

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 23D1010918	(X3) Date Survey Completed 10/01/2018
Name of Provider or Supplier City Medical	Street Address, City, State 30581 Stephenson Hwy, Madison Heights, MI	
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
D2016	<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 review of the CMS database and review of the American Proficiency Institute (API) final graded proficiency testing reports, it was determined that the laboratory failed to successfully participate in a CMS approved proficiency testing program for the hematology analyte: cell identification (ID) or white blood cell (WBC) diff. Findings include: Review of the CMS database and the API proficiency testing reports showed unsatisfactory performance for two of three proficiency testing events for the hematology analyte: cell ID or WBC diff. Refer to D2130</p>
D2130	HEMATOLOGY

CFR(s): 493.851(f)

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 review of the CMS database and review of the American Proficiency Institute (API) final graded proficiency testing reports, the laboratory failed to achieve satisfactory performance in two out of three consecutive proficiency testing events for the following hematology analyte: Cell Identification (ID) or White Blood Cell (WBC) Differential (Diff).(HCT). Findings include: 1. Unsatisfactory performance in two out of three proficiency testing events constitutes subsequent unsuccessful proficiency testing performance. WBC diff PT Event Score 3rd event 2017 53% 2nd event 2018 47% 2. Unsatisfactory performance in four out of five proficiency testing events constitutes subsequent unsuccessful proficiency testing performance. WBC diff PT Event Score 1st event 2017 7% 2nd event 2017 67% 3rd event 2017 53% 2nd event 2018 47%