L2) 964657400		5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9) 01/01/2009		7. PROVIDER/SUPPLIER CATEGORY <u>02</u> (L7) 01 Hospital 05 HHA 09 ESRD 13 PTIP 22 CLIA 02 SNF/NF/Dual 06 PRTF 10 NF 14 CORF 03 SNF/NF/Distinct 07 X-Ray 11 ICF/IID 15 ASC 04 SNF 08 OPT/SP 12 RHC 16 HOSPICE	
6. DATE OF SURVEY 10/23/2020 (L34)		8. ACCREDITATION STATUS: <u> </u> (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other		FISCAL YEAR ENDING DATE: (L35) 12/31	
11. LTC PERIOD OF CERTIFICATION From (a) : To (b) :		10.THE FACILITY IS CERTIFIED AS: X A. In Compliance With <u>And/Or Approved Waivers Of The Following Requirements:</u> Program Requirements <u> </u> 2. Technical Personnel <u> </u> 6. Scope of Services Limit Compliance Based On: <u> </u> 3. 24 Hour RN <u> </u> 7. Medical Director <u> </u> 1. Acceptable POC <u> </u> 4. 7-Day RN (Rural SNF) <u> </u> 8. Patient Room Size <u> </u> 5. Life Safety Code <u> </u> 9. Beds/Room B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: A (L12)			
12.Total Facility Beds 50 (L18)		13.Total Certified Beds 50 (L17)		14. LTC CERTIFIED BED BREAKDOWN 18 SNF 18/19 SNF 19 SNF ICF IID 50 (L37) (L38) (L39) (L42) (L43)	
		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)			

16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):

17. SURVEYOR SIGNATURE <u>Kathleen Lucas, Unit Supervisor</u>	Date : 11/03/2020 (L19)	18. STATE SURVEY AGENCY APPROVAL <u>Douglas Larson, Enforcement Specialist</u>	Date: 11/03/2020 (L20)
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PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY

19. DETERMINATION OF ELIGIBILITY <u>X</u> 1. Facility is Eligible to Participate <u> </u> 2. Facility is not Eligible (L21)		20. COMPLIANCE WITH CIVIL RIGHTS ACT:		21. 1. Statement of Financial Solvency (HCFA-2572) 2. Ownership/Control Interest Disclosure Stmt (HCFA-1513) 3. Both of the Above : <u> </u>	
22. ORIGINAL DATE OF PARTICIPATION 08/01/1985 (L24)	23. LTC AGREEMENT BEGINNING DATE (L41)	24. LTC AGREEMENT ENDING DATE (L25)	26. TERMINATION ACTION: (L30) <u>VOLUNTARY</u> <u>00</u> <u>INVOLUNTARY</u> 01-Merger, Closure 05-Fail to Meet Health/Safety 02-Dissatisfaction W/ Reimbursement 06-Fail to Meet Agreement 03-Risk of Involuntary Termination <u>OTHER</u> 04-Other Reason for Withdrawal 07-Provider Status Change 00-Active		
25. LTC EXTENSION DATE: (L27)	27. ALTERNATIVE SANCTIONS A. Suspension of Admissions: (L44) B. Rescind Suspension Date: (L45)				
28. TERMINATION DATE: (L28)		29. INTERMEDIARY/CARRIER NO. 03001 (L31)		30. REMARKS	
31. RO RECEIPT OF CMS-1539 (L32)		32. DETERMINATION OF APPROVAL DATE 10/21/2020 (L33)		DETERMINATION APPROVAL	



Protecting, Maintaining and Improving the Health of All Minnesotans

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Administrator
Pierz Villa Inc
119 Faust Street Southeast
Pierz, MN 56364

RE: CCN: 245286
Cycle Start Date: August 20, 2020

Dear Administrator:

On September 14, 2020, we informed you of imposed enforcement remedies.

- Mandatory Denial of Payment for new Medicare and/or Medicaid admissions effective November 20, 2020. (42 CFR 488.417 (b))

On September 3, 2020, the Minnesota Department(s) of Health completed a survey and it has been determined that your facility continues to not to be in substantial compliance. The most serious deficiencies in your facility were found to be isolated deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level D), as evidenced by the electronically attached CMS-2567, whereby corrections are required.

As a result of the survey findings:

- Mandatory Denial of Payment for new Medicare and/or Medicaid Admissions, Federal regulations at 42 CFR § 488.417(b), effective November 20, 2020, will remain in effect.

This Department continues to recommend that CMS impose a civil money penalty. (42 CFR 488.430 through 488.444).

