Agency for Health Care Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	AL11953622	A. BUILDING: _____ B. WING _____	R-C 02/02/2023

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
THE POINTE	9797 BAY PINES BLVD SAINT PETERSBURG, FL 33708

(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 second revisit to survey and 6-Month Survey w/LNS/Generator survey was completed on at The Pointe. The deficiencies were corrected at the time of the survey. There were new deficiencies found at the time of the survey.	{A 000}		
A 165 SS=D	429.23(&) 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	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 resident's condition before the incident; or 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) [illegible], neglect, or [illegible] 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	A 165		

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A 165	Continued From page 2 by a health care professional who is subject to 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 file an adverse incident report (AIRs) within 1 business day of an event over which facility personnel could exercise control, rather than as a result of the resident's condition, resulting in a and admission to a higher level of care or one (Resident #1) of one sampled. Findings include: Record review of the facility's incident reports on , revealed that Resident #1 had on and her , causing her to be admitted to the hospital. (Photo evidence obtained) An attempted record review of the facility's	A 165		

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A 165	Continued From page 3 adverse incident reports on . . . . . , revealed the Administrator was unable to produce an adverse incident report for the incident occurring on with Resident #1. An interview with the Administrator on at 11:15am, confirmed the facility did not file an adverse incident report for the incident occurring on . . . . . with Resident #1. The Administrator stated, "No we did not file one". Class III	A 165		