



*Protecting, Maintaining and Improving the Health of All Minnesotans*

Electronically Delivered  
May 1, 2025

Administrator  
The Villas At Roseville  
1000 Lovell Avenue  
Roseville, MN 55113

RE: CCN: 245326  
Cycle Start Date: March 12, 2025

Dear Administrator:

On April 28, 2025, the Minnesota Departments of Health completed a revisit to verify that your facility had achieved and maintained compliance. Based on our review, we have determined that your facility has achieved substantial compliance; therefore no remedies will be imposed.

Feel free to contact me if you have questions.

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

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



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Electronically delivered  
March 20, 2025

Administrator  
The Villas At Roseville  
1000 Lovell Avenue  
Roseville, MN 55113

RE: CCN: 245326  
Cycle Start Date: March 12, 2025

Dear Administrator:

On March 12, 2025, a survey was completed at your facility by the Minnesota Departments of Health and Public Safety, to determine if your facility was in compliance with Federal participation requirements for skilled nursing facilities and/or nursing facilities participating in the Medicare and/or Medicaid programs.

This survey found the most serious deficiencies in your facility to be isolated deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level D), as evidenced by the electronically attached CMS-2567 whereby corrections are required.

#### **ELECTRONIC PLAN OF CORRECTION (ePoC)**

Within **ten (10) calendar days** after your receipt of this notice, you must submit an acceptable ePOC for the deficiencies cited. An acceptable ePOC will serve as your allegation of compliance. Upon receipt of an acceptable ePOC, we will authorize a revisit to your facility to determine if substantial compliance has been achieved.

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

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

The Villas At Roseville

March 20, 2025

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The state agency may, in lieu of an onsite revisit, determine correction and compliance by accepting the facility's ePoC if the ePoC is reasonable, addresses the problem and provides evidence that the corrective action has occurred.

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

- Denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417);
- Civil money penalty (42 CFR 488.430 through 488.444).
- Termination of your facility's Medicare and/or Medicaid agreement (488.456(b)).

#### DEPARTMENT CONTACT

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

Renee McClellan, Regional Operations Supervisor  
Metro A District Office  
Health Regulation Division  
Minnesota Department of Health  
625 Robert Street N  
P.O. Box 64975  
Saint Paul, Minnesota 55164-0975  
Email: renee.mcclellan@state.mn.us  
Office: 651-201-4391 Mobile: 651-328-9282

#### PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

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

#### VERIFICATION OF SUBSTANTIAL COMPLIANCE

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

If substantial compliance has been achieved, certification of your facility in the Medicare and/or Medicaid program(s) will be continued and remedies will not be imposed. Compliance is certified as of

the latest correction date on the approved ePoC, unless it is determined that either correction actually occurred between the latest correction date on the ePoC and the date of the first revisit, or correction occurred sooner than the latest correction date on the ePoC.

#### **FAILURE TO ACHIEVE SUBSTANTIAL COMPLIANCE BY THE THIRD OR SIXTH MONTH AFTER THE LAST DAY OF THE SURVEY**

If substantial compliance with the regulations is not verified by June 12, 2025 (three months after the identification of noncompliance), the CMS Region V Office must deny payment for new admissions as mandated by the Social Security Act (the Act) at Sections 1819(h)(2)(D) and 1919(h)(2)(C) and Federal regulations at 42 CFR Section 488.417(b).

In addition, if substantial compliance with the regulations is not verified by September 12, 2025 (six months after the identification of noncompliance) your provider agreement will be terminated. This action is mandated by the Social Security Act at Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

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

#### **INFORMAL DISPUTE RESOLUTION (IDR)**

In accordance with 42 CFR 488.331 and Minnesota Statute 144A.10 subd 15, 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: <https://forms.web.health.state.mn.us/form/NHDisputeResolution>

This request must be sent within the same ten calendar days you have for submitting an ePoC for the cited deficiencies. 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.

A copy of the Department's informal dispute resolution policies is posted on the MDH Information Bulletin website at: [https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04\\_8.html](https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html)

#### **INDEPENDENT INFORMAL DISPUTE RESOLUTION (INDEPENDENT IDR)**

In accordance with 42 CFR § 488.431 and Minnesota Statute 144A.10 subd 16, when a CMP subject to being collected and placed in an escrow account is imposed, you have one opportunity to question cited deficiencies through an Independent IDR process. You may also contest scope and severity assessments for deficiencies which resulted in a finding of SQC or immediate jeopardy. 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: <https://forms.web.health.state.mn.us/form/NHDisputeResolution>

The Villas At Roseville

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A facility may not use both IDR and independent IDR for the same deficiency citation(s) arising from the same survey unless the IDR process was completed prior to the imposition of the CMP. This request must be sent within ten calendar days of receipt of this offer. An incomplete Independent IDR process will not delay the effective date of any enforcement action.

Questions regarding all documents submitted as a response to the Life Safety Code deficiencies (those preceded by a "K" tag), i.e., the plan of correction, request for waivers, should be directed to:

Travis Z. Ahrens  
State Fire Safety Supervisor  
Health Care & Correctional Facilities  
MN Department of Public Safety-Fire Marshal Division  
445 Minnesota St., Suite 145  
St. Paul, MN 55101  
Email: [travis.ahrens@state.mn.us](mailto:travis.ahrens@state.mn.us)  
Web: [www.sfm.dps.mn.gov](http://www.sfm.dps.mn.gov)  
Cell: 1-507-308-4189

Feel free to contact me if you have questions.

Sincerely,



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

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

PRINTED: 04/01/2025  
FORM APPROVED  
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:  <b>245326</b>	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____  B. WING _____	(X3) DATE SURVEY COMPLETED  <b>C</b> <b>03/12/2025</b>
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NAME OF PROVIDER OR SUPPLIER  <b>THE VILLAS AT ROSEVILLE</b>	STREET ADDRESS, CITY, STATE, ZIP CODE <b>1000 LOVELL AVENUE ROSEVILLE, MN 55113</b>
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(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETION DATE
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E 000	Initial Comments  On 3/10/25-3/12/25, a survey for compliance with §483.73, Appendix Z, Emergency Preparedness Requirements for Long Term Care Facilities was conducted during a standard recertification survey. The facility was in compliance.	E 000		
F 000	INITIAL COMMENTS  On 3/10/25-3/12/25, a standard recertification survey was conducted at your facility. A complaint investigation was also conducted. Your facility was not in compliance with the requirements of 42 CFR 483, Subpart B, Requirements for Long Term Care Facilities.  The following complaints were reviewed with no deficiencies cited: H53269402C (MN00110873) H53269461C (MN00110821) H53269423C (MN00110628)) H53269421C (MN00110184) H53269422C (MN00108301) H53269424C (MN00106927) H53269425C (MN00106850) H53269426C (MN00103891) H53269427C (MN00100217)  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	F 000		

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE  <b>Electronically Signed</b>	TITLE	(X6) DATE <b>03/29/2025</b>
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Any deficiency statement ending with an asterisk (\*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

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F 000	Continued From page 1 form. Your electronic submission of the POC will be used as verification of compliance.	F 000			
F 641 SS=D	<p>Accuracy of Assessments CFR(s): 483.20(g)</p> <p>§483.20(g) Accuracy of Assessments. The assessment must accurately reflect the resident's status. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to accurately code the Minimum Data Set (MDS) for 2 of 2 residents (R40) reviewed for pressure ulcers, and (R15) reviewed for psychotropic medications.</p> <p>Findings include:</p> <p>The Centers for Medicare and Medicaid (CMS) Long-Term Care Facility Resident Assessment Instrument (RAI) 3.0 user's manual dated 10/2024, identified the intent of section M, Skin Conditions, documented the risk, presence, appearance, and change of pressure ulcers and injuries. Further, it was important to recognize and evaluate each resident's risk factors and identify and evaluate all areas at risk of constant pressure and a complete assessment of skin was essential to an effective pressure ulcer prevention and skin treatment program. The RAI manual directed staff to review the medical record including skin care flow sheets, or other skin tracking forms, nurses' notes, and pressure ulcer</p>	F 641	<p>A modified MDS was submitted for R40 on 3/26/25 and R15 on 3/12/25. R40's MDS was modified to correct the pressure ulcer staging, and R15's MDS was modified to accurately document anti-anxiety medication use.</p> <p>Residents with pressure ulcers and residents taking anxiolytic medication have the potential to be affected. All residents with pressure ulcers and residents taking anxiolytic medications will have their most recent MDS reviewed to ensure accurate document.</p> <p>MDS coordinator will be educated on pressure ulcer staging and psychotropic medication classification, as well as how to accurately identify and code them on the MDS.</p> <p>Audits will be done on quarterly, annual, and significant change MDS assessments</p>	4/16/25	

