

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 25D2160279	(X3) Date Survey Completed 08/14/2019
Name of Provider or Supplier Mea Medical Clinic Flora	Street Address, City, State 740 Hwy 49 N Ste V, Flora, MS	
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	A revisit survey (completed by mail) was conducted on 11/25/19 for all previous deficiencies cited on 8/14/19. All deficiencies have been corrected, and no new noncompliance was found. The facility is in compliance with all regulations surveyed.
D6015	<p>LABORATORY DIRECTOR RESPONSIBILITIES CFR(s): 493.1407(e)(4)</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) Ensure that the laboratory is enrolled in an HHS approved proficiency testing program for the testing performed.</p> <p>This STANDARD is not met as evidenced by: Based on surveyor review of the laboratory proficiency testing records, Centers of Medicare and Medicaid Services (CMS) database proficiency testing report and confirmation with technical consultant at 1:00 pm on 8/14/19, the laboratory director failed to ensure the laboratory was enrolled and participated in an HHS approved proficiency testing (PT) program for CBC (complete blood count) performed on the Boule Medonic M Series hematology analyzer, for the second testing event of 2019. The laboratory director must ensure the laboratory is enrolled and participates in all 3 events in an approved proficiency program for the testing performed by the laboratory. Findings include: 1. Observation of the CMS database proficiency testing report revealed no scores for 2019. 2. Observation of laboratory records since installation of Medonic M Series on 2/7/19 through the day of survey, 8/14/19, revealed no evidence of proficiency testing participation prior to survey. 3. Interview with technical consultant at 1:00 pm on day of survey and observation of recent</p>

proficiency enrollment forms revealed the laboratory did not enroll in proficiency in time to participate in the 2nd event of 2019 when installation of the analyzer was 2/7/19 and CBC patient testing began 3/5/19.