

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 676449	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 02/07/2026
NAME OF PROVIDER OR SUPPLIER Ignite Medical Resort Fort Worth, LLC		STREET ADDRESS, CITY, STATE, ZIP CODE 6301 Oakmont Blvd Fort Worth, TX 76132	
For information on the nursing home's plan to correct this deficiency, please contact the nursing home or the state survey agency.			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (Each deficiency must be preceded by full regulatory or LSC identifying information)		
<p>F 0698</p> <p>Level of Harm - Actual harm</p> <p>Residents Affected - Few</p>	<p>Provide safe, appropriate dialysis care/services for a resident who requires such services.</p> <p>**NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on interview and record review the facility failed to ensure that a resident who required dialysis received such services, consistent with professional standards of practice for 1 (Resident #1) of 1 resident reviewed peritoneal dialysis. The facility failed to ensure nursing staff received training to administer, monitor and intervene for Resident #1 who required PD (During peritoneal dialysis, a cleansing fluid flows through a tube into part of the stomach area, also called the abdomen. The inner lining of the abdomen, known as the peritoneum, acts as a filter and removes wastes from blood. After a set amount of time, the fluid with the filtered waste flows out of the abdomen and is thrown away). Resident #1 was admitted to the hospital with peritonitis (bacteria into the peritoneum). The facility failed to ensure Resident #1 had dialysis supplies available on 01/25/26 and 01/26/26 to perform PD. Resident #1 was not dialyzed on 01/25/26 and 01/26/26 because she did not have supplies (cycler) available. This failure could place residents at risk of not receiving life-sustaining medical treatment which could diminish the quality of life. Findings included: Record review of Resident #1's face sheet dated 02/05/26 reflected she was a [AGE] year-old female admitted to the facility on [DATE] with diagnoses of cerebral infarction (stroke caused by a blockage in a blood vessel supplying the brain), end stage renal disease (permanent stage of kidney failure), type 2 diabetes (a chronic condition where the body develops insulin resistance and fails to produce enough insulin causing high blood sugar levels), and heart failure (the heart cannot pump blood efficiently causing fluid buildup, fatigue, and shortness of breath). Record review of Resident #1's care plan dated 1/20/26 reflected she needed dialysis. The goal was for Resident #1 to have immediate intervention should any s/sx complications from dialysis occur. Resident #1 would have no s/sx complications from dialysis. Intervention was to monitor PD catheter site redness/drainage, report cloudy effluent, inadequate drainage or inflows problems, sudden weight gain or loss, shortness of breath, abdominal pain, fever. Monitor/document report PRN any s/sx of infection to access site, redness, swelling, warmth or drainage, renal insufficiency. PD dialysis 6p-6am, guest to complete independently and supply own supplies. Record review of Resident #1's comprehensive MDS assessment dated [DATE] reflected that she had a BIMS score of 15 (Cognitively Intact). Hemodialysis was indicated while a resident. Record review of hospital H&P dated 2/01/26 reflected, admission date 01/31/26 chief complaint: Abdominal pain, nausea/vomiting. Patient was recently admitted at [local hospital] from 01/11 to 01/20 for LLE weakness + possible peritonitis from PD catheter-cranial imaging showed bilateral MCA + left PCA watershed infarcts, neurology was consulted + started patient on DAPT. ID was consulted and obtained PD fluid studies which were negative for peritonitis. Patient reports during this hospitalization had generalized abdominal pain which was thought to be opioid-induced constipation improved with PRN antiemetics + bowel regimen. Patient was eventually discharged to [facility] but reports during the time she was there she was not eating much food due to continued abdominal pain. Patient's [family] at bedside reports patient's</p> <p>(continued on next page)</p>		

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
FORM CMS-2567 (02/99) Previous Versions Obsolete	Event ID: 676449	Facility ID: 676449 If continuation sheet Page 1 of 4

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