

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11910478	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED R-C 10/12/2023
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NAME OF PROVIDER OR SUPPLIER

STREET ADDRESS, CITY, STATE, ZIP CODE

ELANCE AT PASADENA

**1255 PASADENA AVENUE, S.
SAINT PETERSBURG, FL 33707**

(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 second revisit to a complaint investigation (# 2023008736 and #2023010690) was conducted on at Elance at Pasadena. Deficiencies were identified at the time of the visit.	{A 000}		
{A 056} SS=D	59A-36.008(7) FAC Medication - Labeling and Orders (7) MEDICATION LABELING AND ORDERS. (a) The facility may not store prescription drugs for self-administration, assistance with self-administration, or administration unless they are properly labeled and dispensed in accordance with Chapters 465 and 499, F.S., and Rule 64B16-28.108, F.A.C. If a customized patient medication package is prepared for a resident, and separated into individual medicinal drug containers, then the following information must be recorded on each individual container: 1. The resident's name; and, 2. The identification of each medicinal drug in the container. (b) Except with respect to the use of pill organizers as described in subsection (2), no individual other than a pharmacist may transfer medications from one storage container to another. (c) If the directions for use are "as needed" or "as directed," the health care provider must be contacted and requested to provide revised instructions. For an "as needed" prescription, the circumstances under which it would be appropriate for the resident to request the medication and any limitations must be specified; for example, "as needed for , . . . not to exceed 4 tablets per day." The revised instructions, including the date they were obtained from the health care provider and the signature of the staff	{A 056}		

AHCA Form 3020-0001

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

TITLE

(X6) DATE

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{A 056}	Continued From page 1 who obtained them, must be noted in the medication record, or a revised label must be obtained from the pharmacist. (d) Any change in directions for use of a medication that the facility is administering or providing assistance with self-administration must be accompanied by a written, faxed, or electronic copy of a medication order issued and signed by the resident's health care provider. The new directions must promptly be recorded in the resident's medication observation record. The facility may then obtain a revised label from the pharmacist or place an "alert" label on the medication container that directs staff to examine the revised directions for use in the medication observation record. (e) A nurse may take a medication order by telephone. Such order must be promptly documented in the resident's medication observation record. The facility must obtain a written medication order from the health care provider within 10 working days. A faxed or electronic copy of a signed order is acceptable. (f) The facility must make every reasonable effort to ensure that prescriptions for residents who receive assistance with self-administration of medication or medication administration are filled or refilled in a timely manner. (g) Pursuant to Section 465.0276(5), F.S., and Rule 61N-1.006, F.A.C., sample or complimentary prescription drugs that are dispensed by a health care provider, must be kept in their original manufacturer's packaging, which must include the practitioner's name, the resident's name for whom they were dispensed, and the date they were dispensed. If the sample or complimentary prescription drugs are not dispensed in the manufacturer's labeled package, they must be kept in a container that bears a	{A 056}	

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{A 056}	Continued From page 2 label containing the following: 1. Practitioner's name, 2. Resident's name, 3. Date dispensed, 4. Name and strength of the drug, 5. Directions for use; and, 6. Expiration date. (h) Pursuant to Section 465.0276(2)(c), F.S., before dispensing any sample or complimentary prescription drug, the resident's health care provider must provide the resident with a written prescription, or a faxed or electronic copy of such order. This Statute or Rule is not met as evidenced by: Based on record review and interview, the facility failed to provide prescribed medications as ordered from residents' health care providers for one Residents (Resident #1) of the one sampled resident that required assistance with self-administration of medications. Findings Included: During observation of assistance with administration of medication pass on at 10:21 am with Staff A, Medication technician, Staff A assisted Resident # 1 with 0.4 milligram capsule in the morning at 10:21 AM. A record review conducted on of medication observation record for resident #1 revealed instructions for 0.4 milligrams take one (1) capsule by daily at Night. A record review conducted on at 12:41 p.m. record review of medication policies	{A 056}		

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{A 056}	Continued From page 3 revealed the facility has no process on receiving and verifying medications delivered from pharmacy. An interview conducted on _____ at 12:43 p.m. with Wellness Director revealed Medication technicians are responsible to restock the cart with the medication , when they pass the medication, they are to compare the card to the medication observation record. An interview conducted on _____ at 1:31pm. the Administrator stated we are discussing with pharmacy to stagger medications, and increase staffing to see if that will help. Class III	{A 056}			
{A 058} SS=D	429.42() FS: 58A-5.033(3)(a) FAC Pharmacy & Dietary; Uncorrected Deficiencies 429.42 Pharmacy and dietary services.- (1) Any assisted living facility in which the agency has documented a class I or class II deficiency or uncorrected class III deficiencies regarding medicinal drugs or over-the-counter preparations, including their storage, use, delivery, or administration, or dietary services, or both, during a biennial survey or a monitoring visit or an investigation in response to a complaint, shall, in addition to or as an alternative to any penalties imposed under s. 429.19, be required to employ the consultant services of a licensed pharmacist, a licensed registered nurse, or a registered or licensed dietitian, as applicable. The consultant shall, at a minimum, provide onsite quarterly consultation until the inspection team from the agency determines that such consultation services are no longer required.	{A 058}			

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{A 058}	Continued From page 4 (2) A corrective action plan for deficiencies related to assistance with the self-administration of medication or the administration of medication must be developed and implemented by the facility within 48 hours after notification of such deficiency, or sooner if the deficiency is determined by the agency to be life-threatening. 58A-5.033(3) (3) EMPLOYMENT OF A CONSULTANT. (a) Medication Deficiencies. 1. If a class I, class II, or uncorrected class III deficiency directly relating to facility medication practices as established in Rule 58A-5.0185, F.A.C., is documented by the agency pursuant to an inspection of the facility, the agency must notify the facility in writing that the facility must employ or contract the services of a pharmacist licensed pursuant to Section 465.0125, F.S., or registered nurse as determined by the agency. 2. After developing and implementing a corrective action plan in compliance with Section 429.42(2), F.S., the initial on-site consultant visit must take place within 7 working days of the notice of the class I or II deficiency and within 14 working days of the notice of an uncorrected class III deficiency. The facility must have available for review by the agency a copy of the license of the consultant pharmacist or registered nurse and the consultant's signed and dated review of the corrective action plan no later than 10 working days subsequent to the initial on-site consultant visit. 3. The facility must provide the agency with, at a minimum, quarterly on-site corrective action plan updates until the agency determines after written notification by the consultant and facility administrator that deficiencies are corrected and staff has been trained to ensure that proper	{A 058}			

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{A 058}	Continued From page 5 medication standards are followed and that such consultant services are no longer required. The agency must provide the facility with written notification of such determination. This Statute or Rule is not met as evidenced by: Based on record review and interview, the facility failed to ensure all medication concerns were corrected. (Cross Reference: A0056) Findings Included: Due to the uncorrected Class III deficiencies concerning the facility's medication practices, the facility shall be required to employ or contract with a licensed Pharmacist or Registered Nurse Consultant. The Consultant shall at a minimum, provide onsite quarterly consultation until the inspection team from the agency determines that such consultation services are no longer required. This written statement serves as notification of the above stated requirement. During an interview with the Administrator, conducted on 10/27/2023 at 1:19 p.m., the Administrator verbalized their understanding that the facility was required to employ or contract with a licensed Pharmacist and stated they are working with a pharmacy and assistance director of clinical services to reach compliance. Class III	{A 058}			