You will receive a formal notice from the CMS RO only if CMS agrees with our recommendation.

The CMS Region V Office will notify your Medicare Administrative Contractor (MAC) that the denial of payment for new admissions is effective November 20, 2020. They will also notify the State Medicaid Agency that they must also deny payment for new Medicaid admissions effective November 20, 2020.

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.

As we notified you in our letter of [First State Notice Date()], in accordance with Federal law, as specified in the Act at Section 1819(f)(2)(B)(iii)(I)(b) and 1919(f)(2)(B)(iii)(I)(b), your facility is prohibited from conducting Nursing Aide Training and/or Competency Evaluation Programs (NATCEP) for two years from November 20, 2020.

ELECTRONIC PLAN OF CORRECTION (ePOC)

Within ten (10) calendar days after your receipt of this notice, you must submit an acceptable plan of correction (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.

If an acceptable ePoC is not received within 10 calendar days from the receipt of this letter, we will recommend to the CMS Region V Office that one or more of the following remedies be imposed:

- Optional denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417 (a));
- Per day civil money penalty (42 CFR 488.430 through 488.444).

DEPARTMENT CONTACT

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

**Kathleen Lucas, Unit Supervisor
St. Cloud B Survey Team
Licensing and Certification Program**

Health Regulation Division
Minnesota Department of Health
Midtown Square
3333 Division Street, Suite 212
Saint Cloud, Minnesota 56301-4557
Email: kathleen.lucas@state.mn.us
Phone: (320) 223-7343
Fax: (320) 223-7348

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 [Cycle Start + 6 Months()] (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 Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

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

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:

Tamika.Brown@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 Tamika Brown, Principal Program Representative by phone at (312) 353-1502 or by e-mail at Tamika.Brown@cms.hhs.gov.

INFORMAL DISPUTE RESOLUTION/ INDEPENDENT INFORMAL DISPUTE RESOLUTION (IIDR)

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

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

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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/ltc_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 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 "Douglas Larson", with a long horizontal flourish extending to the right.

Douglas Larson, Enforcement Specialist
Minnesota Department of Health
Licensing and Certification Program
Program Assurance Unit
Health Regulation Division
Telephone: 651-201-4118 Fax: 651-215-9697
Email: doug.larson@state.mn.us

cc: Licensing and Certification File

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

PRINTED: 10/05/2020
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245286	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 09/03/2020
NAME OF PROVIDER OR SUPPLIER PIERZ VILLA INC			STREET ADDRESS, CITY, STATE, ZIP CODE 119 FAUST STREET SOUTHEAST PIERZ, MN 56364		
(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 9/2/20 through 9/3/20, an abbreviated survey was completed at your facility to conduct complaint investigations. Pierz Villa was found not to be in compliance with 42 CFR Part 483, Requirements for Long Term Care Facilities. The following complaints were found to be SUBSTANTIATED: H5286032C, H5286033C, H5286034C. No deficiencies were issued related to the complaints, due to actions implemented by the facility prior to the abbreviated survey. However, as a result of the investigation, other deficiencies were identified. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of the CMS-2567 form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.	F 000			
F 761 SS=D	Label/Store Drugs and Biologicals CFR(s): 483.45(g)(h)(1)(2) §483.45(g) Labeling of Drugs and Biologicals Drugs and biologicals used in the facility must be labeled in accordance with currently accepted professional principles, and include the appropriate accessory and cautionary instructions, and the expiration date when applicable. §483.45(h) Storage of Drugs and Biologicals §483.45(h)(1) In accordance with State and Federal laws, the facility must store all drugs and biologicals in locked compartments under proper	F 761			10/15/20