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F 641	<p>Continued From page 2</p> <p>injury risk assessments, speak with the treatment nurse and direct care staff on all shifts to confirm conclusions from the medical record review and observations of the resident, examine the resident and determine whether any ulcers, injuries, scars, or non-removable dressings, devices were present. Additionally, pressure ulcer staging was determined by the deepest anatomical stage and if a stageable pressure ulcer had been classified at a higher numerical stage than what was observed it should continue to be classified at the higher numerical stage and pressure ulcers do not heal in a reverse sequence. Stage three and four pressure ulcers fill with granulation tissue that is never as strong as the tissue that was lost and hence is more prone to future breakdown. Clinical standards do not support reverse staging or back staging as a way to document healing as it does not accurately characterize what is occurring physiologically as the ulcer heals. Once a pressure ulcer has healed, it is documented as a healed pressure ulcer at its highest numerical stage and a previously closed pressure ulcer that opens again should be reported at its worst stage, unless currently presenting at a higher stage or unstageable.</p> <p>The State Operations Manual (SOM) provided guidance on pressure ulcer staging that defined the following pressure ulcers/injuries: Stage 1 Pressure Injury is intact skin with localized redness. Stage 2 Pressure ulcer is partial-thickness loss of skin with exposed dermis, presenting as a shallow open ulcer. The wound bed is viable, pink or red, moist, and may also present as an intact or open or ruptured blister. Adipose (fat) is not visible and deeper tissues are not visible.</p>	F 641	to ensure pressure ulcers and psychotropic medications are accurately documented. DON or designees will conduct audits on 5 MDS assessments each week x2 weeks, then 3 MDS assessments each week x2 weeks, then 1 MDS assessment each week x2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.	

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F 641	<p>Continued From page 3</p> <p>Granulation tissue (new connective tissue), slough, (non-viable tissue) and eschar (dead tissue) are not present.</p> <p>Stage 3 Pressure Ulcer is a full thickness loss of skin in which subcutaneous fat may be visible in the ulcer and granulation tissue and epibole (rolled wound edges) are often present. Slough and or eschar may be visible but does not obscure the depth of tissue loss. If slough or eschar obscures the wound bed, it is an unstageable pressure ulcer/injury.</p> <p>Stage 4 Pressure Ulcer is a full thickness skin and tissue loss with exposed or directly palpable fascia, muscle, tendon, ligament, cartilage or bone in the ulcer. Slough and or eschar may be visible on some parts of the wound bed. Epibole (rolled edges), undermining and or tunneling often occur. Depth varies by anatomical location. If slough or eschar obscures the wound bed, it is an unstageable pressure ulcer/pressure injury.</p> <p>Unstageable Pressure Ulcer is a full thickness skin and tissue loss in which the extent of tissue damage within the ulcer cannot be confirmed because the wound bed is obscured by slough or eschar. If the slough or eschar is removed, a stage 3 or stage 4 pressure ulcer will be revealed. If the anatomical depth of the tissue damage involved can be determined, then the reclassified stage should be assigned.</p> <p>Deep Tissue Pressure Injury (DTPI) is intact skin with a localized area of persistent non-blanchable deep red, maroon, purple discoloration due to damage of underlying soft tissue. This injury results from intense and or prolonged pressure and shear forces at the bone-muscle interface. The wound may evolve rapidly to reveal the actual extent of tissue injury, or may resolve without tissue loss. Once a deep tissue injury opens to an ulcer, reclassify the ulcer into the</p>	F 641		

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F 641	<p>Continued From page 4 appropriate stage.</p> <p>R40's annual MDS dated 1/3/25, indicated R40 had one or more unhealed pressure ulcers/injuries, 0 stage 1 pressure injuries, one stage 2 pressure ulcer/injury, 0 stage 3 pressure ulcers, 0 stage 4 pressure ulcers, 0 unstageable pressure ulcers with slough or eschar, and 0 unstageable DTPI's. The MDS directed staff to report based on the highest stage of existing ulcers/injuries at their worst; and do not "reverse" stage.</p> <p>R40's progress notes dated 1/1/25, indicated, R40's "pressure ulcer 2 right Achilles-improving".</p> <p>R40's Skin and Wound Evaluation form dated 12/26/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 2.9 centimeters (CM) long by 2.1 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 70% granulation tissue, 10% slough, and 20% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 1/1/25, indicated R40 had an in-house acquired stage two pressure ulcer to the right Achilles that had been present for one week measuring 4.1 (cm) long by 2.7 cm wide. Further, the note indicated the wound bed contained 80% granulation tissue, 20% slough, and no eschar.</p> <p>R40's Integrated Wound Care note dated 12/26/24, indicated R40 had a right heel unstageable pressure ulcer. Further, the wound was 2 cm long by 4.7 cm wide, by 0.4 cm deep and contained 10% slough, 20% necrotic tissue, and 70% granulation tissue.</p>	F 641		

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F 641	<p>Continued From page 5</p> <p>R40's Integrated Wound Care note dated 1/2/24, indicated R40 had a right heel unstageable pressure ulcer. Further, the wound was 2.5 cm long, by 4 cm wide by 0.4 cm deep and contained 20% slough, and 80% granulation.</p> <p>During interview on 3/10/25 at 5:18 p.m., R40 stated her heel was painful and pointed to her right heel and stated that was why it was elevated. R40's wheelchair foot rest on the right was pulled down and R40 had a sock on with her right heel resting directly on the foot rest. R40 stated she had a sore on her heel and stated staff usually changed the dressing in the morning, but stated it was changed mid morning on 3/10/25.</p> <p>During interview on 3/12/25 at 12:14 p.m., registered nurse (RN)-A stated he was the MDS coordinator and scheduled the MDS assessments, completed assessments, and made sure everyone completed their sections. RN-A stated he completed sections A, E, G, H, GG, I, J, K, L, M, N, O, P, and S. RN-A stated they were supposed to schedule skin assessments each week and when looking for skin assessments, looked at the Forms in the electronic medical record (EMR). RN-A stated R40 had an Annual MDS on 1/3/25, and when completing the MDS, looked for a skin assessment for 1/3/25, or around that date. RN-A stated they used to have a wound care manager who documented notes directly into progress notes but did not locate notes related to staging during the window for the MDS after reviewing the progress notes. RN-A stated he could not recall where he went to find the stage of the wound and knew it had been a stage two pressure ulcer and stated he used the RAI</p>	F 641		

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F 641	<p>Continued From page 6</p> <p>manual for guidance. RN-A stated according to the RAI manual, a stage two pressure ulcer was the first layer of skin removed and could also be a blister. RN-A further stated a stage two pressure ulcer did not contain slough, eschar, or granulation tissue. RN-A stated he viewed the Wound Evaluation form 1/1/25, for the MDS and stated the form indicated R40 had a stage two pressure ulcer but was staged incorrectly because a stage two pressure ulcer did not have granulation tissue, or slough and stated the MDS would need to be modified. RN-A stated he had been stating that staging of wounds was not being completed correctly and has completed education on staging of wounds. RN-A further stated he looked at wound pictures, but had to be more critical in what he saw and had not been doing that and further added it was important to have correct staging in order to know whether there has been a decline in the pressure ulcer which could indicate the wound was not being managed correctly.</p> <p>During interview on 3/12/25 at 2:14 p.m., the director of nursing (DON) stated she expected documentation align with the provider documentation and the documentation should be accurate. The DON verified the wound documentation note was not accurate for the wound staging according to the wound staging definitions.</p> <p>A policy was requested, however an email from the DON dated 3/12/25 at 3:01 p.m., indicated they utilized the RAI manual dated October 2024 for accuracy of assessments.</p>	F 641		

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F 641	<p>Continued From page 7</p> <p>R15's quarterly Minimum Data Set (MDS) dated 1/13/25 indicated he had intact cognition and identified his diagnoses of anxiety, depression, and bipolar disorder (a mental health condition characterized by extreme periods of elevated mood, or mania, and low mood, or depression). The MDS reported he was taking an antipsychotic medication on a routine basis and indicated a gradual dose reduction (GDR) had not been attempted and there was no physician documentation the GDR was clinically contraindicated. Furthermore, the MDS reported he was not taking an antianxiety medication.</p> <p>R15's active pharmacy orders were reviewed on dated 3/12/25 at 8:12 a.m., and revealed the following orders:</p> <ul style="list-style-type: none"> <li>- hydroxyzine hydrochloride (HCl) oral tablet 50 milligrams (mg), Give 50mg by mouth two times a day for anxiety," dated 12/15/23.</li> <li>- quetiapine fumarate oral tablet 100mg, Give 100mg by mouth three times a day for depression," dated 7/25/23.</li> </ul> <p>R15's medication administration record (MAR) dated 1/25 was reviewed 3/12/25 and reflected the following administered medication orders:</p> <ul style="list-style-type: none"> <li>- hydroxyzine hydrochloride (HCl) oral tablet 50 milligrams (mg), Give 50mg by mouth two times a day for anxiety," dated 12/15/23.</li> <li>- "Seroquel oral tablet (quetiapine fumarate), Give 300mg by mouth at bedtime," dated 10/30/24.</li> </ul> <p>R15's care plan dated 7/26/23, identified his use of psychotropic drug medications and his</p>	F 641		

