

Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: AL11968683	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____		(X3) DATE SURVEY COMPLETED R 10/21/2022
NAME OF PROVIDER OR SUPPLIER BANYAN PLACE			STREET ADDRESS, CITY, STATE, ZIP CODE 2950 NW 5TH AVENUE BOCA RATON, FL 33431		
(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 unannounced Revisit to Complaint (#2022011996) survey was conducted on 10-21-2022 at Banyan Place. A previously cited deficiency was found uncorrected at the time of the survey.	{A 000}			
{A 165}	429.23(1-4 &) FS Risk Mgmt & QA 429.23 Internal risk management and quality assurance program; adverse incidents and reporting requirements.- (1) Every facility licensed under this part may, as part of its administrative functions, voluntarily establish a risk management and quality assurance program, the purpose of which is to assess resident care practices, facility incident reports, deficiencies cited by the agency, adverse incident reports, and resident grievances and develop plans of action to correct and respond quickly to identify quality differences. (2) Every facility licensed under this part is required to maintain adverse incident reports. For purposes of this section, the term, "adverse incident" means: (a) An event over which facility personnel could exercise control rather than as a result of the resident's condition and results in: 1. ; 2. or , damage; 3. Permanent ; 4. or of bones or joints; 5. Any condition that required medical attention to which the resident has not given his or her consent, including failure to honor advanced directives; 6. Any condition that requires the transfer of the resident from the facility to a unit providing more acute care due to the incident rather than the resident's condition before the incident; or	{A 165}			

AHCA Form 3020-0001
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

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{A 165}	Continued From page 1 7. An event that is reported to law enforcement or its personnel for investigation; or (b) Resident elopement, if the elopement places the resident at risk of harm or injury. (3) Licensed facilities shall provide within 1 business day after the occurrence of an adverse incident, through the agency's online portal, or if the portal is offline, by electronic mail, a preliminary report to the agency on all adverse incidents specified under this section. The report must include information regarding the identity of the affected resident, the type of adverse incident, and the status of the facility's investigation of the incident. (4) Licensed facilities shall provide within 15 days, through the agency's online portal, or if the portal is offline, by electronic mail, a full report to the agency on all adverse incidents specified in this section. The report must include the results of the facility's investigation into the adverse incident. (6) , neglect, or , must be reported to the Department of Children and Families as required under chapter 415. (7) The information reported to the agency pursuant to subsection (3) which relates to persons licensed under chapter 458, chapter 459, chapter 461, chapter 464, or chapter 465 shall be reviewed by the agency. The agency shall determine whether any of the incidents potentially involved conduct by a health care professional who is subject to disciplinary action, in which case the provisions of s. 456.073 apply. The agency may investigate, as it deems appropriate, any such incident and prescribe measures that must or may be taken in response to the incident. The agency shall review each incident and determine whether it potentially involved conduct by a health care professional who is subject to	{A 165}			

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{A 165}	Continued From page 2 disciplinary action, in which case the provisions of s. 456.073 apply. (8) If the agency, through its receipt of the adverse incident reports prescribed in this part or through any investigation, has reasonable belief that conduct by a staff member or employee of a licensed facility is grounds for disciplinary action by the appropriate board, the agency shall report this fact to such regulatory board. (9) The adverse incident reports and preliminary adverse incident reports required under this section are confidential as provided by law and are not discoverable or admissible in any civil or administrative action, except in disciplinary proceedings by the agency or appropriate regulatory board. (10) The agency may adopt rules necessary to administer this section. This Statute or Rule is not met as evidenced by: Based on record review and interview the facility failed to timely submit an Adverse Incident Report as required within one business day after the occurrence of an adverse incident involving Resident #2 on . The findings included: Review of the facility records indicated that the facility created and submitted an Adverse Incident Report to the Agency on . Review of this Adverse Incident Report indicated that Resident #2 was injured while being assisted with ambulation in a wheelchair on . . . at approximately 2:00 PM. Review of this report revealed that the resident was wheeled into the dining room awaiting emergency response personnel when she subsequently . again while sitting in a wheelchair in the dining room. Further	{A 165}			

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{A 165}	Continued From page 3 review of this report indicated that the resident required to be transported to the hospital Emergency Room via 911 for evaluation of her / injury due to the 2 she sustained while in the facility. In an interview conducted with the Director of Nursing on at 3:00 PM, she stated that she was not aware and was not informed by the Administrator of any other Adverse Incident Reports which were submitted to the Agency after . She stated that upon learning of the adverse incident that occurred on , it was required to submit a preliminary report to the Agency within 1 business day and she confirmed that this was not done by the facility. Class III	{A 165}			