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

TITLE

(X6) DATE

Electronically Signed

09/24/2020

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 761	Continued From page 1 temperature controls, and permit only authorized personnel to have access to the keys. §483.45(h)(2) The facility must provide separately locked, permanently affixed compartments for storage of controlled drugs listed in Schedule II of the Comprehensive Drug Abuse Prevention and Control Act of 1976 and other drugs subject to abuse, except when the facility uses single unit package drug distribution systems in which the quantity stored is minimal and a missing dose can be readily detected. This REQUIREMENT is not met as evidenced by: Based on observation, interview, and document review, the facility failed to ensure insulin was given per manufacturer's instructions for 1 of 5 residents (R2), reviewed for medication storage. Findings include: R2's Face Sheet with a print date of 9/3/20, included diagnoses bacteremia, depressive episodes, nutritional deficiency, and type 2 diabetes mellitus without complications. R2's annual Care Area Assessment (CAA) dated 2/7/20, indicated R2 was dependent with activities of daily living (ADLs) including bed mobility, transfers, toileting, bathing, and dressing. R2's physician's orders, printed date of 9/3/20, directed staff to administer Novolog Flexpen U-100 Insulin (short acting insulin), according to the sliding scale provided, subcutaneously (under the skin), three times a day. During interview on 9/2/20, at 2:50 p.m. licensed practical nurse (LPN)-A was reading insulin vials and pens on the south medication cart to verify	F 761	Pierz Villa strives to ensure drugs and biologicals use in our facility are labeled in accordance with currently accepted professional principles and include the appropriate accessory and cautionary instructions and expiration when applicable. On 8/24/2020 Pierz Villa implemented labels with open date/expiration date to be placed on medications with shortened expiration dates. On 9/2/2020 R2's expired insulin was disposed of and new insulin was retrieved out of refrigerator and marked with open date and expiration date. On 9/3/2020 DON checked medications cart for open/expiration dates on insulin vials and pens with 100% compliance found. The policy and procedure for medication labeling was reviewed and revised on 9/17/2020 to include the list of Medication Expiration after Opening. On 9/17/2020 the Medication Expiration after Opening list was placed on the north and south		

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F 761	Continued From page 2 they matched the orders for each resident receiving insulin. LPN-A grabbed R2's Novolog Flexpen and read the open date and expiration date that was hand written on the insulin pen and verified that the handwritten open date was 7/28/20, and expiration date was 8/30/20, indicating the Novolog Flexpen had expired three days prior. R2's Insulin Administration History record from 8/1/20 to 8/31/20 indicated R2 received two units of Novolog on 8/31/20 and 9/1/20 during the 12:00 p.m. medication pass. During phone interview on 9/3/20, at 1:53 p.m. LPN-A indicated she disposed of the expired insulin after discovery. LPN-A confirmed that she gave R2 the two units of insulin on 8/31/20 and 9/1/20 from the Novolog insulin pen with the expiration date of 8/30/20. Further, LPN-A stated "It gets extremely busy and it's easy to miss." During interview on 9/3/20, at 2:35 p.m. director of nursing (DON) confirmed she was informed of medication error on 9/2/20 after the discovery of expired insulin. DON stated that her expectation is the nurses administering medication should be checking the open and expiration dates on the vial before administering insulin. DON stated that the trained medication assistants (TMAs) go through each medication cart and look at the expiration dates and give to the licensed nurse to dispose of them. Further, the DON indicated that there is no policy specifically for insulin administration but expects the nurse to follow the Medication Administration policy. Review of the Medications with Shortened Expiration Dates chart, at the south nursing	F 761	medication carts and nurse stations for referencing. On 9/17/2020 DON again checked resident insulins for open dates and expiration dates with 100% compliance found. RNs/LPNs and TMA's were educated via email on 9/18/2020. On 9/18/2020 a system was established to check medication carts weekly for expiration dates. Further education will be provided to the Pierz Villa Nursing Team on 10/5/2020; 10/7/2020; 10/8/2020; 10/11/2020; 10/12/2020. Audits will be conducted on R2 and all resident with insulin pens or vials 1 time weekly for 1 month; then every other week for 1 month, then 1 time a month for 1 month or until resolved. Results of audits will be reviewed and discussed with QA committee. Pierz Villa will be in compliance on 10/15/2020.		

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F 761	Continued From page 3 station, indicated Novolog expired 28 days after first use. Review of the Medication Administration policy, dated 2/13, directed staff to administer medications to the resident in accordance with the "Five Rights", however it did not direct staff on how to administer insulin according to manufacture's instructions. In addition, the policy did not address shortened expiration dates for insulin.	F 761			



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered
September 18, 2020

Administrator
Pierz Villa Inc
119 Faust Street Southeast
Pierz, MN 56364

Re: State Nursing Home Licensing Orders
Event ID: OKWH11

Dear Administrator:

The above facility was surveyed on September 2, 2020 through September 3, 2020 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

Page 2

"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.

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:

Kathleen Lucas, Unit Supervisor
St. Cloud B Survey Team
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Midtown Square
3333 Division Street, Suite 212
Saint Cloud, Minnesota 56301-4557
Email: kathleen.lucas@state.mn.us
Phone: (320) 223-7343
Fax: (320) 223-7348

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 note it is your responsibility to share the information contained in this letter and the results of this visit with the President of your facility's Governing Body.

