

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11966019	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED
NAME OF PROVIDER OR SUPPLIER MARIA'S HOME		STREET ADDRESS, CITY, STATE, ZIP CODE 16131 SW 104 COURT MIAMI, FL 33157		
(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 Standard Biennial survey was conducted on Maria's Home Assisted Living Facility had deficiencies found at the time of the visit.	A 000		
A 055 SS=D	58A-5.0185(6) FAC Medication - Storage and Disposal (6) MEDICATION STORAGE AND DISPOSAL. (a) In order to accommodate the needs and preferences of residents and to encourage residents to remain as independent as possible, residents may keep their medications, both prescription and over-the-counter, in their possession both on or off the facility premises; or in their _____ or apartments, which must be kept locked when residents are absent, unless the medication is in a secure place within the or apartments or in some other secure place which is out of sight of other residents. However, both prescription and over-the-counter medications for residents shall be centrally stored if: 1. The facility administers the medication; 2. The resident requests central storage. The facility shall maintain a list of all medications being stored pursuant to such a request; 3. The medication is determined and documented by the health care provider to be hazardous if kept in the personal possession of the person for whom it is prescribed; 4. The resident fails to maintain the medication in a safe manner as described in this paragraph; 5. The facility determines that because of physical arrangements and the conditions or habits of residents, the personal possession of medication by a resident poses a safety hazard to other residents; or 6. The facility's rules and regulations require central storage of medication and that policy has	A 055		

AHCA Form 3020-0001

TITLE

(X6) DATE

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

STATE FORM

0099

95T011

If continuation sheet 1 of 6

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A 055	Continued From page 1 been provided to the resident prior to admission as required under Rule 58A-5.0181, F.A.C. (b) Centrally stored medications must be: 1. Kept in a locked cabinet, locked cart, or other locked storage receptacle, _____, or area at all times; 2. Located in an area free of dampness and abnormal temperature, except that a medication requiring refrigeration shall be refrigerated. Refrigerated medications shall be secured by being kept in a locked container within the refrigerator, by keeping the refrigerator locked, or by keeping the area in which refrigerator is located locked; 3. Accessible to staff responsible for filling pill-organizers, assisting with self-administration, or administering medication. Such staff must have ready access to keys to the medication storage areas at all times; and 4. Kept separately from the medications of other residents and properly closed or sealed. (c) Medication which has been discontinued but which has not expired shall be returned to the resident or the resident's representative, as appropriate, or may be centrally stored by the facility for future resident use by the resident at the resident's request. If centrally stored by the facility, it shall be stored separately from medication in current use, and the area in which it is stored shall be marked "discontinued medication." Such medication may be reused if re-prescribed by the resident's health care provider. (d) When a resident's stay in the facility has ended, the administrator shall return all medications to the resident, the resident's family, or the resident's guardian unless otherwise prohibited by law. If, after notification and waiting at least 15 days, the resident's medications are still at the facility, the	A 055		

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A 055	Continued From page 2 medications shall be considered abandoned and may disposed of in accordance with paragraph (e). (e) Medications which have been abandoned or which have expired must be disposed of within 30 days of being determined abandoned or expired and disposition shall be documented in the resident's record. The medication may be taken to a pharmacist for disposal or may be destroyed by the administrator or designee with one witness. (f) Facilities that hold a Special-ALF permit issued by the Board of Pharmacy may return dispensed medicinal drugs to the dispensing pharmacy pursuant to Rule 64B16-28.870, F.A.C. This Statute or Rule is not met as evidenced by: Based on observations and interview the facility failed to ensure medications which have expired must be disposed of within 30 days of being determined expired. The findings Include: During medication reconciliation on the following medications belonging to resident #2 was found: - 50 mg had an expiration date of and Triamt/HCTZ had an expiration date of _____ Interview conducted with the administrator on at approximately 1:37 p.m. acknowledged the findings. Class III		A 055		
A 056 SS=D	58A-5.0185(7) FAC Medication - Labeling and Orders		A 056		
(7) MEDICATION LABELING AND ORDERS.					

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A 056	Continued From page 3 (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 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	A 056		

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A 056	Continued From page 4 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; 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.	A 056		

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A 056	Continued From page 5 This Statute or Rule is not met as evidenced by: Based on record review, observation, and interview, the facility failed to ensure that prescription drugs were properly labeled for 1 out of 2 sampled residents (Resident #2). The findings include: Resident #2's Medication Observation Record (MOR) revealed that the medications 0.125mg take one tablet by mouth daily, 1 mg take one tablet by mouth daily and CD HR take one capsule by mouth daily had been provided to Resident #2 by the facility. During the medication reconciliation the prescriptions labels revealed that 0.125 mg must have been finished on 1 mg must have been finished on and Diltiazem must have been finished on, however the bottles had a lot of pills. Interview conducted with the administrator on at approximately 1:30 p.m. revealed that was unaware that the prescription labels were not correct. He stated that he will contact Resident's daughter to review all the resident's #2 medications. Class III		A 056		



RICK SCOTT
GOVERNOR

Better Health Care for all Floridians

ELIZABETH DUDEK
SECRETARY

_____, 2012

Administrator
Maria's Home
16131 Sw 104 Court
Miami, FL 33157

Dear Administrator:

This letter reports the findings of a state licensure survey that was conducted on _____, 2012 by representative(s) of 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 days of the date of this letter unless the Agency has approved another timeframe.

Staff from this office will conduct a review after _____ 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 surveyor(s). Should you have any questions please call Kristal Branton, Health Facility Evaluator Supervisor at (305) 593-3100.

Sincerely,

for Arlene Mayo-Davis
Field Office Manager

AMD:sy
Enclosure
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