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
	AL11964856	A. BUILDING: _____ B. WING _____	07/22/2019

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
BROOKDALE TEQUESTA	211 VILLAGE BLVD TEQUESTA, FL 33469

(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 licensure complaint survey, #2019007751, was conducted on [redacted] at Brookdale Tequesta, License #9321. The facility had a deficiency at the time of the survey.	A 000		
A 165 SS=B	429.23(&) FS; 58A-5.0241 FAC Risk Mgmt & QA; Adverse Incident Report 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. [redacted]; 2. or , damage; 3. Permanent [redacted]; 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, by electronic mail, facsimile, or United States 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, by electronic mail, facsimile, or United States 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 disciplinary action, in which case the provisions of s. 456.073 apply.	A 165		

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A 165	Continued From page 2 (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 Department of Elderly Affairs may adopt rules necessary to administer this section. 58A-5.0241 Adverse Incident Report. (1) INITIAL ADVERSE INCIDENT REPORT. The preliminary adverse incident report required by Section 429.23(3), F.S., must be submitted within 1 business day after the incident pursuant to Rule 59A-35.110, F.A.C., which requires online reporting. (2) FULL ADVERSE INCIDENT REPORT. For each adverse incident reported in subsection (1) above, the facility must submit a full report within 15 days of the incident. The full report must be submitted pursuant to Rule 59A-35.110, F.A.C., which requires online reporting. This Statute or Rule is not met as evidenced by: Based on record review and interviews, the facility failed to ensure fifteen day Adverse Incident Reports were completed for 4 out of 4 sampled residents (Residents #1-#4). The findings included: Review of the facility's Adverse Incident Reports,	A 165		

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A 165	Continued From page 3 submitted to the Agency as required for any event over which facility personnel could exercise control rather than as a result of the resident's condition and results in a , revealed the following reports had not been completed. The "one day reports" contained all relevant information showing the facility's investigation into the events, but the "fifteen day report" showing the facility's conclusion of the incidents had not been submitted. a. Resident #1-Event dated b. Resident #2-Event dated c. Resident #3-Event dated d. Resident #4-Event dated During interview with the Administrator on at 11:30 AM, she stated that fifteen day Adverse Incident Report had not been completed for these events because she was advised that this was not necessary, and that she had closed the reports. She confirmed the information in the one day reports for these events was accurate and showed the facility's investigation, but that the fifteen day reports documenting that the events were not "adverse" had not been submitted. It was discussed at this time that the fifteen day reports were required to be completed within fifteen days of the events to show the results of the facility's investigation. The facility provided no additional documentation for review at this time. Class	A 165		