Please feel free to call me with any questions.

Sincerely,



Douglas Larson, Enforcement Specialist
Minnesota Department of Health

Pierz Villa Inc

Page 3

Licensing and Certification Program

Program Assurance Unit

Health Regulation Division

Telephone: 651-201-4118 Fax: 651-215-9697

Email: doug.larson@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: 00384	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED C 09/03/2020
NAME OF PROVIDER OR SUPPLIER PIERZ VILLA INC		STREET ADDRESS, CITY, STATE, ZIP CODE 119 FAUST STREET SOUTHEAST PIERZ, MN 56364		
(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 9/2/20 through 9/3/20, an abbreviated survey was conducted at Pierz Villa to determine compliance with State Licensure. Your facility was found not to be compliance with the MN State Licensure. The following complaints were found to be	2 000		

Minnesota Department of Health

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

TITLE

(X6) DATE

Electronically Signed

09/24/20

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00384	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED C 09/03/2020
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2 000	Continued From page 1 SUBSTANTIATED: H5350111C, H5350116C, H5350107C. NO licensing orders were issued related to the complaints However, as a result of the investigation other orders were identified. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of state form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.	2 000			
21620	MN Rule 4658.1345 Labeling of Drugs Drugs used in the nursing home must be labeled in accordance with part 6800.6300. This MN Requirement is not met as evidenced by: Based on observation, interview, and document review, the facility failed to ensure insulin was given per manufacturer's instructions for 1 of 5 residents (R2), reviewed for medication storage. Findings include: R2's Face Sheet with a print date of 9/3/20, included diagnoses bacteremia, depressive episodes, nutritional deficiency, and type 2 diabetes mellitus without complications. R2's annual Care Area Assessment (CAA) dated 2/7/20, indicated R2 was dependent with activities of daily living (ADLs) including bed mobility, transfers, toileting, bathing, and dressing.	21620	Corrected		10/15/20

Minnesota Department of Health

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21620	Continued From page 2 R2's physician's orders, printed date of 9/3/20, directed staff to administer Novolog Flexpen U-100 Insulin (short acting insulin), according to the sliding scale provided, subcutaneously (under the skin), three times a day. During interview on 9/2/20, at 2:50 p.m. licensed practical nurse (LPN)-A was reading insulin vials and pens on the south medication cart to verify they matched the orders for each resident receiving insulin. LPN-A grabbed R2's Novolog Flexpen and read the open date and expiration date that was hand written on the insulin pen and verified that the handwritten open date was 7/28/20, and expiration date was 8/30/20, indicating the Novolog Flexpen had expired three days prior. R2's Insulin Administration History record from 8/1/20 to 8/31/20 indicated R2 received two units of Novolog on 8/31/20 and 9/1/20 during the 12:00 p.m. medication pass. During phone interview on 9/3/30, at 1:53 p.m. LPN-A indicated she disposed of the expired insulin after discovery. LPN-A confirmed that she gave R2 the two units of insulin on 8/31/20 and 9/1/20 from the Novolog insulin pen with the expiration date of 8/30/20. Further, LPN-A stated "It gets extremely busy and it's easy to miss." During interview on 9/3/20, at 2:35 p.m. director of nursing (DON) confirmed she was informed of medication error on 9/2/20 after the discovery of expired insulin. DON stated that her expectation is the nurses administering medication should be checking the open and expiration dates on the vial before administering insulin. DON stated that the trained medication assistants (TMAs) go through each medication cart and look at the	21620			

Minnesota Department of Health

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21620	Continued From page 3 expiration dates and give to the licensed nurse to dispose of them. Further, the DON indicated that there is no policy specifically for insulin administration but expects the nurse to follow the Medication Administration policy. Review of the Medications with Shortened Expiration Dates chart, at the south nursing station, indicated Novolog expired 28 days after first use. Review of the Medication Administration policy, dated 2/13, directed staff to administer medications to the resident in accordance with the "Five Rights", however it did not direct staff on how to administer insulin according to manufacture's instructions. In addition, the policy did not address shortened expiration dates for insulin. SUGGESTED METHOD OF CORRECTION: The director of nursing (DON) and consulting pharmacist could review and revise policies and procedures for proper storage and disposition of medications. Nursing staff could be educated as necessary to the importance of labeling, auditing medications for expiration and removing any expired medications from the supply. The DON or designee, could audit medications on a regular basis to ensure compliance. TIME PERIOD FOR CORRECTION: Twenty one (21) days.	21620			