Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	37D0862392	04/14/2022
Name of Provider or Supplier	**Street Address, City, State**	
Ascension St John Medical Center Labor & Delivery	1923 S Utica, Tulsa, OK	

For information on the provider's plan to correct this deficiency, please contact the provider or the state survey agency.

(X4) ID Prefix Tag	Summary Statement of Deficiencies
D0000	The recertification survey was performed on 04/14/2022. The findings were reviewed with the technical consultant and point of care technician at the conclusion of the survey. The laboratory was found in compliance with standard-level deficiency cited.
D5807	TEST REPORT CFR(s): 493.1291(d) Pertinent "reference intervals" or "normal" values, as determined by the laboratory performing the tests, must be available to the authorized person who ordered the tests and, if applicable, the individual responsible for using the test results. This STANDARD is not met as evidenced by: Based on a review of a patient report and interview with the technical consultant, the laboratory failed to provide normal reference intervals for one of one Amnisure patient test report. Findings include: (1) On 04/14/2022 at 10:20 am, the technical consultant stated the following to the surveyor: (a) The laboratory used the Amnisure ROM test, to aid in the detection of the rupture of the fetal membrane. (2) A review of one test report for a patient tested on 02/08/2022 at 12:58 pm. The report did not include a normal reference range; (3) The report was reviewed with the technical consultant, who stated on 04/14/2022 at 01:40 pm the patient report did not include a normal reference range.