

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11967434	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 09/20/2023
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NAME OF PROVIDER OR SUPPLIER

STREET ADDRESS, CITY, STATE, ZIP CODE

ORLANDO LUTHERAN TOWERS

**404 MARIPOSA STREET
ORLANDO, FL 32801**

(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 complaint investigation #2023009633 was conducted at Orlando Lutheran Towers on . The facility had deficiencies at the time of the survey.	A 000		
A 025 SS=D	429.26(7) FS; 59A-36.007(1) FAC Resident Care - Supervision 429.26 (7) The facility shall notify a licensed physician when a resident exhibits signs of or or has a change of condition in order to rule out the presence of an underlying physiological condition that may be contributing to such or . The notification must occur within 30 days after the acknowledgment of such signs by facility staff. If an underlying condition is determined to exist, the facility must notify the resident's representative or designee of the need for health care services and must assist in making , for the necessary care and services to treat the condition. If the resident does not have a representative or designee or if the resident's representative or designee cannot be located or is unresponsive, the facility shall arrange with the appropriate health care provider for the necessary care and services to treat the condition. 59A-36.007 Resident Care Standards. An assisted living facility must provide care and services appropriate to the needs of residents accepted for admission to the facility. (1) SUPERVISION. Facilities must offer personal supervision as appropriate for each resident, including the following: (a) Monitoring of the quantity and quality of resident diets in accordance with Rule	A 025		

AHCA Form 3020-0001

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

TITLE

(X6) DATE

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A 025	Continued From page 1 59A-36.012, F.A.C. (b) Daily observation by designated staff of the activities of the resident while on the premises, and awareness of the general health, safety, and physical and emotional well-being of the resident. (c) Maintaining a general awareness of the resident's whereabouts. The resident may travel independently in the community. (d) Contacting the resident's health care provider and other appropriate party such as the resident's family, guardian, health care surrogate, or case manager if the resident exhibits a significant change. (e) Contacting the resident's family, guardian, health care surrogate, or case manager if the resident is discharged or moves out. (f) Maintaining a written record, updated as needed, of any significant changes, any illnesses that resulted in medical attention, changes in the method of medication administration, or other changes that resulted in the provision of additional services. This Statute or Rule is not met as evidenced by: Based on record reviews and interviews, the facility failed to ensure a medication was held based on parameters set by a health care provider for 1 of 3 sampled residents (#1). Findings: A review of resident #1's record revealed a facility admission date of . The resident's medical history included . The resident required assistance with self-administration of medications. A review of the resident's . medication administration record (MAR) revealed the	A 025		

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A 025	Continued From page 2 following entries: CD ER 24-hour capsule 120 milligram (mg) 1 capsule by one time daily for Hold for () <105 or rate () < . 25 mg tablet, 1 tablet by daily for blocker. Hold for < . /or <110 and call the doctor. On at 6 am the resident's was 96 and the was signed as given by nurse E. On at 9am the resident's was 108 and the was signed as given by nurse F. A review of the resident's MAR revealed the following: On at 9am the resident's was 109 and the was signed as given by nurse G. On at 9am the resident's was 107 and the was signed as given by nurse F. There was no documentation on either the MARs or in the resident's record to explain the medications being given despite the B/P readings and doctor notification. On at 3:15 pm, nurse D provided a "24-hour" report, which was not part of the resident's medical record, and said she documented a re-check (B/P) of on it. She then said the result of the	A 025		

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A 025	Continued From page 3 re-check was why she gave the She said she made a mistake by not documenting the new B/P on either the MAR or in the medical record. On at 3:30 pm, the director of nursing said she could not comment on nurse G giving the on because the nurse no longer worked for the facility. On at 3:50 pm, nurse E said the resident's B/P on was "probably a typo. His B/P is never low. That's 11pm-7am. I get to him last and by the time I get to him I'm half asleep. I start on the 6th floor and work my way down". Class III	A 025			
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	A 056			

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A 056	Continued From page 4 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, 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	A 056			

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A 056	Continued From page 5 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: Based on record reviews and interviews, the facility failed to ensure a boost supplement drink was refilled in a timely manner to ensure the resident had enough for 1 of 3 sampled residents (#1). Findings: A review of resident #1's 1823 revealed the resident required assistance with self-administration of medications.	A 056			

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A 056	Continued From page 6 On _____ a health care provider ordered Boost supplement drink three times per day. On _____ and _____ the Boost was documented as not given because "awaiting on delivery". On _____ at 2:45 pm, nurse D said the boost was ordered by the facility from a pharmacy. On _____ at 3:20 pm, the finding was discussed with the facility's manager, who said she would look into it and let the surveyor know. On _____ at 3:30 pm, the finding was discussed with the director of nursing. No documentation was provided to demonstrate the Boost was ordered in a timely manner to prevent running out. Class III	A 056		