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F 641	<p>Continued From page 8</p> <p>alteration in psychosocial well-being related to his diagnoses of schizophrenia (chronic mental health condition characterized by disruption in thought processes, perceptions, emotions, and social interactions), agoraphobia with panic (anxiety disorder characterized by intense fear of being in situations that might cause panic, or being trapped), and anxiety.</p> <p>A pharmacist's recommendation to prescriber dated 9/16/24, recommended a GDR to R15's psychotropic medications. The prescriber ordered a reduction of his evening Seroquel (quetiapine).</p> <p>A provider progress note dated 11/12/24 indicated under the "treatment and plan: no medication changes at this time. GDR attempts are not recommend[sic] at this time due to failed GDR and mental decompensation."</p> <p>Per interview on 3/12/25 at 9:19 a.m., licensed practical nurse (LPN)-A, confirmed R15 was taking hydroxyzine and verified this was an antianxiety medication.</p> <p>Per interview on 3/12/25 at 2:37 p.m., the director of nursing (DON) confirmed R15 was taking an antianxiety medication and had a failed GDR. The DON expected the MDS data to be accurate based on a review of a resident's recent medication list, progress notes, and provider notes.</p> <p>Per interview on 3/12/25 at 2:46 p.m., registered nurse (RN)-A confirmed accountability for completing R15's quarterly MDS dated 1/13/25. RN-A verified answering "no" him taking an antianxiety on the MDS and reviewed his MAR dated 1/25 and stated, "I missed the</p>	F 641		

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F 641	Continued From page 9 hydroxyzine." Furthermore, RN-A reviewed the provider progress note dated 11/12/24 and the pharmacist's recommendation to prescriber dated 9/16/24 and verified there had been a GDR performed as well as a clinical contraindication to GDR provided. RN-A stated the facility's procedure was to make a modification to erroneous MDS and submit the corrected one.	F 641			
F 645 SS=D	A policy pertaining to MDS accuracy was requested but not received. PASARR Screening for MD & ID CFR(s): 483.20(k)(1)-(3)  §483.20(k) Preadmission Screening for individuals with a mental disorder and individuals with intellectual disability.  §483.20(k)(1) A nursing facility must not admit, on or after January 1, 1989, any new residents with: (i) Mental disorder as defined in paragraph (k)(3)(i) of this section, unless the State mental health authority has determined, based on an independent physical and mental evaluation performed by a person or entity other than the State mental health authority, prior to admission, (A) That, because of the physical and mental condition of the individual, the individual requires the level of services provided by a nursing facility; and (B) If the individual requires such level of services, whether the individual requires specialized services; or (ii) Intellectual disability, as defined in paragraph (k)(3)(ii) of this section, unless the State intellectual disability or developmental disability authority has determined prior to admission- (A) That, because of the physical and mental	F 645		4/16/25	

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F 645	<p>Continued From page 10</p> <p>condition of the individual, the individual requires the level of services provided by a nursing facility; and</p> <p>(B) If the individual requires such level of services, whether the individual requires specialized services for intellectual disability.</p> <p>§483.20(k)(2) Exceptions. For purposes of this section-</p> <p>(i) The preadmission screening program under paragraph(k)(1) of this section need not provide for determinations in the case of the readmission to a nursing facility of an individual who, after being admitted to the nursing facility, was transferred for care in a hospital.</p> <p>(ii) The State may choose not to apply the preadmission screening program under paragraph (k)(1) of this section to the admission to a nursing facility of an individual-</p> <p>(A) Who is admitted to the facility directly from a hospital after receiving acute inpatient care at the hospital,</p> <p>(B) Who requires nursing facility services for the condition for which the individual received care in the hospital, and</p> <p>(C) Whose attending physician has certified, before admission to the facility that the individual is likely to require less than 30 days of nursing facility services.</p> <p>§483.20(k)(3) Definition. For purposes of this section-</p> <p>(i) An individual is considered to have a mental disorder if the individual has a serious mental disorder defined in 483.102(b)(1).</p> <p>(ii) An individual is considered to have an intellectual disability if the individual has an intellectual disability as defined in §483.102(b)(3)</p>	F 645		

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F 645	<p>Continued From page 11</p> <p>or is a person with a related condition as described in 435.1010 of this chapter. This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility failed to ensure a Level 1 Pre-Admission Screening (PAS) and, if needed, a a Level II Pre-Admission Screening and Resident Review (PASARR) were completed, retained in the medical record, and readily available to ensure continuity of care with mental health needs for 1 of 1 resident (R1) reviewed for PASARR.</p> <p>Findings include:</p> <p>R1's admission Minimum Data Set (MDS) dated 1/6/25, indicated R1 was not considered by the state level II PASRR process to have a serious mental illness and or intellectual disability or a related condition, had intact cognition, and had the following active diagnoses: bipolar disorder and schizophrenia.</p> <p>R1's physician's orders indicated the following order: 12/30/24, aripiprazole (an antipsychotic medication) 2 milligrams (MG) daily, take along with 5 mg for a total dose of 7 mg every day for schizoaffective disorder. 12/30/24, aripiprazole 5 mg daily, take 5 mg along with 2 mg for a total of 7 mg per day. 2/21/25, resident targeted behaviors of yelling, refusal of cares, isolation and interventions to redirect, remove from environment, and see notes, monitor resident for signs and symptoms of medication side effects and notify the physician if noted.</p> <p>R1's History and Physical dated 12/24/24,</p>	F 645	<p>R1's PAS referral was received by senior linkage line on 12/30/24 and was sent to lead agency. Facility reached out to lead agency to receive R1's level 1 and level 2 PASRR.</p> <p>All residents have the potential to be affected and will be reviewed to ensure a level 1 PAS, and if needed, a level 2 PASRR are uploaded in their electronic chart.</p> <p>Social service director and admissions coordinator will be educated on PASRR's and required documentation.</p> <p>Audits will be done on all new admissions to ensure a level 1 PAS, and if needed, a level 2, is on file for each resident. NHA or designee will conduct audits 5 days a week x2 weeks, then 3 days a week x2 weeks, then 1 day a week x2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 645	<p>Continued From page 12</p> <p>indicated R1 had active diagnoses of schizoaffective disorder and bipolar 1 disorder.</p> <p>R1's care plan, no date identified, indicated R1 had an alteration in mood and behavior due to bipolar and schizoaffective diagnoses and R1's goals included a stable mood and behavioral state, and R1 would respond to interventions.</p> <p>R1's medical record was reviewed and under a Miscellaneous form in the electronic medical record (EMR), indicated under the heading, "PASRR" included a document titled, "ER 12-30-24 PAS.pdf" The document, was opened and included a Preadmission Screening Referral form. The reason for the referral included two checked boxes indicating managed care, and waiver/AC/ECS. The form indicated the preadmission screen (PAS) was not completed by Senior LinkAge Line and the PAS required action. Further, the form indicated an OBRA Level II and level of care face to face assessment must be completed before nursing home admission. R1's medical record lacked evidence a Level 1 or, if needed, a Level II PASARR had been completed for R1 despite admitting to the nursing home a few months prior and having mental health-related diagnoses (i.e., schizophrenia and bipolar disorder) which could require active treatment. Further, the form lacked a determination of the PAS.</p> <p>During interview on 3/11/25 at 2:01 p.m., the admission coordinator (AC)-E stated she worked at the facility for over two years and her role included requesting preadmission screens through the Senior LinkAge Line. AC-E stated the PAS described what a resident was admitted for and showed the resident's medical and living</p>	F 645		

