

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  27D0411039	<b>(X3) Date Survey Completed</b>  12/01/2025
<b>Name of Provider or Supplier</b>  Bitterroot Health Daly	<b>Street Address, City, State</b>  1200 Westwood Dr, Hamilton, MT	
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 December 1, 2025. At the time of the desk review, it was determined that the laboratory was not in compliance with all conditions required by the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 Code of Federal Regulations, Part 493 (42 C.F.R. 493). The following condition level deficiency was cited: 493.803 Condition: Successful participation.
<b>D2016</b>	<p>SUCCESSFUL PARTICIPATION 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 American Association of Bioanalysts (AAB) 2025 records, the laboratory failed to successfully participate in a proficiency</p>

	<p>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 subspecialties of Endocrinology and Routine Chemistry for analytes Folate and Hemoglobin A1c. Refer to D2096, D2107</p>
<p><b>D2096</b></p>	<p><b>ROUTINE CHEMISTRY</b> CFR(s): 493.841(f)</p> <p>(f) Failure to achieve satisfactory performance for the same analyte or test in 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 2025 American Association of Bioanalysts (AAB) proficiency testing records, the laboratory failed to achieve satisfactory performance (80% or better) for two consecutive proficiency testing events in the subspecialty of Routine Chemistry for the analyte Hemoglobin A1c. Findings: 1. Review of the CASPER 0155 report revealed the following results: Routine Chemistry 2025-2nd Event, the laboratory received an unsatisfactory score of 60% for Hemoglobin A1c Routine Chemistry 2025-3rd Event, the laboratory received an unsatisfactory score of 0% for Hemoglobin A1c 2. A review of the 2025 AAB Proficiency Testing records confirmed the laboratory received the above results.</p>
<p><b>D2107</b></p>	<p><b>ENDOCRINOLOGY</b> CFR(s): 493.843(f)</p> <p>(f) Failure to achieve satisfactory performance for the same analyte or test in 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 2025 American Association of Bioanalysts (AAB) proficiency testing records, the laboratory failed to achieve satisfactory performance (80% or better) for two consecutive proficiency testing events in the subspecialty of Endocrinology for the analyte Folate. Findings: 1. Review of the CASPER 0155 report revealed the following results: Endocrinology 2025-2nd Event, the laboratory received an unsatisfactory score of 60% for Folate Endocrinology 2025-3rd Event, the laboratory received an unsatisfactory score of 0% for Folate 2. A review of the 2025 AAB Proficiency Testing records confirmed the laboratory received the above results.</p>