

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  17D1052648	<b>(X3) Date Survey Completed</b>  04/02/2026
<b>Name of Provider or Supplier</b>  Kansas Medical Center, Llc	<b>Street Address, City, State</b>  1124 W 21st Street, Andover, KS	
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 Centers for Medicare &amp; Medicaid Services (CMS) Proficiency Testing (PT) Certification and Survey Provider Enhanced Reporting system and the American Proficiency Institute (API) PT summary reports, the laboratory failed to successfully participate in the CMS approved PT 2025/2026 program under the specialty of Chemistry, for two of three testing events. Refer D2087.</p>
<b>D2087</b>	<p><b>ROUTINE CHEMISTRY</b> CFR(s): 493.841(a)</p>

(a) Failure to attain a score of at least 80 percent of acceptable responses for each analyte in each testing event is unsatisfactory analyte performance for the testing event.

This STANDARD is not met as evidenced by:

Based on a desk review of proficiency testing (PT) on the CMS CASPER Reports and the provider American Proficiency Institute (API) for Routine Chemistry, the laboratory failed to achieve an acceptable score of 80% or higher for two of three testing events for the regulated routine chemistry analyte: 0340 - B Natriuretic Peptide (BNP). Findings: 1. Review of the API PT scores for Event 2 2025 for the analyte BNP revealed a 60% performance score. 2. Review of the API PT scores for Event 1 2026 for the analyte BNP revealed a 20% performance score. 3. A phone interview with the assistant laboratory supervisor on April 2, 2026 at 10:45 a.m. and review of the CMS CASPER Reports confirmed, the laboratory failed to achieve an acceptable score of 80% or higher for two of three testing events for the regulated routine chemistry analytes: 0340 - B Natriuretic Peptide (BNP).