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F 645	<p>Continued From page 13</p> <p>situation. AC-E viewed R1's chart and initially stated R1 had a PAS, but after opening the form, stated R1 had a referral and the form was not the PAS. AC-E stated no other staff worked on preadmission screenings and verified the PAS were located in the EMR and a PAS would let staff know if a resident needed more than a level one and possibly a level II. AC-E stated R1 would stay at the nursing facility until they could find placement and further stated she would add R1 to her list to find out why she didn't have a PAS and stated it was probably overlooked and R1 should have had a PAS completed before being admitted to the facility.</p> <p>During interview on 3/11/25 at 2:11 p.m., the social services director (SSD) stated she was not a licensed social worker. SSD stated AC-E followed up on the PAS screens and further stated the purpose of the PAS was to help determine services and level II PAS included mental illness diagnoses like schizoaffective disorders would trigger a resident to need a level two PASARR. SSD stated the PAS screenings were supposed to be completed prior to admission and were uploaded into the miscellaneous tab. SSD viewed R1's miscellaneous tab and verified no PAS was uploaded and stated it was important to be completed for the whole team to assess and assist the resident and stated R1 could trigger for a level II PASARR and viewed the form and stated she thought AC-E uploaded the form at this time. SSD viewed a PAS scanned that indicated under a heading, "Does the person have a current diagnosis of a mental illness" that indicated R1 did not have a mental illness and stated she would ask their consultant if the answer for the question should be documented</p>	F 645		

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F 645	<p>Continued From page 14 as yes.</p> <p>During interview on 3/11/25 at 2:22 p.m., AC-E stated she contacted Senior LinkAge right away and stated they never attached the PAS and had just sent the PAS.</p> <p>During interview on 3/11/25 at 2:58 p.m., SSD stated she reviewed the PAS with the consultant and was directed to submit a new PAS with the diagnosis of R1 having mental illness to further determine if R1 needed a Level II PASARR.</p> <p>During interview on 3/12/25 at 11:04 a.m., the director of nursing (DON) stated she expected the PAS to be completed and stated it was important to have the PAS in order to know how to take care of residents and create an individualized care plan.</p> <p>During interview and observation on 3/12/25 at 11:14 a.m., registered nurse (RN)-C viewed R1's paper chart and verified there was no PAS in the paper chart.</p> <p>A care plan with revision history was requested, but was not provided.</p> <p>A policy, Pre-Admission Screening (PASSR), dated 6/2023, indicated social services would check for preadmission screening and OBRA Level II requirements and would ensure the initial Pre-Admission Screening results state that the resident meets level of care for purposes of medical assistance payment prior to the resident being admitted to the facility. The initial Pre-Admission Screening must be completed by a medical professional. If the requirements from the initial Pre-Admission Screening cannot be</p>	F 645		

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F 645	Continued From page 15 determined, the admission will need to be postponed until the county can complete a face to face assessment and the county can confirm the consumer meets requirements for nursing facility level of care under Medicaid. Upon receipt of the hard copy of the preadmission form(s) from the Senior Linkage Line and or Lead Agency, social services or designated staff person will upload documentation to the resident's medical record. For residents discharging to a nursing facility from the hospital, assisted living, clinic and out of state, before accepting the admission you must receive a copy of the initial PAS from the referring agency which notes that consumer meets criteria for placement.	F 645			
F 657 SS=D	Care Plan Timing and Revision CFR(s): 483.21(b)(2)(i)-(iii)  §483.21(b) Comprehensive Care Plans §483.21(b)(2) A comprehensive care plan must be- (i) Developed within 7 days after completion of the comprehensive assessment. (ii) Prepared by an interdisciplinary team, that includes but is not limited to-- (A) The attending physician. (B) A registered nurse with responsibility for the resident. (C) A nurse aide with responsibility for the resident. (D) A member of food and nutrition services staff. (E) To the extent practicable, the participation of the resident and the resident's representative(s). An explanation must be included in a resident's medical record if the participation of the resident and their resident representative is determined not practicable for the development of the resident's care plan.	F 657		4/16/25	

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F 657	<p>Continued From page 16</p> <p>(F) Other appropriate staff or professionals in disciplines as determined by the resident's needs or as requested by the resident.</p> <p>(iii) Reviewed and revised by the interdisciplinary team after each assessment, including both the comprehensive and quarterly review assessments.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review the facility failed to ensure care conferences were completed in a timely manner for 1 of 1 resident (R14) reviewed for care planning.</p> <p>Findings include:</p> <p>R14's annual Minimum Data Set (MDS) dated 2/11/25, indicated intact cognition and diagnosis of chronic kidney disease. If further indicated R14 required staff assistance with most activities of daily living (ADL) and mobility.</p> <p>During interview on 3/10/25 at 2:53 p.m. R14 stated he couldn't remember if he'd had a care conference or not.</p> <p>R14's progress note dated 8/27/2024, indicated the following: care conference held today with resident. Resident was in a good mood and willing to participate. Interdisciplinary team (IDT) went over cares, nursing needs, therapy, and other questions as needed. Resident had no questions. IDT will continue to assist resident as needed.</p> <p>R14's medical record lacked documentation of a care conference since 8/27/24.</p> <p>During interview on 3/12/25 at 2:53 p.m., the</p>	F 657	<p>R14 has a care conference scheduled for 3/31/25.</p> <p>All residents have potential to be affected and will be reviewed to ensure they have had a care conference within the last 90 days.</p> <p>Social services will be educated on care conference timing/scheduling.</p> <p>Audits will be done by NHA or designee to ensure care conferences are being held for residents each quarter or with a significant change. Audits will be done 5 days a week x2weeks, 3 days a week x2 weeks, and then 1 day a week x2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 657	Continued From page 17 director of social services stated the MDS nurse provided a list of who needed a care conference each month based off of the MDS assessments' schedule. When it was time for a resident to have a care conference she would talk to the team and then schedule it. The director of social services stated they were working on developing a better system in regards to care conferences and they're probably things that have been missed. She also verified the last time R14 had a care conference was on 8/27/25.  A facility policy on care conferences was requested but not received.	F 657		
F 686 SS=D	Treatment/Svcs to Prevent/Heal Pressure Ulcer CFR(s): 483.25(b)(1)(i)(ii)  §483.25(b) Skin Integrity §483.25(b)(1) Pressure ulcers. Based on the comprehensive assessment of a resident, the facility must ensure that- (i) A resident receives care, consistent with professional standards of practice, to prevent pressure ulcers and does not develop pressure ulcers unless the individual's clinical condition demonstrates that they were unavoidable; and (ii) A resident with pressure ulcers receives necessary treatment and services, consistent with professional standards of practice, to promote healing, prevent infection and prevent	F 686		4/16/25

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F 686	<p>Continued From page 18</p> <p>new ulcers from developing. This REQUIREMENT is not met as evidenced by:</p> <p>Based on observation, interview, and document review, the facility failed to ensure and identify accurate wound care assessments for 1 of 1 resident (R40) reviewed for pressure ulcers.</p> <p>Findings include:</p> <p>The Centers for Medicare and Medicaid (CMS) Long-Term Care Facility Resident Assessment Instrument (RAI) 3.0 user's manual dated 10/2024, identified the intent of section M, Skin Conditions, documented the risk, presence, appearance, and change of pressure ulcers and injuries. Further, it was important to recognize and evaluate each resident's risk factors and identify and evaluate all areas at risk of constant pressure and a complete assessment of skin was essential to an effective pressure ulcer prevention and skin treatment program. The RAI manual directed staff to review the medical record including skin care flow sheets, or other skin tracking forms, nurses' notes, and pressure ulcer injury risk assessments, speak with the treatment nurse and direct care staff on all shifts to confirm conclusions from the medical record review and observations of the resident, examine the resident and determine whether any ulcers, injuries, scars, or non-removable dressings, devices were present. Additionally, pressure ulcer staging was determined by the deepest anatomical stage and if a stageable pressure ulcer had been classified at a higher numerical stage than what was observed it should continue to be classified at the higher numerical stage and pressure ulcers do not heal in a reverse sequence. Stage three and four pressure ulcers</p>	F 686	<p>R40's skin and wound assessments now correctly show wound classified as Unstageable.</p> <p>Residents with pressure ulcers have the potential to be affected. All residents with pressure ulcers have been reviewed to ensure all current pressure ulcers are staged appropriately.</p> <p>Nurse management will be educated on pressure ulcer staging.</p> <p>DON or designee will audit the new skin and wound assessments each week and ensure the staging matches the Integrated Wound Care notes and is accurate. Audits will be done once a week x6 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 686	<p>Continued From page 19</p> <p>fill with granulation tissue that is never as strong as the tissue that was lost and hence is more prone to future breakdown. Clinical standards do not support reverse staging or back staging as a way to document healing as it does not accurately characterize what is occurring physiologically as the ulcer heals. Once a pressure ulcer has healed, it is documented as a healed pressure ulcer at its highest numerical stage and a previously closed pressure ulcer that opens again should be reported at its worst stage, unless currently presenting at a higher stage or unstageable.</p> <p>The State Operations Manual (SOM) provided guidance on pressure ulcer staging that defined the following pressure ulcers/injuries:            Stage 1 Pressure Injury is intact skin with localized redness.            Stage 2 Pressure ulcer is partial-thickness loss of skin with exposed dermis, presenting as a shallow open ulcer. The wound bed is viable, pink or red, moist, and may also present as an intact or open or ruptured blister. Adipose (fat) is not visible and deeper tissues are not visible. Granulation tissue (new connective tissue), slough, (non-viable tissue) and eschar (dead tissue) are not present.            Stage 3 Pressure Ulcer is a full thickness loss of skin in which subcutaneous fat may be visible in the ulcer and granulation tissue and epibole (rolled wound edges) are often present. Slough and or eschar may be visible but does not obscure the depth of tissue loss. If slough or eschar obscures the wound bed, it is an unstageable pressure ulcer/injury.            Stage 4 Pressure Ulcer is a full thickness skin and tissue loss with exposed or directly palpable fascia, muscle, tendon, ligament, cartilage or</p>	F 686		

