

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11967590	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED
NAME OF PROVIDER OR SUPPLIER A SECOND HOME OF RIVIERA BEACH INC		STREET ADDRESS, CITY, STATE, ZIP CODE 120 WEST 22ND STREET RIVIERA BEACH, FL 33404			
(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 An Assisted Living Facility standard biennial licensure survey was conducted on A Second Home of Riviera Beach, had a licensure deficiency found at the time of the visit.	A 000			
A 056 SS=D	58A-5.0185(7) FAC Medication - Labeling and Orders (7) MEDICATION LABELING AND ORDERS. (a) No prescription drug shall be kept or administered by the facility, including assistance with self-administration of medication, unless it is 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. Identification of each medicinal drug product in the container. (b) Except with respect to the use of pill organizers as described in subsection (2), no person 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 shall 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 shall be specified; for example, "as needed for pain, 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, shall be noted in the	A 056			

AHCA Form 3020-0001

TITLE

(X6) DATE

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

STATE FORM

6899

5H9611

If continuation sheet 1 of 3

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A 056	Continued From page 1 medication record, or a revised label shall be obtained from the pharmacist. (d) Any change in directions for use of a medication for which the facility is providing assistance with self-administration or administering medication must be accompanied by a written medication order issued and signed by the resident ' s health care provider, or a faxed copy of such order. The new directions shall promptly be recorded in the resident ' s medication observation record. The facility may then place an " alert " label on the medication container which directs staff to examine the revised directions for use in the MOR, or obtain a revised label from the pharmacist. (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 copy of a signed order is acceptable. (f) The facility shall 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 64F-12.006, 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 shall also 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 shall be kept in a container that bears a label containing the following: 1. Practitioner ' s name;	A 056		

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A 056	Continued From page 2 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 shall provide the resident with a written prescription, or a fax copy of such order. This Statute or Rule is not met as evidenced by: Based on record review and an interview, the facility failed to request specific instructions for a medication that staff assisted a resident with that had instructions to give "as needed" for 1 out of 1 sampled resident (Resident #1). The findings include: On _____ at 1 PM, Resident #1's _____ 2013 MORs (medication observation record) was reviewed. _____ eye ointment was listed to be given "as needed" but did not indicate for what. The label was read and indicated the same. The Administrator was interviewed at this time and stated she gives the resident this medication once a day and a home health nurse administers it to her the second time, so she is providing the ointment twice a day. She was unable to state during interview what the medication was for, only that it was for treatment following a procedure on the resident's skin above her eye. She had not contacted the health care provider to obtain specific instructions to follow for this "PRN/as needed" medication. Class III	A 056			

RICK SCOTT
GOVERNOR



ELIZABETH DUDEK
SECRETARY

2013

Administrator
A Second Home Of Riviera Beach Inc
120 West 22nd Street
Riviera Beach, FL 33404

Dear Administrator:

This letter reports the findings of a state licensure survey that was conducted on 2013 by a representative from this office.

Attached is the provider's copy of the State (3020) Form, which indicates the deficiencies that were identified on the day of the visit. Section 408.811(4), Florida Statutes, requires that you correct these deficiencies within **thirty (30) days** of the date of this letter unless the Agency has approved another timeframe. Staff from this office will conduct a review after 2013 to verify that the necessary corrections are in place to correct the deficiencies identified on your survey.

The Quality Assurance Questionnaire has long been employed to obtain your feedback following survey activity. This form has been placed on the Agency's website at <http://ahca.myflorida.com/Publications/Forms.shtml> as a first step in providing a web-based interactive consumer satisfaction survey system. You may access the questionnaire through the link under Health Facilities and Providers on this page. Your feedback is encouraged and valued, as our goal is to ensure the professional and consistent application of the survey process.

Thank you for the assistance provided to the representative. Should you have any questions please call this office at (561) 381-5840.

Sincerely,


Arlene Mayo-Davis
Field Office Manager

AMD/dso
Enclosure
XG90

