

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  31D1036684	<b>(X3) Date Survey Completed</b>  01/24/2018
<b>Name of Provider or Supplier</b>  Atlantic Dermatology & Laser Center	<b>Street Address, City, State</b>  1401 New Road, Linwood, NJ	
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>D5601</b>	<p><b>HISTOPATHOLOGY</b> CFR(s): 493.1273(a)(f)</p> <p>(a) As specified in 493.1256(e)(3), fluorescent and immunohistochemical stains must be checked for positive and negative reactivity each time of use. For all other differential or special stains, a control slide of known reactivity must be stained with each patient slide or group of patient slides. Reactions of the control slide with each special stain must be documented. (f) The laboratory must document all control procedures performed, as specified in this section.</p> <p>This STANDARD is not met as evidenced by: Based on observation of Quality Control (QC) records, accession log, and interview with the Office Manager (OM), the laboratory failed to document the reactions of the QC slides for Periodic acid-Schiff (PAS) stains and Grocott methenamine-silver (GM) used for Histopathology slides on 10/30/17. The findings include: 1. There was no QC documented for PAS and GM stains on 10/30/17. 2. Approximately 19 Patients were affected. 3. The OM confirmed on 1/24/18 at 10:00 am that the QC reactions were not documented.</p>