

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11965359	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED 10/23/2024
NAME OF PROVIDER OR SUPPLIER YESENIA ALF		STREET ADDRESS, CITY, STATE, ZIP CODE 15608 SW 63 TERRACE MIAMI, FL 33193			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETE DATE
A 000	Initial Comments A re-licensure and a generator monitoring survey was conducted at Yesenia ALF on . A deficiency was identified at the time of the survey.	A 000			
A 033 SS=D	59A-36.007(B) PHYSICAL (B) PHYSICAL . Residents for whom a physician has prescribed a physical must have a written care plan for the use of the physical . The care plan must be developed within 14 days of the device being prescribed, and prior to use on the resident. (a) The care plan must specify: 1. The device prescribed for use; 2. The maximum amount of time the resident is to have the applied each day; and, 3. In what manner and frequency staff will monitor, observe, and report to the physician any injuries, increase in agitation, signs and symptoms of , or decline in mobility or function related to the use of the prescribed . (b) Facility staff must ensure that the device is applied appropriately and safely. (c) The resident's physician must review the appropriateness of the continued use of the physical annually, and documentation of this review must be maintained in the resident's record. If the resident's ability to independently remove or avoid the device fluctuates, the device must be considered a physical and all requirements of this subsection apply. This Statute or Rule is not met as evidenced by: Based on observation, record review and interview the facility failed to develop a plan of care for the use of physician (full bedrail's) for one out of six sampled resident (Resident #3).	A 033			

AHCA Form 3020-0001

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

TITLE

(X8) DATE

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A 033	Continued From page 1 Findings as follow: On at 8:00 Am Resident #3 was observed lying in bed with a physical (full bedrail's) attached to the bed in an upright position. The resident could not put the bed rails down. Review of Resident #3's record showed there was no written care plan developed for the use of the physical (full bedrail's) in file. On at 10:35 AM the Administrator stated she would complete a bedrail's plan of care for Resident #3. Class III	A 033			