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F 686	<p>Continued From page 20</p> <p>bone in the ulcer. Slough and or eschar may be visible on some parts of the wound bed. Epibole (rolled edges), undermining and or tunneling often occur. Depth varies by anatomical location. If slough or eschar obscures the wound bed, it is an unstageable pressure ulcer/pressure injury. Unstageable Pressure Ulcer is a full thickness skin and tissue loss in which the extent of tissue damage within the ulcer cannot be confirmed because the wound bed is obscured by slough or eschar. If the slough or eschar is removed, a stage 3 or stage 4 pressure ulcer will be revealed. If the anatomical depth of the tissue damage involved can be determined, then the reclassified stage should be assigned.</p> <p>Deep Tissue Pressure Injury (DTPI) is intact skin with a localized area of persistent non-blanchable deep red, maroon, purple discoloration due to damage of underlying soft tissue. This injury results from intense and or prolonged pressure and shear forces at the bone-muscle interface. The wound may evolve rapidly to reveal the actual extent of tissue injury, or may resolve without tissue loss. Once a deep tissue injury opens to an ulcer, reclassify the ulcer into the appropriate stage.</p> <p>R40's 5 day Minimum Data Set (MDS) assessment dated 6/27/24, indicated diagnoses of anemia, heart failure, peripheral vascular disease, diabetes mellitus, was at risk of developing pressure ulcers, had one stage 2 pressure ulcer that was not present upon admission. Additionally, R40 did not have any stage 1, 3, 4, pressure ulcers or unstageable, or deep tissue injuries.</p> <p>R40's annual MDS dated 1/3/25, indicated R40 had intact cognition, did not reject care, had</p>	F 686		

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F 686	<p>Continued From page 21</p> <p>anemia, coronary artery disease, heart failure, peripheral vascular disease, diabetes mellitus, one or more unhealed pressure ulcers/injuries, 0 stage 1 pressure injuries, one stage 2 pressure ulcer/injury, 0 stage 3 pressure ulcers, 0 stage 4 pressure ulcers, 0 unstageable pressure ulcers with slough or eschar, and 0 unstageable DTPI's. The MDS directed staff to report based on the highest stage of existing ulcers/injuries at their worst; and do not "reverse" stage.</p> <p>R40's Optional State Assessment (OSA) dated 1/3/25, indicated R40 required extensive assist with bed mobility, transfers, and toileting.</p> <p>R40's Care Area Assessment (CAA) for pressure ulcers indicated R40 had a stage two pressure ulcer and required partial assistance with lying to sitting, had a foley catheter which could increase the risk for pressure due to the tubing and the pressure ulcer would be addressed in the care plan.</p> <p>R40's care plan undated, indicated R40 had limited physical mobility related to non weight bearing to right lower extremity and interventions included to turn and reposition every 3-4 hours, uses wheelchair and ensure foot peddles are in place. R40's care plan indicated a self care deficit and required assist of one with grooming, bathing, and was encouraged to participate in dressing by placing arms and legs into clothing and staff were to assist with placing and removing socks and shoes. Further, R40's care plan indicated R40 was admitted with a right heel surgical wound and diabetic ulcer and on 3/9/23, had a right heel blister/suspected deep tissue injury and a pressure injury on the right Achilles and interventions included evaluating and treating</p>	F 686		

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F 686	<p>Continued From page 22</p> <p>per physician's orders, evaluate for signs and symptoms of possible infections, was followed by the wound care team weekly, took prostat (a protein supplement), was to be turned and repositioned every 3 to 4 hours, monitor skin during cares, provide a pressure reducing mattress.</p> <p>R40's physician's orders included the following order: 1/29/25, cleanse right heel wound with generic wound cleanser or Vashe, apply collagen until Santyl arrives, cover with foam dressing, change dressing daily and as needed, inform the provider with any signs or symptoms of infection. Dressings need dates and initials and change daily for wound care.</p> <p>R40's Re-Admission History and Physical note dated 6/11/24, located in the progress notes indicated R40 had an unstageable pressure ulcer to the right heel.</p> <p>R40's Re-Admission History and Physical note dated 6/25/24, located in the progress notes indicated R40 had an unstageable pressure ulcer of the right heel.</p> <p>R40's progress notes dated 1/1/25, indicated, R40's "pressure ulcer 2 right Achilles-improving".</p> <p>R40's Skin and Wound Evaluation form dated 11/21/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 2.9 centimeters (CM) long by 2.6 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 30% granulation tissue, 20% slough, and 50% eschar.</p>	F 686		

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F 686	<p>Continued From page 23</p> <p>R40's Skin and Wound Evaluation form dated 12/3/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 3.6 cm long by 2 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 60% granulation tissue, and 40% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 12/10/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 3.9 centimeters (CM) long by 2.6 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 30% granulation tissue, 60% slough, and 10% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 12/17/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 3.1 centimeters (CM) long by 2.0 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 30% granulation tissue, 60% slough, and 10% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 12/26/24, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 2.9 centimeters (CM) long by 2.1 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 70% granulation tissue, 10% slough, and 20% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 1/1/25, indicated R40 had an in-house acquired</p>	F 686		

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F 686	<p>Continued From page 24</p> <p>stage two pressure ulcer to the right Achilles that had been present for one week measuring 4.1 (cm) long by 2.7 cm wide. Further, the note indicated the wound bed contained 80% granulation tissue, 20% slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 1/7/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 4.1 centimeters (CM) long by 2.5 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 90% granulation tissue, and 10% slough.</p> <p>R40's Skin and Wound Evaluation form dated 1/14/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 7.4 cm long by 2.8 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 90% granulation tissue, 10% slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 1/21/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 2.4 cm long by 1.3 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 50% granulation tissue, no slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 1/28/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 5.5 cm long by 2.4 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the</p>	F 686		

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F 686	<p>Continued From page 25</p> <p>wound bed contained 70% granulation tissue, no slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 2/6/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 5 cm long by 3.9 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 70% granulation tissue, 10% slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 2/13/25, indicated R40 had a stage two pressure ulcer to the right Achilles that had been present for one week measuring 4.5 cm long by 2.7 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 70% granulation tissue, 10% slough, and 20% eschar.</p> <p>R40's Skin and Wound Evaluation form dated 2/20/25, indicated R40 had an unstageable pressure ulcer to the right Achilles that had been present for one week measuring 4.5 cm long by 2.7 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 70% granulation tissue, 10% slough, and no eschar.</p> <p>R40's Skin and Wound Evaluation form dated 2/25/25, indicated R40 had an unstageable pressure ulcer to the right Achilles that had been present for one week measuring 4.3 cm long by 2.4 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 20% granulation tissue, 20% slough, and no eschar.</p>	F 686		

