

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11968706	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED R 10/20/2020
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NAME OF PROVIDER OR SUPPLIER

STREET ADDRESS, CITY, STATE, ZIP CODE

ENCORE AT AVALON PARK

**13798 CYGNUS DR
ORLANDO, FL 32828**

(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 complaint investigation # 2020014570 was conducted at Encore At Avalon Park on . A deficiency was 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 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	{A 056}		

AHCA Form 3020-0001

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

TITLE

(X6) DATE

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NAME OF PROVIDER OR SUPPLIER ENCORE AT AVALON PARK			STREET ADDRESS, CITY, STATE, ZIP CODE 13798 CYGNUS DR ORLANDO, FL 32828		
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{A 056}	Continued From page 1 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 label containing the following:	{A 056}			

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{A 056}	Continued From page 2 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: DEFICIENCY REMAINED UNCORRECTED Based on observations, record review and interview the facility failed to ensure a revised label from the pharmacist or an "alert" label was placed on the medication container that directed staff to examine the revised directions for use in the medication observation record, when the directions for use changed for 1 of sampled residents. Findings: Resident #1 had a health assessment dated that listed high and Prescription order dated indicated 1 mg at bedtime and 0.5 mg one twice a day. The Medication Observation Record (MOR) listed 1 mg at bedtime and was marked given as ordered. Review of the resident's medications revealed a prescription labels dated and that indicated 1 mg three times daily as needed for agitation. Continued	{A 056}			

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{A 056}	Continued From page 3 observations revealed no evidence that a revised label from the pharmacist or an "alert" label was placed on the medication container that directed staff to examine the revised directions for use in the medication observation record. In an interview with the nurse on at 2:30 PM who said she knew the resident took 1 mg at bedtime. Class III /	{A 056}		