

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  28D0455760	<b>(X3) Date Survey Completed</b>  11/10/2025
<b>Name of Provider or Supplier</b>  Grand Island Clinic Inc	<b>Street Address, City, State</b>  2444 W Faidley Ave, Grand Island, NE	
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 proficiency testing desk review was completed on November 10, 2025. At the time of the review, the laboratory was not in compliance with the Clinical Laboratory Improvement Amendments of 1988, 42 CFR 493.1 through 42 CFR 493.1780. The following condition deficiencies were cited: D2016 - 42 C.F.R. 493.803 Condition: Successful participation [proficiency testing] D6076 - 42 C.F.R. 493.1441 Condition: Laboratories performing high complexity testing; laboratory director
<b>D2016</b>	<p><b>SUCCESSFUL PARTICIPATION</b> CFR(s): 493.803(a)(b)(c)</p> <p>(a) Each laboratory performing nonwaived testing must successfully participate in a proficiency testing program approved by CMS, if applicable, as described in subpart I of this part for each specialty, subspecialty, and analyte or test in which the laboratory is certified under CLIA. (b) Except as specified in paragraph (c) of this section, if a laboratory fails to participate successfully in proficiency testing for a given specialty, subspecialty, analyte or test, as defined in this section, or fails to take remedial action when an individual fails gynecologic cytology, CMS imposes sanctions, as specified in subpart R of this part. (c) If a laboratory fails to perform successfully in a CMS-approved proficiency testing program, for the initial unsuccessful performance, CMS may direct the laboratory to undertake training of its personnel or to obtain technical assistance, or both, rather than imposing alternative or principle sanctions except when one or more of the following conditions exists: (1) There is immediate jeopardy to patient health and safety. (2) The laboratory fails to provide CMS or a CMS agent with satisfactory evidence that it has taken steps to correct the problem identified by the unsuccessful proficiency testing performance. (3) The laboratory has a poor compliance history.</p> <p>This CONDITION is not met as evidenced by: Based on a proficiency testing desk review from the Certification and Survey Provider Enhanced Reporting (CASPER) 0155 report and College of American Pathologists</p>

	<p>(CAP) 2025 records, the laboratory did not successfully participate in a proficiency testing program approved by HHS, for each specialty, subspecialty, and analyte or test in which the laboratory is certified under CLIA. The laboratory failed to successfully participate in the specialty of Bacteriology. Refer to D2028.</p>
<b>D2028</b>	<p><b>BACTERIOLOGY</b> CFR(s): 493.823(e)</p> <p>(e) Failure to achieve an overall testing event score of satisfactory performance for two consecutive testing events or two out of three consecutive testing events is unsuccessful performance.</p> <p>This STANDARD is not met as evidenced by: Based on a proficiency testing desk review of CASPER 0155 report and College of American Pathologists (CAP) 2025 proficiency testing records and phone interview with the laboratory manager, the laboratory failed to achieve satisfactory performance (80% or greater) for two consecutive testing events or two out of three consecutive testing events in the subspecialty of Bacteriology. Findings are: 1. Review of the CASPER 0155 report revealed the following results: Bacteriology 2025 - 1st Event the laboratory received an unsatisfactory score of 78% for Bacteriology. Bacteriology 2025 - 2nd Event the laboratory received an unsatisfactory score of 66% for Bacteriology. 2. Review of the CAP 2025 proficiency testing records confirmed the laboratory received the above results. 3. Phone interview with the laboratory manager on November 10, 2025 at 12:09 PM, confirmed the above results.</p>
<b>D6076</b>	<p><b>LABORATORY DIRECTOR</b> CFR(s): 493.1441</p> <p>The laboratory must have a director who meets the qualification requirements of 493.1443 of this subpart and provides overall management and direction in accordance with 493.1445 of this subpart.</p> <p>This CONDITION is not met as evidenced by: Based on a proficiency testing desk review of CASPER 0155 report and College of American Pathologists 2025 (1st and 2nd events) records, the laboratory director failed to manage successful proficiency testing participation. Refer to D6089.</p>
<b>D6089</b>	<p><b>LABORATORY DIRECTOR RESPONSIBILITIES</b> CFR(s): 493.1445(e)(4)(i)</p> <p>(e)(4)(i) The proficiency testing samples are tested as required under subpart H of this part;</p> <p>This STANDARD is not met as evidenced by: Based on a proficiency testing desk review of CASPER 0155 report and College of American Pathologists 2025 (1st and 2nd events) records, the laboratory director failed to ensure successful participation in an HHS approved proficiency testing program. Refer to D2028.</p>