

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  11D1020334	<b>(X3) Date Survey Completed</b>  02/28/2020
<b>Name of Provider or Supplier</b>  Kayal Dermatology And Skin Cancer	<b>Street Address, City, State</b>  141 Lacy Street, Suite 200, Marietta, GA	
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>	A Clinical Laboratory Improvement Amendments (CLIA) recertification survey was completed on February 28, 2020. The laboratory was not in compliance with applicable CLIA requirements found at 42 CFR 493.1 through 42 CFR 493.1780. The following deficiencies were cited:
<b>D3011</b>	<p><b>FACILITIES</b> CFR(s): 493.1101(d)</p> <p>Safety procedures must be established, accessible, and observed to ensure protection from physical, chemical, biochemical, and electrical hazards, and biohazardous materials.</p> <p>This STANDARD is not met as evidenced by: Based on observation and staff interview, the laboratory failed to observe safety procedures to ensure protection from physical, chemical, and electrical hazards as required. Findings include: 1. Observation during the laboratory tour and subsequent office tour on 2/28/2020 at approximately 11:30 a.m. revealed all three fire extinguishers in the facility were last inspected on 9/06/2017. 2. An interview with the lead histotech in the laboratory on 2/28/2020 at approximately 11:45 a.m. confirmed all three fire extinguishers in the facility were last inspected on 9/06/2017.</p>
<b>D5217</b>	<p><b>EVALUATION OF PROFICIENCY TESTING PERFORMANCE</b> CFR(s): 493.1236(c)(1)</p> <p>At least twice annually, the laboratory must verify the accuracy of any test or procedure it performs that is not included in subpart I of this part.</p> <p>This STANDARD is not met as evidenced by: Based on testing personnel (TP) document review and staff interview, the laboratory</p>

failed to verify twice annually the accuracy of any unregulated test or procedure it performs as required. Findings include: 1. Based on TP document review, the laboratory failed to have available at the time of survey 2018 twice annual peer review documents for Staff #5 (CMS 209). 2. An interview with the lead histotech in the laboratory on 2/28/2020 at approximately 11:45 a.m. confirmed the lack of 2018 twice annual peer reviews for Staff #5 (CMS 209).