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F 686	<p>Continued From page 26</p> <p>R40's Skin and Wound Evaluation form dated 3/7/25, indicated R40 had an unstageable pressure ulcer to the right Achilles that had been present for one week measuring 4.4 cm long by 2.8 cm wide. Further, and contrary to the definition of a stage two pressure ulcer, the note indicated the wound bed contained 20% granulation tissue, 20% slough, and no eschar.</p> <p>R40's Integrated Wound Care note dated 12/3/24, indicated a right lateral heel unstageable pressure ulcer measuring 3.8 cm long by 2.7 cm wide by 0.3 cm deep with 30% necrotic tissue, 20% eschar, and 50% granulation tissue.</p> <p>R40's Integrated Wound Care Note dated 12/17/24, indicated a right lateral heel unstageable pressure ulcer measuring 3 cm long by 3.5 cm wide by 0.5 cm deep with 10% necrotic tissue, 30% granulation tissue, and 60% slough.</p> <p>R40's Integrated Wound Care note dated 12/26/24, indicated R40 had a right heel unstageable pressure ulcer. Further, the wound was 2 cm long by 4.7 cm wide, by 0.4 cm deep and contained 10% slough, 20% necrotic tissue, and 70% granulation tissue.</p> <p>R40's Integrated Wound Care note dated 1/2/24, indicated R40 had a right heel unstageable pressure ulcer. Further, the wound was 2.5 cm long, by 4 cm wide by 0.4 cm deep and contained 20% slough, and 80% granulation.</p> <p>R40's Integrated Wound Care note dated 1/2/25, indicated a right lateral heel unstageable pressure ulcer measuring 4 cm long by 2.5 cm wide by 0.5 cm deep with 10% slough and 90% granulation tissue.</p>	F 686		

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F 686	<p>Continued From page 27</p> <p>R40's Integrated Wound Care note dated 1/14/25, indicated a right lateral heel unstageable pressure ulcer measuring 7.3 cm long by 2.8 cm wide by 0.4 cm deep with 10% slough and 90% granulation tissue.</p> <p>R40's Integrated Wound Care note dated 1/21/25, indicated a right lateral heel unstageable pressure ulcer measuring 2.3 cm long by 1.3 cm wide by 0.3 cm deep with 50% necrotic tissue, and 50% granulation tissue.</p> <p>R40's Integrated Wound Care note dated 1/28/25, indicated a right lateral heel unstageable pressure ulcer measuring 5.5 cm by 2.4 cm by 0.3 cm with 30% necrotic tissue and 70% granulation tissue.</p> <p>R40's Integrated Wound Care note dated 2/6/25, indicated a right lateral heel unstageable pressure ulcer measuring 5 cm long by 3.9 cm wide by 0.2 cm deep with 20% necrotic tissue, 70% granulation tissue, and 10% slough.</p> <p>R40's Integrated Wound Care note dated 2/13/25, indicated a right lateral heel unstageable pressure ulcer measuring 4.5 cm long by 2.7 cm wide by 0.2 cm deep with 20% necrotic tissue, 70% granulation tissue and 10% slough.</p> <p>R40's Integrated Wound Care note dated 2/20/25, indicated a right lateral heel unstageable pressure ulcer measuring 4.5 cm long by 2.7 cm wide by 0.2 cm deep with 20% necrotic tissue, 70% granulation tissue and 10% slough.</p> <p>R40's Integrated Wound Care note dated 2/25/25, indicated a right lateral heel unstageable</p>	F 686		

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F 686	<p>Continued From page 28</p> <p>pressure ulcer measuring 4.3 cm long by 2.4 cm wide by 0.2 cm deep with 30% necrotic tissue, 20% granulation, and 20% slough.</p> <p>R40's Integrated Wound Care note dated 3/7/25, indicated a right lateral heel unstageable pressure ulcer measuring 4.4 cm long by 2.8 cm wide by 0.2 cm deep with 30% necrotic tissue, 20% granulation, and 20% slough.</p> <p>During interview on 3/10/25 between 5:16 p.m., and 5:18 p.m., R40 stated she was going to the hospital on 3/11/25 to have an X-RAY of her leg to determine blood flow and further stated her heel was painful and pointed to her right heel and stated that was why it was elevated. R40's wheelchair foot rest on the right was pulled down and R40 had a sock on with her right heel resting directly on the foot rest. R40 stated she had a sore on her heel and stated staff usually changed the dressing in the morning, but stated it was changed mid morning on 3/10/25.</p> <p>During interview on 3/12/25 at 7:52 a.m., R40 stated she had a procedure the day prior to increase the blood flow and help heal her foot.</p> <p>During interview on 3/12/25 at 8:26 a.m., registered nurse (RN)-C stated R40 had a diabetic sore on her heel that was a stage II and had some slough but no eschar.</p> <p>During interview on 3/12/25 at 11:19 a.m., RN-B stated she conducted wound rounds with the nurses weekly and stated the wound assessment was documented under the Skin and Wound in the Forms tab. RN-B stated R40 had an unstageable pressure ulcer on the right Achilles heel. RN-B viewed the 2/13/25, note and verified</p>	F 686		

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F 686	<p>Continued From page 29</p> <p>the wound was documented as a stage two with slough and eschar and stated it could be documented as a stage two because of improvement in the heel. RN-B stated she was not sure if a stage two pressure ulcer presented with eschar and slough and was not sure about the recent staging because sometimes there was improvement in the wound and then the wound comes back.</p> <p>During interview on 3/12/25 at 12:14 p.m., registered nurse (RN)-A stated he was the MDS coordinator and scheduled the MDS assessments, completed assessments, and made sure everyone completed their sections. RN-A stated he completed sections A, E, G, H, GG, I, J, K, L, M, N, O, P, and S. RN-A stated they were supposed to schedule skin assessments each week and when looking for skin assessments, looked at the Forms in the electronic medical record (EMR). RN-A stated R40 had an Annual MDS on 1/3/25, and when completing the MDS, looked for a skin assessment for 1/3/25, or around that date. RN-A stated they used to have a wound care manager who documented notes directly into progress notes but did not locate notes related to staging during the window for the MDS after reviewing the progress notes. RN-A stated he could not recall where he went to find the stage of the wound and knew it had been a stage two pressure ulcer and stated he used the RAI manual for guidance. RN-A stated according to the RAI manual, a stage two pressure ulcer was the first layer of skin removed and could also be a blister. RN-A further stated a stage two pressure ulcer did not contain slough, eschar, or granulation tissue. RN-A stated he viewed the Wound Evaluation form 1/1/25, for the MDS and</p>	F 686		

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F 686	<p>Continued From page 30</p> <p>stated the form indicated R40 had a stage two pressure ulcer but was staged incorrectly because a stage two pressure ulcer did not have granulation tissue, or slough and stated the MDS would need to be modified. RN-A stated he had been stating that staging of wounds was not being completed correctly and has completed education on staging of wounds. RN-A further stated he looked at wound pictures, but had to be more critical in what he saw and had not been doing that and further added it was important to have correct staging in order to know whether there has been a decline in the pressure ulcer which could indicate the wound was not being managed correctly.</p> <p>During interview on 3/12/25 at 12:14 p.m., RN-A stated he spoke with the director of nursing and was told they had documentation the wound was almost healed so it would have been a stage two at that point, and added he did not want to retract anything he said.</p> <p>During observation and interview on 3/12/25 at 1:43 p.m., RN-C changed R40's dressing and completed measurements. RN-C stated R40's wound measured 1 cm long by 2.2 cm wide by 0.1 cm deep. RN-C stated he had training on how to stage wounds previously but not at the facility and stated R40's wound was at a stage three.</p> <p>During interview on 3/12/25 at 2:14 p.m., the director of nursing (DON) stated staff rounded with a wound nurse practitioner who changed orders or made recommendations and staged the wounds and nurses at the facility completed wound assessments and completed the Skin and Wound Evaluation and added if you don't click</p>	F 686		

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F 686	Continued From page 31 and change information, the information stays in place in the note. The DON stated R40 admitted with a wound vac (a treatment for the wound) and the wound healed and reopened and added R40 was non compliant and added R40 had an angiogram 3/11/25. The DON stated she expected the provider stage the wound and nursing documentation should align with the provider and should be accurate and verified wound documentation notes were not accurate for the wound staging according to the staging definitions.  A care plan with the revision history was requested, but was not provided.  A policy, Skin Assessment and Wound Management dated 2/2025, indicated a weekly skin inspection was completed by licensed staff. When a new pressure ulcer is identified the following actions will be taken, notify the provider/treatment ordered, notify the resident representative, complete education with the resident and resident representative including risks and benefits, initiate the skin and wound evaluation, notify the nurse manager/wound nurse.	F 686			
F 689 SS=D	Free of Accident Hazards/Supervision/Devices CFR(s): 483.25(d)(1)(2)  §483.25(d) Accidents. The facility must ensure that - §483.25(d)(1) The resident environment remains as free of accident hazards as is possible; and  §483.25(d)(2) Each resident receives adequate supervision and assistance devices to prevent accidents.	F 689		4/16/25	

