

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER / SUPPLIER / CLIA IDENTIFICATION NUMBER 155572	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED 07/28/2020
NAME OF PROVIDER OF SUPPLIER APERION CARE DEMOTTE		STREET ADDRESS, CITY, STATE, ZIP 10352 N 600 E COUNTY LINE RD DEMOTTE, IN 46310	
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)		
F 0692 Level of harm - Minimal harm or potential for actual harm Residents Affected - Few	<p>Provide enough food/fluids to maintain a resident's health.</p> <p>**NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY**</p> <p>Based on record review and interview, the facility failed to ensure a Physician order [REDACTED]. Hospice services were in place. The 3/5/2020 Minimum Data Set assessment indicated the resident's cognitive skills for decision making were moderately impaired. No behaviors or rejection of care occurred. Assistance of one staff was needed for eating. Current Care plans indicated the resident was at risk for aspiration and on a fluid restriction of 1500 cc of liquids per day. Interventions included, but were not limited to, provide diet as ordered, monitor weight and set up and assist the resident with meals. A physician's orders [REDACTED]. BUN (blood urea nitrogen) level was 94 (normal level range 7-30). Creatinine level was 5.77 (normal level range 0.7- 2.0). The April 2020 Nursing Progress Notes indicated the following: 4/26/20 at 5:30 p.m. NP called with new orders for laboratory results. Daughter was aware and agreed with IV orders. 4/27/20 at 12:22 p.m. Resident's left hand poked x 2 for IV start using a 22 gauge needle. Resident tolerated well. Spoke to the Family and they stated they would like the resident to be as comfortable as possible. There was no documentation of any attempts to insert the IV or start the ordered IV fluids between 4/26/20 at 5:30 p.m. and 4/27/20 at 12:22 p.m. When interviewed on 7/28/20 at 12:30 p.m., LPN 1 indicated she had worked the Day shift (6AM - 6PM) on 4/26/20. She did not recall if she was told about the IV in report that day. When interviewed on 7/28/20 at 12:45 p.m., the ADON (Assistant Director of Nursing) indicated she worked the Day Shift on 4/27/20 and inserted the IV at that time. The ADON was unsure why the IV catheter and fluids were not started prior to that. When interviewed on 7/28/20 at 2:30 p.m. the facility Administrator indicated Resident B was on Hospice services and the IV fluids should have been initiated at the time of the physician's orders [REDACTED].</p>		
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE		TITLE	(X6) DATE

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.