

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  14D2264781	<b>(X3) Date Survey Completed</b>  10/06/2025
<b>Name of Provider or Supplier</b>  Hmcs Inc - Home Med Diagnostic & Lab Svcs	<b>Street Address, City, State</b>  3500 W Peterson Ave - Unit 404, Chicago, IL	
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>	Based on a desk review of proficiency testing records from the Certification and Survey Provider Enhanced Reporting (CASPER) database and verified with the proficiency testing provider the laboratory was found to be out of compliance with the following CONDITION level deficiencies: D2016 - 42 C.F.R. 493.803 Condition: Successful participation [proficiency testing] D6000 - 42 C.F.R. 493.1403 Condition: Laboratories performing moderate 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 of the Certification and Survey Provider Enhanced Reporting (CASPER) 0155D Report and American Proficiency Institute</p>

	<p>(API) Proficiency Testing (PT) records confirmed the initial unsuccessful PT performance for the analyte Erythrocyte Count for two consecutive PT events in 2025 (events 1 and 2). Refer to D2130.</p>
<p><b>D2130</b></p>	<p><b>HEMATOLOGY</b> CFR(s): 493.851(f)</p> <p>(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.</p> <p>This STANDARD is not met as evidenced by: Based on a proficiency testing desk review of the Certification and Survey Provider Enhanced Reporting (CASPER) 0155D Report and American Proficiency Institute (API) Proficiency Testing (PT) records, the laboratory failed to achieve satisfactory performance for the hematology analyte Erythrocyte Count for two consecutive PT events in 2025 (events 1 and 2) resulting in the initial unsuccessful PT performance. Findings include: 1. Review of the CASPER Report 0155D, generated on 10-01-2025, the laboratory received the following unsatisfactory scores for the hematology analyte Erythrocyte Count. Erythrocyte Count EVENT 1, 2025 - 40% Unsatisfactory EVENT 2, 2025 - 40% Unsatisfactory 2. Review of API PT evaluation reports (2025 Hematology/Coagulation - 1st &amp; 2nd Event) confirmed the above unsatisfactory scores that resulted in the initial unsuccessful PT performance for the hematology analyte Erythrocyte Count.</p>
<p><b>D6000</b></p>	<p><b>MODERATE COMPLEXITY LABORATORY DIRECTOR</b> CFR(s): 493.1403</p> <p>The laboratory must have a director who meets the qualification requirements of 493.1405 of this subpart and provides overall management and direction in accordance with 493.1407 of this subpart.</p> <p>This CONDITION is not met as evidenced by: Based on a proficiency desk review of the Certification and Survey Provider Enhanced Reporting (CASPER) 0155D Report and American Proficiency Institute (API) Proficiency Testing (PT) records, the laboratory director failed to ensure successful participation in a Health and Human Services (HHS) approved PT program for the hematology analyte Erythrocyte Count resulting in the laboratory's initial unsuccessful PT performance. Refer to D6016.</p>
<p><b>D6016</b></p>	<p><b>LABORATORY DIRECTOR RESPONSIBILITIES</b> CFR(s): 493.1407(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 the Certification and Survey Provider Enhanced Reporting (CASPER) 0155D Report and American Proficiency Institute (API) Proficiency Testing (PT) records, the laboratory director failed to ensure successful participation in a Health and Human Services (HHS) approved PT program</p>

for the hematology analyte Erythrocyte Count resulting in the laboratory's initial unsuccessful PT performance. Refer to D2130.