

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11965917	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED R 11/19/2025
NAME OF PROVIDER OR SUPPLIER CABOT COVE ASSISTED LIVING LLC			STREET ADDRESS, CITY, STATE, ZIP CODE 455 BELCHER RD S LARGO, FL 33771		
(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 fourth revisit to a Biennial survey from in conjunction with a complaint investigation (#2025016260) was conducted at Cabot Cove Assisted Living Facility on . The deficiencies were not found to be corrected at the time of survey. Additional deficiencies were also identified at the time of survey.	{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	{A 056}			

AHCA Form 3020-0001

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

TITLE

(X8) DATE

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{A 056}	Continued From page 1 tablets per day." The revised instructions, including the date they were obtained from the health care provider and the signature of the staff 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	{A 056}			

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{A 056}	Continued From page 2 or complimentary prescription drugs are not dispensed in the manufacturer's labeled package, they must be kept in a container that bears a 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 ensure medications are refilled in a timely manner for residents who received assistance with self-administration of medication for five residents (Resident #1, Resident #2, Resident #3, Resident #7, and Resident #8) of five sampled residents. Findings included: During a medication pass observation for Resident #1, with Staff A (unlicensed Medication Technician) conducted on _____ at 9:03 a.m., Resident #1 had several medications that were not available and were not given to Resident #1 including _____ tablet 40 mg (milligram), _____ tablet 50 mg, and Jardiance tablet 25 mg. During a medication pass observation for Resident #2, with Staff A conducted on _____	{A 056}			

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STREET ADDRESS, CITY, STATE, ZIP CODE

CABOT COVE ASSISTED LIVING LLC

**455 BELCHER RD S
LARGO, FL 33771**

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{A 056}	Continued From page 3 at 9:09 a.m., Resident #2 had several medications that were not available and were not given to Resident #2 including Suspension 0.5mg/2 and Spray 50 mcg (microgram). A Medication Observation Review (MORs) review for Resident #1, conducted on revealed that Resident #1 did not receive the following medications due to not being available: - capsule mg on , and - tablet 40 mg on , and - tablet 50 mg on , and - Jardiance tablet 25 mg on , and - tablet 100 mg on , and - tablet 10 mg on , and - tablet 100 mg on , and - D3 capsule 125 mcg (microgram) on , and	{A 056}		

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{A 056}	Continued From page 4 A Medication Observation Review (MORs) review for Resident #2, conducted on _____, revealed that Resident #2 did not receive the following medications due to not being available: - _____ Extended-Release tablet 500 mg on _____, and _____ - _____ tablet 200 mcg on _____, and _____ - _____ Suspension 0.5 mg/2 on _____, and _____ - _____ Spray 50 mcg on _____, and _____ A Medication Observation Review (MORs) review for Resident #3, conducted on _____, revealed that Resident #3 did not receive the following medications due to not being available: - _____ tablet 88 mcg on _____, and _____ - Polyethylene _____ Powder 3350 on _____, and _____ - _____ tablet 20 mg on _____ A Medication Observation Review (MORs) review for Resident #7, conducted on _____, revealed that Resident #7 did not receive the following medications due to not being available: - _____ tablet 0.5 mg on _____, and _____ - _____ D capsule 5000 unit on _____, and _____ A Medication Observation Review (MORs) review for Resident #8, conducted on _____, revealed that Resident #8 did not receive the following medications due to not being available:	{A 056}		

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{A 056}	Continued From page 5 - tablet 100 mg on _____, and During an interview on _____ at 11:30 a.m., with the Administrative Assistant, conducted the Administrative Assistant agreed the MOR's show that the medications for five Residents #1, #2, #3, #7 and #8 did not receive medications due to not being available. (Photographic evidence obtained) 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. (2) A corrective action plan for deficiencies related to assistance with the self-administration of medication or the administration of medication	A 058			

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A 058	Continued From page 6 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 medication standards are followed and that such consultant services are no longer required. The agency must provide the facility with written	A 058			

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A 058	Continued From page 7 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 documented Class III deficiencies regarding medicinal drugs were corrected as required (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 the consultation services of a licensed Pharmacist Consultant or a Registered Nurse Consultant. The Consultant shall, at a minimum, provide quarterly onsite 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 Administrative Assistant, conducted on at 11:30 a.m., the Administrative Assistant verbalized understanding that the facility was required to employ or contract with a licensed Pharmacist Consultant or Registered Nurse Consultant due to the facility's uncorrected medication Class III deficiencies. Class III	A 058			