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F 689	<p>Continued From page 32</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review the facility failed to demonstrate safe patient handling to reduce the risk for accidents for 1 of 1 residents (R13) reviewed for safety with mechanical lift assisted transfers.</p> <p>Findings include:</p> <p>R13's annual Minimum Data Set (MDS) dated 12/12/24, indicated he had intact cognition and was dependent on staff for chair/bed-to-chair transfers. The MDS reported diagnoses of hemiplegia (one-sided weakness) following a cardiovascular accident (CVA, stroke), and hemiplegia (one-sided paralysis).</p> <p>R13's Care Area Assessment for functional abilities dated 12/19/24, indicated he was dependent on staff for transfers.</p> <p>R13's care plan revised 1/3/24, identified his activities of daily living (ADL) self-care deficit and directed staff to provide 2-person assistance using the "EZstand".</p> <p>A therapy to nursing communication form dated 5/10/24, indicated R13's transfer status under "Transfers: EZ Stand".</p> <p>During interview on 3/10/25 at 4:52 p.m., R13 reported using the "EZ Stand" for transfers and stated it caused him pain to his ribs and lungs. He stated during some transfers, he would "hang there" and when he would attempt to tell staff to lower him down "because I can't breathe and its pushing on my lungs", staff would tell him to stop "hollering" at him.</p>	F 689	<p>R13's transfer status was downgraded from EZ stand to hoyer lift. R13 was evaluated by therapy on 3/13/25 and continues to work with PT on transfers.</p> <p>All residents that require staff assistance with transfers, but are not fully dependent (hoyer lift) have the potential to be affected and will be reviewed to ensure current transfer status is safe and appropriate.</p> <p>All nursing staff will be educated on identifying unsafe transfers and the appropriate steps to ensure resident safety, including downgrading transfer status when necessary.</p> <p>DON or designee will audit staff-assisted transfers to ensure resident is transferring safely. Audits will be done on at least 1 transfer per day, 5 days a week x2 week, then 3 days a week x2 weeks, then 1 time a weekx2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 689	<p>Continued From page 33</p> <p>During observation and interview on 3/12/25 at 12:54 p.m., nursing assistant (NA)-A and NA-B were in R13's room to transfer him from his wheelchair to his bed. NA-B stood behind the mechanical standing lift with the remote control in hand. NA-A put the lift sling behind R13's back and ensured all the sling loops were fastened to the standing lift. NA-A stated because R13 had weakness to his left side, they would need to make sure he was holding onto the standing lift with his hand properly. After the NAs ensured his feet were secured on the standing lift's platform, NA-B used the remote control to lift R13 from the wheelchair into a semi-upright position. R13 was unable to stand straight up; his knees were bent at approximately 80 degrees while his arms were bent at approximately 90 degrees. He was hanging by both shoulders from the standing lift with the sling behind his back and shoulders. NA-A told NA-B to "stop going up" and began to move the standing lift over towards R13's bed. The surveyor asked R13 if he was able to stand up any straighter. He attempted and was able to straighten his legs, but his arms remained bent, and his upper extremity posture remained unchanged. NA-A pushed the standing lift over his bed and began to lower R13 onto the bed. R13 stated the transfer felt "okay" but said if he had to transfer like that repeatedly, he would be "in a lot of pain." NA-A stated sometimes R13 would say he did not want to get up or that he was not feeling well, and they would leave him be and notify the nurse. When asked if the observed transfer was safe and positioning was appropriate, NA-A said it was okay because he was able to get into bed, but if he was not okay, they would report that to the nurse.</p>	F 689		

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F 689	Continued From page 34 During interview on 3/12/25 at 1:11 p.m., physical therapist assistant (PTA) stated if a resident using a mechanical standing lift was observed using bad posturing, like bowing of the arms out or "sort of hanging there", it would be an indication the resident should be evaluated by therapy. PTA expected NAs to report such observations to nurses or nurse managers so it could be passed along to the interdisciplinary team (IDT) to discuss. PTA stated residents struggling to use mechanical standing lifts should be reviewed during IDT so they can get on the therapy caseload for evaluation. PTA stated the residents exhibiting bad posturing techniques during transfers could present risks such cutting off blood flow or circulation to their shoulder joints or risk slipping out of the mechanical standing lift during a transfer. PTA denied being notified of any concerns regarding R13's transfers.  During interview on 3/12/25 at 2:30 p.m., the director of nursing (DON) stated a safe mechanical standing lift transfer would be one in which a resident could stand for eight seconds, they could fully hold onto or grasp the lift bars and could come to a complete stand without hanging in the mechanical standing lift. The DON expected if staff observed an unsafe transfer to complete a lift and mobility assessment, change the transfer status and update the care plan accordingly. Finally, the DON expected this to be communicated during morning standup so therapy could get involved.	F 689			
F 757 SS=D	Drug Regimen is Free from Unnecessary Drugs CFR(s): 483.45(d)(1)-(6)	F 757		4/16/25	

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F 757	<p>Continued From page 35</p> <p>§483.45(d) Unnecessary Drugs-General. Each resident's drug regimen must be free from unnecessary drugs. An unnecessary drug is any drug when used-</p> <p>§483.45(d)(1) In excessive dose (including duplicate drug therapy); or</p> <p>§483.45(d)(2) For excessive duration; or</p> <p>§483.45(d)(3) Without adequate monitoring; or</p> <p>§483.45(d)(4) Without adequate indications for its use; or</p> <p>§483.45(d)(5) In the presence of adverse consequences which indicate the dose should be reduced or discontinued; or</p> <p>§483.45(d)(6) Any combinations of the reasons stated in paragraphs (d)(1) through (5) of this section.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on observation, interview and document review the facility failed to ensure antifungal medications without an end date were monitored and evaluated for the appropriateness of continued use for 1 of 1 residents (R3) reviewed who were prescribed antifungal medications.</p> <p>Findings include:</p> <p>R3's quarterly Minimum Data Set (MDS) dated 12/11/24, indicated she was rarely or never understood and identified diagnoses of non-traumatic brain dysfunction, neurosyphilis (a form of syphilis, a sexually transmitted infection,</p>	F 757	<p>R3's order for antifungal medication was updated by provider with clarification to continue medication due to resident's history rash recurring.</p> <p>All residents with order for antifungal medications have potential to be affected. All residents with current orders for antifungal medications will be reviewed to ensure order specifies duration.</p> <p>All licensed nurses will be educated on requesting stop dates and/or clarification for antifungal medication orders.</p>	

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F 757	<p>Continued From page 36</p> <p>that affects the brain and spinal cord), non-Alzheimer's dementia (a condition that causes a decline in cognitive function, including memory, thinking, language, and problem-solving). The MDS indicated she required two-person extensive assistance with personal and toileting cares.</p> <p>R13's order summary dated 3/13/25, included the order:</p> <p>- "Nyamyc external powder 100000 unit/gram (gm) (Nystatin Topical), Apply to abdominal folds topically two times a day for candidiasis," dated 10/28/24 with no end date.</p> <p>R13's medication administration record (MAR) dated 11/24 was reviewed on 3/11/25 and reflected administration of the Nystatin powder as ordered.</p> <p>R13's MAR dated 12/24 was reviewed on 3/11/25 and reflected administration of the Nystatin powder as ordered.</p> <p>R13's MAR dated 1/25 was reviewed on 3/11/25 and reflected administration of the Nystatin powder as ordered.</p> <p>R13's MAR dated 2/25 was reviewed on 3/11/25 and reflected administration of the Nystatin powder as ordered.</p> <p>R13's MAR dated 3/1/25 through 3/10/25 was reviewed on 3/11/25 and reflected administration of the Nystatin powder as ordered.</p> <p>A weekly skin inspection dated 11/5/24, indicated "previously noted resident had redness/rash on</p>	F 757	<p>DON or designee will audit new orders for antifungal medication to ensure there is a specified duration of use. Audits will be done 5 days a week x2 weeks, then 3 days a week x2 weeks, then 1 day a week x2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 757	<p>Continued From page 37</p> <p>left lower abdomen folds" related to "moisture". The inspection indicated "treatment is in place."</p> <p>A weekly skin inspection dated 11/12/24, indicated redness/rash on left lower abdomen folds was clearing.</p> <p>A weekly skin inspection dated 11/26/24, indicated "slight redness/rash on left lower abdomen folds is clearing. Nystatin applied."</p> <p>A weekly skin inspection dated 12/17/24, indicated "skin is intact".</p> <p>A weekly skin inspection dated 12/24/24, indicated no skin concerns noted, "Nystatin applied under adm [sic] folds/groin."</p> <p>During observation on 3/12/25 at 12:45 p.m., licensed practical nurse (LPN)-B was at R3's bedside and assessed her abdominal folds and groin skin. LPN-B confirmed there was no redness and stated her skin was intact. LPN-B stated R3 used to have redness and moisture to this area, but it had cleared now.</p> <p>During interview on 3/12/25 at 12:25 p.m., certified physician assistant (PA-C) confirmed familiarity with R3's care but stated the Nystatin powder order was "before my time" since PA-C started in December and the order was dated 10/28/24. PA-C stated when treating a true fungal infection, the normal procedure would be to prescribe it with and end date after two weeks and then reassess to determine if it was effective and reconsider the necessity of the topical antifungal and question if it could be an as needed (PRN) order. PA-C stated, "if she was continuing on it, I would want to know why. I</p>	F 757		

