

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  39D2021686	<b>(X3) Date Survey Completed</b>  02/02/2018
<b>Name of Provider or Supplier</b>  Perll Diagnostics Inc	<b>Street Address, City, State</b>  5010 Ritter Road, Mechanicsburg, PA	
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>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 review of the CASPER Report 155 and performance evaluations from the proficiency testing organization American Proficiency Institute (API), the laboratory failed to successfully participate in a proficiency testing program approved by CMS for the analyte: White Blood Cell Differential (WBC) which is of the specialty Hematology. The laboratory had unsatisfactory scores for the 1st event of 2017 and the 3rd event of 2017. See D2130.</p>
<b>D2130</b>	<p><b>HEMATOLOGY</b> CFR(s): 493.851(f)</p>

Failure to achieve satisfactory performance for the same analyte in two consecutive events or two out of three consecutive testing events is unsuccessful performance.

This STANDARD is not met as evidenced by:

Based on a review of the CASPER 155 report and performance evaluations from the proficiency testing organization American Proficiency Institute (API), the laboratory failed to successfully participate in a proficiency testing program approved by CMS for the analyte: White Blood Cell Differential which is in the specialty of Hematology in which the laboratory is certified under CLIA. The laboratory had unsatisfactory scores for the 1st event of 2017, and 3rd event 2017 for the analyte listed above.

Findings include: 1. API 2017 Event 1 for White Blood Cell Differential the score was 0% and was unsatisfactory. 2. API 2017 Event 3 for White Blood Cell Differential the score was 48% and was unsatisfactory.