

Department of Health & Human Services
Centers for Medicare & Medicaid Services

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No. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 325111	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 05/06/2026
NAME OF PROVIDER OR SUPPLIER Northrise Wellness & Rehabilitation		STREET ADDRESS, CITY, STATE, ZIP CODE 2884 North Road Runner Parkway Las Cruces, NM 88011	
For information on the nursing home's plan to correct this deficiency, please contact the nursing home or the state survey agency.			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (Each deficiency must be preceded by full regulatory or LSC identifying information)		
F 0656 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Some	Develop and implement a complete care plan that meets all the resident's needs, with timetables and actions that can be measured. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on record review and interview, the facility failed to develop and implement accurate, person-centered comprehensive care plans for 8 (R #13, R #15, R #28, R #30, R # 31, R #41, R #54, and R #59) of 8 (R #13, R #15, R #28, R #30, R # 31, R #41, R #54, and R #59) residents reviewed for comprehensive care plans. This deficient practice could likely result in staff being unaware of the current and actual needs of the residents and worsening of the pressure ulcers. The findings are: R #13 A. Record review of R #13's admission Record, no date, revealed R #13 was admitted to the facility on [DATE]. B. Record review of R #13's physician's orders revealed the following orders; 1. Dated 03/15/26 and discontinued 04/13/26, for hydrocodone-acetaminophen (a prescription opioid combination used to treat moderate to severe pain) 5-325 mg every 6 hours PRN for pain. 2. Dated 04/13/26, for hydrocodone-acetaminophen 5-325 mg every 4 hours as needed for pain. 3. Dated 03/16/26 for Lovenox (a prescription low molecular weight heparin (LMWH) injection used to treat and prevent deep vein thrombosis (DVT, a serious condition where a blood clot forms in a deep vein) and pulmonary embolism (PE)) injection 40mg/0.4mL once a day for DVT prophylaxis (to prevent diseases). C. Record review of R #13's admission MDS, dated [DATE], revealed staff documented R #13 was taking the following high-risk medications (drugs with a heightened risk of causing significant patient harm, or even death): 1. Anticoagulants (often called blood thinners, prevent or reduce blood clot formation by interfering with the body's coagulation process). 2. Opioid medication (powerful, regulated pain-relievers used for moderate-to-severe pain). D. Record review of R #13's care plan, dated 03/16/26, revealed staff did not document that R #13 was taking anticoagulant or opioid medications. E. On 05/04/26 at 2:21 PM, during an interview, the DON and Regional Clinical Nurse (RCN) confirmed the following: 1. R #13 had orders for anticoagulants and opioid medications. (continued on next page)		

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.

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

TITLE

(X6) DATE

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