Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	AL11967208	A. BUILDING: _____ B. WING _____	R 06/22/2026

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
VENETIAN ISLES ALF	3003 SW 156 PLACE MIAMI, FL 33185

(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 revisit to a re-licensure survey was conducted at Venetian Isles ALF on . Deficiencies were found to be uncorrected at the time of the survey.	{A 000}		
{A 054} SS=D	59A-36.008(5) FAC Medication - Records (5) MEDICATION RECORDS. (a) For residents who use a pill organizer managed in subsection (2), the facility must keep either the original labeled medication container; or a medication listing with the prescription number, the name and address of the issuing pharmacy, the health care provider's name, the resident's name, the date dispensed, the name and strength of the drug, and the directions for use. (b) The facility must maintain a daily medication observation record for each resident who receives assistance with self-administration of medications or medication administration. A medication observation record must be immediately updated each time the medication is offered or administered and include: 1. The name of the resident and any known the resident may have; 2. The name of the resident's health care provider and the health care provider's telephone number; 3. The name, strength, and directions for use of each medication; and, 4. A chart for recording each time the medication is taken, any missed dosages, refusals to take medication as prescribed, or medication errors. (c) For medications that serve as chemical , the facility must, pursuant to Section 429.41, F.S., maintain a record of the prescribing physician's annual evaluation of the use of the medication.	{A 054}		

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{A 054}	Continued From page 1 This Statute or Rule is not met as evidenced by: Based on interview and record review the facility failed to maintain an accurate medication observation record (MOR) that specified the direction of use of each medication for two out of five (Residents #4 and 5) sampled residents. Findings Included: A review of Resident #4's MOR revealed that the medication was documented with the administration time of "bedtime." The review also revealed that the medication had a blank entry for the scheduled dose on . A review of Resident #5's MOR revealed that the medication was documented with the administration time of "AM." Interviewed on at 11:26 AM the Administrator acknowledged and confirmed that the MOR's for Residents #4 and 5 had been complete by and included the handwritten entries "bedtime" and "AM" for the medication administration times. Class III	{A 054}		
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	A 056		

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A 056	Continued From page 2 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 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, faxes, 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	A 056		

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A 056	Continued From page 3 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 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:	A 056		

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A 056	Continued From page 4 Based on interview and record review the facility failed to ensure that medication observation records (MOR) were maintained by the pharmacist in accordance with medication labeling and order requirements for one out of five sampled residents (Resident #5). Findings Included: A review of Resident 5's Medication Observation Log (MOR) revealed a facility created handwritten medication list. The document was not provided by the pharmacy and did not contain pharmacy generated medical information. Interviewed on at 11:26 AM the Administrator acknowledged and confirmed that the MOR was handwritten by the facility. Class III	A 056		
A 058 SS=D	429.42( ) FS; 59A-36.023(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	A 058		

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A 058	Continued From page 5 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 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. 59A-36.023(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 59A-36.008, F.A.C., is documented by agency personnel 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 a class I or class 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	A 058		

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A 058	Continued From page 6 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 notification of such determination. This Statute or Rule is not met as evidenced by: Based on interview and record review the facility is required to employ consultant services of a licensed registered nurse, or pharmacist who will provide onsite quarterly consultations until it is determined that such consultation services are no longer required. The facility received a Class III uncorrected deficiency of Medication-Records. Findings Included: Uncorrected Deficiency A review of Resident #4's MOR revealed that the medication was documented with the administration time of "bedtime." The review also revealed that the medication had a blank entry for the scheduled dose on A review of Resident #5's MOR revealed that the medication was documented with the administration time of "AM." Interviewed on at 11:26 AM the Administrator acknowledged and confirmed that the MOR's for Residents #4 and #5 had been complete by and included the handwritten entries "bedtime" and "AM" for the medication administration times.	A 058		

