

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11970392	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED R-C 07/30/2025
NAME OF PROVIDER OR SUPPLIER HEARTHSTONE NONA		STREET ADDRESS, CITY, STATE, ZIP CODE 10298 SAVANNAH PARK DR ORLANDO, FL 32832			
(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
{CZ821}	59A-35.110() FAC Reporting Requirements; Electronic Submission 59A-35.110 Reporting Requirements; Electronic Submission. (1) During the two year licensure period, any change or expiration of any information that is required to be reported under Chapter 408, Part II, F.S., or authorizing statutes for the provider type as specified in Section 408.803(3), F.S., during the license application process must be reported to the Agency within 21 days of occurrence of the change, including: (a) Insurance coverage renewal; (b) Bond renewal; (c) Change of administrator or the similarly titled person who is responsible for the day-to-day operation of the provider; (d) Annual sanitation inspections; (e) Fire inspections. (2) Electronic submission of information. (a) The following required information must be submitted electronically through the Agency's Single Sign On Portal located at https://apps.ahca.myflorida.com/SingleSignOnPo rtal. 1. Nursing homes: Adverse incident reports must be submitted electronically to the Agency within 15 calendar days after the occurrence of the incident as required in Section 400.147, F.S. on Nursing Home Adverse Incident, AHCA Form 3110-0010 OL, , which is hereby incorporated by reference and available at: https://www.flrules.org/Gateway/reference.asp?N o=Ref-08777, and through the Agency's adverse incident reporting system which can only be accessed through the Agency's Single Sign On Portal located at: https://apps.ahca.myflorida.com/SingleSignOnPo rtal.	{CZ821}			

AHCA Form 3020-0001

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

TITLE

(X8) DATE

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{CZ821}	Continued From page 1 2. Assisted living facilities: Adverse incident reports must be submitted electronically to the Agency within 1 business day after the occurrence of the incident, and within 15 calendar days after the occurrence of the incident as required in Section 429.23, F.S., on Assisted Living Facility Adverse Incident, AHCA Form 3180-1025 OL, , which is hereby incorporated by reference and available at: https://www.flrules.org/Gateway/reference.asp?No=Ref-08778, and through the Agency's adverse incident reporting system which can only be accessed through the Agency's Single Sign On Portal located at: https://apps.ahca.myflorida.com/SingleSignOnPortal. 3. Hospitals: Adverse incident reports must be submitted electronically to the Agency within 15 calendar days after the occurrence of the incident as required in Section 395.0197, F.S., on Hospital Adverse Incident, AHCA Form 3140-5001 OL, , which is hereby incorporated by reference and available at: https://www.flrules.org/Gateway/reference.asp?No=Ref-08779, and through the Agency's adverse incident reporting system which can only be accessed through the Agency's Single Sign On Portal located at: https://apps.ahca.myflorida.com/SingleSignOnPortal. 4. Ambulatory Surgical Centers: Adverse incident reports must be submitted electronically to the Agency within 15 calendar days after the occurrence of the incident as required in Section 395.0197, F.S., on Ambulatory Surgical Center Adverse Incident, AHCA Form 3140-5004 OL, , which is hereby incorporated by reference and available	{CZ821}			

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{CZ821}	Continued From page 2 at: https://www.flrules.org/Gateway/reference.asp?No=Ref-08780 , and through the Agency's adverse incident reporting system which can only be accessed through the Agency's Single Sign On Portal located at: https://apps.ahca.myflorida.com/SingleSignOnPortal . 5. Hospitals and ambulatory surgical centers must submit annual reports pursuant to Section 395.0197 F.S., electronically to the Agency on Annual Report, AHCA Form 3140-5005 OL, , which is hereby incorporated by reference and available at: http://www.flrules.org/Gateway/reference.asp?No=Ref-12123 , and through the Agency's Single Sign On Portal located at: https://apps.ahca.myflorida.com/SingleSignOnPortal . 6. For the purposes of this rule, the following applies for submitting adverse incident reports: a. A business day means any day other than a Saturday, Sunday, or legal holiday as designated in Section 110.117, F.S. b. A preliminary (1-Day) report is deemed late when submitted more than 1 business day after the day of the incident. c. Nursing Homes, Hospitals, and Ambulatory Surgical Centers: A full report (15- Day) is deemed late when submitted more than 15 calendar days after the day of the incident. d. Assisted Living Facilities: A full report (15-Day) is deemed late when submitted more than 15 calendar days after the day of the incident or more than 3 business days after the Agency issues the reminder pursuant to Section 429.23(5), F.S., whichever is later. e. Assisted Living Facilities and Nursing Homes that submit reports deemed late may be fined up	{CZ821}			

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{CZ821}	Continued From page 3 to \$50 per day late not to exceed \$500. (b) The licensee must retain a copy of all documentation generated at time of reporting as confirmation of successful electronic submission. (c) If the Agency's Single Sign On Portal or the online adverse incident reporting system is temporarily out of service the licensee may contact the Agency directly at 1(888)419-3456 for assistance. Reporting will resume as soon as online access is restored. This Statute or Rule is not met as evidenced by: DEFICIENCY REMAINED UNCORRECTED Based on record reviews and interviews, the facility failed to electronically submit a one-day report and a 15 day (full report) adverse incident report (AIRS) to the agency for 1 of 4 sampled residents (#3) as required. Findings: On at 8:35 AM during the entrance conference, the administrator was asked to provide copies of the incident reports that were required for the revisit survey. On at 10:30 AM a follow up occurred requesting the administrator to provide copies of all adverse incident reports since , along with incident report for resident #3. On at 11:37 AM the director of nursing confirmed what was needed for the adverse reports stating the administrator is working on them now. Review of the adverse incident reports provided did not include the submission report for resident	{CZ821}			

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{CZ821}	Continued From page 4 #3's with injury. On at 3:14 PM the director of nursing stated an adverse incident report was not submitted for resident #3's on the AHCA website. The information was submitted in a secured email and the administrator is working to get it. On at 12:30 PM during the exit conference, no further documentation was provided for resident #3 and the administrator confirmed the findings. Unclassified	{CZ821}			
{A 000}	Initial Comments A revisit to complaint investigation #2025006857 and generator monitoring were conducted at Hearthstone Nona Assisted Living & Memory Care on . The facility had uncorrected 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	{A 025}			

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{A 025}	Continued From page 5 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 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	{A 025}			

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{A 025}	Continued From page 6 changes that resulted in the provision of additional services. This Statute or Rule is not met as evidenced by: DEFICIENCY REMAINED UNCORRECTED Based on record reviews and interview, the facility failed to follow a health care provider's medication order for 1 of 12 sampled residents (#11). Findings: Review of resident #11's health assessment form (1823) revealed the resident needed assistance with self-administration of medication. The resident's medical history included Review of the health care provider visit note revealed 20 milligram (mg) was discontinued in . However, review of the residents' and medication observation records (MORs) revealed the resident was given 20 mg on , , and , and . An order for 40 mg was written on . However, the resident was not given 40 mg from to . The resident had lower extremity , and , for several days. As a result, 40 mg was resumed on . In addition, the resident was given both 40 mg and 20 mg on , and . On at 9:18 AM the findings were	{A 025}			

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{A 025}	Continued From page 7 discussed with the director of nursing (DON), who said there was an issue with the pharmacy when the milligrams were changed. Class III	{A 025}			