

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 13D0964970	(X3) Date Survey Completed 06/06/2022
Name of Provider or Supplier Oga	Street Address, City, State 3520 E Louise Dr, Meridian, ID	
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
D5805	<p>TEST REPORT CFR(s): 493.1291(c)</p> <p>The test report must indicate the following: (c)(1) For positive patient identification, either the patient's name and identification number, or a unique patient identifier and identification number. (c)(2) The name and address of the laboratory location where the test was performed. (c)(3) The test report date. (c)(4) The test performed. (c)(5) Specimen source, when appropriate. (c)(6) The test result and, if applicable, the units of measurement or interpretation, or both. (c)(7) Any information regarding the condition and disposition of specimens that do not meet the laboratory's criteria for acceptability.</p> <p>This STANDARD is not met as evidenced by: Based on a review of patient laboratory reports and an interview with the laboratory lead on 6/6/2022, the laboratory failed to indicate the address of the performing laboratory for wet prep testing. The findings include: 1. A review of patient laboratory test reports for vaginal wet prep testing identified that the laboratory failed to indicate the address of the performing laboratory. 2. An interview with the laboratory lead on 6/6/2022 at 2:16 pm confirmed that the address of the performing laboratory for vaginal wet prep testing is not indicated on the test report. 3. The laboratory reports performing 1,016 vaginal wet prep tests annually.</p>
D5807	<p>TEST REPORT CFR(s): 493.1291(d)</p> <p>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.</p>

This STANDARD is not met as evidenced by:

Based on a review of patient laboratory test reports and an interview with the laboratory lead on 6/6/2022, the laboratory failed to include the normal values for the reported vaginal wet prep analytes on patient laboratory test reports. The findings include: 1. A review of patient laboratory test reports identified that the laboratory failed to include the normal values for the analytes of a vaginal wet prep; yeast, bacteria, trichomonas vaginalis, epithelial cells, clue cells, white blood cells and the whiff test. 2. An interview with the laboratory lead on 6/6/22 at 2:16 pm confirmed the above findings. 3. The laboratory reports performing 1,016 vaginal wet prep tests annually.