

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  10D2109688	<b>(X3) Date Survey Completed</b>  11/07/2023
<b>Name of Provider or Supplier</b>  Florida Institute For Reproductive Medicine	<b>Street Address, City, State</b>  302 Norton Drive Suite 101, Tallahassee, FL	
For information on the provider's plan to correct this deficiency, please contact the provider or the state survey agency.		

<b>(X4) ID Prefix Tag</b>	<b>Summary Statement of Deficiencies</b>
<b>D0000</b>	A recertification survey was conducted on November 7, 2023. Florida Institute for Reproductive Medicine clinical laboratory was not in compliance with 42 CFR 493, requirements for clinical laboratories.
<b>D5221</b>	<p>EVALUATION OF PROFICIENCY TESTING PERFORMANCE CFR(s): 493.1236(d)</p> <p>All proficiency testing evaluation and verification activities must be documented.</p> <p>This STANDARD is not met as evidenced by: Based on record review and staff interview, the facility failed to document corrective action for one unsatisfactory proficiency testing event in 2023. Findings include: The record review of the American Association of Bioanalysts (AAB) proficiency testing event titled "Embryology, Andrology &amp; Fetal 21 2023" showed a score of 50% for the test "Sperm Count, Video" using the Makler method. There was no documentation of corrective action. The interview with testing person A on 11/7/23 a 10:10am confirmed the documentation was missing.</p>