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A 058	Continued From page 7 Class III	A 058		
{A 162} SS=D	59A-36.015(3) FAC Records - Resident (3) RESIDENT RECORDS. Resident records must be maintained on the premises and include: (a) Resident demographic data as follows: 1. Name, 2. , 3. Race, 4. Date of birth, 5. Place of birth, if known, 6. Social security number, 7. Medicaid and/or Medicare number, or name of other health insurance , 8. Name, address, and telephone number of next of kin, legal representative, or individual designated by the resident for notification in case of an emergency; and, 9. Name, address, and telephone number of the resident's health care practioner and case manager, if applicable. (b) A copy of the Resident Health Assessment form, AHCA Form 1823 or the health care practitioner's medical examination form described in Rule 59A-36.006, F.A.C. (c) Any orders for medications, nursing services, therapeutic diets, , or other services to be provided, supervised, or implemented by the facility that require a health care provider's order. (d) Documentation of a resident's refusal of a therapeutic diet pursuant to Rule 59A-36.012, F.A.C., if applicable. (e) The resident care record described in paragraph 59A-36.007(1)(f), F.A.C. (f) A record that is initiated on admission. Information may be taken from AHCA Form 1823	{A 162}		

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{A 162}	Continued From page 8 or the resident's health assessment. Residents receiving assistance with the activities of daily living must have their recorded semi-annually. This subsection does not apply to residents who are receiving licensed hospice services when such residents, their representatives, or their physicians request in writing that not be taken. (g) For facilities that will have unlicensed staff assisting the resident with the self-administration of medication, a copy of the written informed consent described in Rule 59A-36.006, F.A.C., if such consent is not included in the resident's contract. (h) For facilities that manage a pill organizer, assist with self-administration of medications or administer medications for a resident, copies of the required medication records maintained pursuant to Rule 59A-36.008, F.A.C. (i) A copy of the resident's contract with the facility, including any addendums to the contract as described in Rule 59A-36.018, F.A.C. (j) For a facility whose owner, administrator, staff, or representative thereof, serves as an attorney in fact for a resident, a copy of the monthly written statement of any transaction made on behalf of the resident as required in Section 429.27, F.S. (k) For any facility that maintains a separate trust fund to receive funds or other property belonging to or due a resident, a copy of the quarterly written statement of funds or other property disbursed as required in Section 429.27, F.S. (l) If the resident is an recipient, a copy of the Department of Children and Families form Alternate Care Certification for , ( ), CF-ES 1006, , which is hereby incorporated by reference and available for review at: http://www.flrules.org/Gateway/reference.asp?No	{A 162}		

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{A 162}	Continued From page 9 =Ref-04004. The absence of this form will not be the basis for administrative action against a facility if the facility can demonstrate that it has made a good faith effort to obtain the required documentation from the Department of Children and Families. (m) Documentation of the , , of a health care surrogate, health care proxy, guardian, or the existence of a power of attorney, where applicable. (n) For hospice patients, the interdisciplinary care plan and other documentation that the resident is a hospice patient as required in Rule 59A-36.006, F.A.C. (o) The resident's , DH Form 1896, if applicable. (p) For independent living residents who receive meals and occupy beds included within the licensed capacity of an assisted living facility, but who are not receiving any personal, limited nursing, or extended congregate care services, record keeping may be limited to a log listing the names of residents participating in this arrangement. (q) Except for resident contracts, which must be retained for 5 years, all resident records must be retained for 2 years following the departure of a resident from the facility unless it is required by contract to retain the records for a longer period of time. Upon request, residents must be provided with a copy of their records upon departure from the facility. (r) Additional resident records requirements for facilities holding a limited mental health, extended congregate care, or limited nursing services license are provided in Rules 59A-36.020, 59A-36.021 and 59A-36.022, F.A.C., respectively. This Statute or Rule is not met as evidenced by: Based on interview and record review the facility	{A 162}		

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{A 162}	Continued From page 10 failed to maintain an accurate and up-to-date health assessment for two out of five sampled residents (Residents #2 and 3). Findings Included: 1) A review of Resident #2's record showed she was admitted to the facility on and had a diagnosis of . The health assessment dated showed the resident required no added salt, low fat/ low diet. A review of Resident #2's hospice record revealed the physician ordered a soft diet on . 2) A review of Resident #3's record showed she was admitted to the facility on and had a diagnosis of , Type 2 , and . The health assessment dated showed the resident required a calorie-controlled diet. A review of Resident #3's hospice record revealed the physician ordered a mechanical soft diet on . During an interview on at 11:26 AM the Administrator stated he would contact Resident #2' s and 3's doctor and hospice services to discuss the residents' diets. Class III	{A 162}		