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F 757	<p>Continued From page 38</p> <p>would probably have stopped [it]. Why would you give something that isn't necessary."</p> <p>During interview on 3/12/24 at 1:41 p.m., registered nurse (RN)-B was not sure if topical antifungals were monitored by the infection preventionist, however, stated nursing staff should be looking for an end date and monitoring for symptoms of improvement or worsening condition. RN-B stated for topical antifungal, like Nystatin powder, if the area was cleared, nursing staff should call the provider to get monitoring orders and consider asking for the medication to be as needed.</p> <p>During interview on 3/12/25 at 2:26 p.m., the director of nursing (DON), also the facility's infection preventionist, stated they did not track topical antifungals in ICAR, a tool used to systematically assess a healthcare facilities infection prevention and control practices. The DON stated because floor nurses were administering the topical Nystatin powder, they would be responsible for assessing the area and were expected to update the provider and ask if the medication order could be re-evaluated once the area had improved. The DON stated the Nystatin powder should have been discontinued once the skin concern improved.</p> <p>Per facility policy titled skin assessment and wound management last revised 2/25, when a significant alteration in skin integrity was noted, staff were directed to notify the provider for treatment orders and perform skin and/or wound evaluations and update the care plan to include interventions. For ongoing skin issues, staff were directed to follow ongoing treatments per provider order and update the provider and care plan as</p>	F 757		

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FORM APPROVED  
OMB NO. 0938-0391

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NAME OF PROVIDER OR SUPPLIER  <b>THE VILLAS AT ROSEVILLE</b>		STREET ADDRESS, CITY, STATE, ZIP CODE <b>1000 LOVELL AVENUE ROSEVILLE, MN 55113</b>		
(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 757  F 883 SS=D	Continued From page 39 needed. Influenza and Pneumococcal Immunizations CFR(s): 483.80(d)(1)(2)  §483.80(d) Influenza and pneumococcal immunizations §483.80(d)(1) Influenza. The facility must develop policies and procedures to ensure that- (i) Before offering the influenza immunization, each resident or the resident's representative receives education regarding the benefits and potential side effects of the immunization; (ii) Each resident is offered an influenza immunization October 1 through March 31 annually, unless the immunization is medically contraindicated or the resident has already been immunized during this time period; (iii) The resident or the resident's representative has the opportunity to refuse immunization; and (iv) The resident's medical record includes documentation that indicates, at a minimum, the following: (A) That the resident or resident's representative was provided education regarding the benefits and potential side effects of influenza immunization; and (B) That the resident either received the influenza immunization or did not receive the influenza immunization due to medical contraindications or refusal.  §483.80(d)(2) Pneumococcal disease. The facility must develop policies and procedures to ensure that- (i) Before offering the pneumococcal immunization, each resident or the resident's representative receives education regarding the benefits and potential side effects of the	F 757  F 883		4/16/25

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F 883	<p>Continued From page 40</p> <p>immunization;</p> <p>(ii) Each resident is offered a pneumococcal immunization, unless the immunization is medically contraindicated or the resident has already been immunized;</p> <p>(iii) The resident or the resident's representative has the opportunity to refuse immunization; and</p> <p>(iv) The resident's medical record includes documentation that indicates, at a minimum, the following:</p> <p>(A) That the resident or resident's representative was provided education regarding the benefits and potential side effects of pneumococcal immunization; and</p> <p>(B) That the resident either received the pneumococcal immunization or did not receive the pneumococcal immunization due to medical contraindication or refusal.</p> <p>This REQUIREMENT is not met as evidenced by:</p> <p>Based on interview and document review, the facility failed to ensure 1 of the 5 residents (R1) reviewed for immunizations was offered and/or provided the pneumococcal vaccination series as recommended by the Centers for Disease Control (CDC) to help reduce the risk of associated infection(s).</p> <p>Findings include:</p> <p>The CDC's PneumoRecs VaxAdvisor for Vaccine Providers dated 9/12/24, recommended based on R1's age and vaccine history to give a dose of PCV15, PCV20 or PCV21 at least 1 year after the last dose of PPSVR23. "Regardless of which vaccine is used (PCV15, PCV20, or PCV21), their pneumococcal vaccinations are complete."</p> <p>R1's Clinical Profile printed on 3/12/25, indicated</p>	F 883	<p>R1 was provided education regarding the benefits and potential side effects of pneumococcal immunization on 3/12/25 and received PCV20 on 3/13/25.</p> <p>All residents have the potential to be affected. Per facility policy, all current residents will be assessed for current immunization status within 5 days of admission, and will be offered the vaccine, when indicated, within 30 days of admission.</p> <p>Nurse management will be educated on facility policy for offering pneumococcal vaccinations.</p> <p>Audits will be done on all new admissions to review vaccination status and to ensure</p>	

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F 883	<p>Continued From page 41</p> <p>R1 was 78 years old. The immunization record, printed 3/12/25, indicated R1 received a PPSV23 on 1/17/13.</p> <p>R1's electronic medical record (EMR) lacked evidence of shared clinical decision making occurring with the physician for giving one dose of PCV15, PCV20, or PCV21 as it was more than 12 years after the previous pneumococcal dose. R26's EMR and paper chart lacked evidence of R1 being offered or receiving any pneumococcal doses or education.</p> <p>During interview on 3/11/25 at 3:33 p.m., R1 stated didn't remember staff talking or offering a pneumococcal vaccine. R1 stated would have accepted a pneumovax vaccine if offered.</p> <p>During interview on 3/12/25 at 11:28 a.m., the infection preventionist/director of nursing (DON) stated besides the duties as the infection preventionist and DON position, had also been fulfilling the nurse manager's duties. DON stated missed auditing R1. DON stated a resident who received the recommended pneumovax vaccination would prevent someone from getting pneumonia, or if they would get pneumonia, the symptoms won't be as severe.</p> <p>Facility's policy titled Pneumococcal Policy dated 2/2024 indicated: It is the practice of the Health Care Facility to offer all residents the pneumococcal vaccines to aid in the prevention of pneumococcal/pneumonia infections. Facility policy indicated the purpose was to follow recommendations of the Advisory Committee on Immunization Practices (ACIP) Centers for Disease Control (CDC) and/or the state Department of Health for prevention of</p>	F 883	<p>pneumovax is offered when indicated. Audits will be done by DON or designee 5 days a week x2 weeks, 3 days a week 2 weeks, and then 1 day a week x2 weeks or until compliance is met. Results of audits will be brought to QAPI committee by NHA for input on the need to increase, decrease or discontinue the audits.</p>	

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F 883	Continued From page 42 Pneumococcal disease by identifying those residents at risk for Pneumococcal disease and offering Pneumococcal vaccination.	F 883		

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K 000	<p><b>INITIAL COMMENTS</b></p> <p><b>FIRE SAFETY</b></p> <p>An annual Life Safety Code survey was conducted by the Minnesota Department of Public Safety, State Fire Marshal Division on 3/12/2025. At the time of this survey, The Villas of Roseville was found in compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2012 edition of National Fire Protection Association (NFPA) 101, Life Safety Code (LSC), Chapter 19 Existing Health Care and the 2012 edition of NFPA 99, the Health Care Facilities Code.</p> <p>(Building Info)</p> <p>The Villas At Roseville is a 2-story building with no basement. The building was constructed at two different times. The original building was built in 1968 and was determined to be of Type II(222) construction. In 1992, an addition was constructed to the Northside that was determined to be of Type II(222) construction. Because the original building and the one addition are of the same type of construction, the facility was surveyed as one building.</p> <p>A complete fire sprinkler system protects the building. The facility has a fire alarm system with entire corridor smoke detection, resident rooms, and spaces open to the corridors that are monitored for automatic fire department notification.</p> <p>The facility has a capacity of 63 beds and had a census of 60 at time of the survey.</p>	K 000			

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

TITLE

(X6) DATE

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

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K 000	Continued From page 1  The requirement at 42 CFR, Subpart 483.70(a) is MET.	K 000		