

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 37D2247441	(X3) Date Survey Completed 09/27/2024
Name of Provider or Supplier Ou Health Er & Urgent Care Czech Hall	Street Address, City, State 709 N Czech Hall Rd, Yukon, OK	
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
D0000	The following deficiencies are a result of a desk review of proficiency testing scores obtained from the national database and from the proficiency testing provider. The laboratory was found out of compliance with the following CLIA Conditions: 493.803; D2016: Successful Participation 493.1407; D6000: Laboratory Director, Moderate Complexity
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 a desk review of proficiency testing scores obtained from the Casper 0155D report and proficiency testing Comparative Evaluation obtained from the laboratory for the first and second 2024 events, the laboratory failed to successfully participate in</p>

	<p>a proficiency testing program for the specialty of Hematology. Findings include: (1) The laboratory failed to achieve satisfactory performance for two consecutive testing events for Cell Identification/White Blood Cell Differential for %Neutrophils, Absolute Neutrophils, %Lymphocytes; and WBC (White Blood Cell) Count. Refer to D2130.</p>
<p>D2130</p>	<p>HEMATOLOGY CFR(s): 493.851(f)</p> <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.</p> <p>This STANDARD is not met as evidenced by: Based on a desk review of proficiency testing scores obtained from the Casper 0155D report and the proficiency testing graded evaluations obtained from the laboratory, the laboratory failed to achieve satisfactory performance for Cell Identification/White Blood Cell Differential for %Neutrophils, Absolute Neutrophils, %Lymphocytes; and WBC (White Blood Cell) Count for two consecutive testing events. Findings include: (1) WBC Count - The laboratory received a score of 0% on the first 2024 event and a score of 60% on the second 2024 event; (2) %Neutrophils - The laboratory received a score of 0% on the first 2024 event and a score of 60% on the second 2024 event; (3) Absolute Neutrophils - The laboratory received a score of 0% on the first 2024 event and a score of 60% on the second 2024 event; (4) %Lymphocytes - The laboratory received a score of 0% on the first 2024 event and a score of 60% on the second 2024 event.</p>
<p>D6000</p>	<p>MODERATE COMPLEXITY LABORATORY DIRECTOR 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 desk review of proficiency testing scores obtained from the Casper 0155D report and the proficiency testing graded evaluations obtained from the laboratory, the laboratory failed to achieve satisfactory performance for Cell Identification/White Blood Cell Differential for %Neutrophils, Absolute Neutrophils, %Lymphocytes; and WBC (White Blood Cell) Count for two consecutive testing events. The laboratory failed to achieve a passing score of 80% for the first and second 2024 events. Refer to D6016.</p>
<p>D6016</p>	<p>LABORATORY DIRECTOR RESPONSIBILITIES CFR(s): 493.1407(e)(4)(i)</p> <p>The laboratory director is responsible for the overall operation and administration of the laboratory, including the employment of personnel who are competent to perform test procedures, and record and report test results promptly, accurate, and proficiently and for assuring compliance with the applicable regulations. (e) The laboratory director must-- (e)(4)(i) Ensure that the proficiency testing samples are tested as required under Subpart H of this part;</p>

This STANDARD is not met as evidenced by:

Based on a desk review of proficiency testing scores obtained from the Casper 0155D report and the proficiency testing graded evaluations obtained from the laboratory, the laboratory failed to achieve satisfactory performance for Cell Identification/White Blood Cell Differential for %Neutrophils, Absolute Neutrophils, %Lymphocytes; and WBC (White Blood Cell) Count for two consecutive testing events. The laboratory failed to achieve a passing score of 80% for the first and second 2024 events. Refer to D2130.