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| <b>Statement of Deficiencies</b>   | <b>(X1) Provider/Supplier/CLIA Identification Number</b><br><br>21D2043930                    | <b>(X3) Date Survey Completed</b><br><br>09/26/2024 |
| <b>Name of Provider or Supplier</b><br><br>Anne Arundel Dermatology  | <b>Street Address, City, State</b><br><br>7671 Quarterfield Road Ste 200 A-B, Glen Burnie, MD |   |
| 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>   |
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| <b>D5413</b>              | <p>TEST SYSTEMS, EQUIPMENT, INSTRUMENTS, REAGENT<br/>CFR(s): 493.1252(b)</p> <p>The laboratory must define criteria for those conditions that are essential for proper storage of reagents and specimens, accurate and reliable test system operation, and test result reporting. The criteria must be consistent with the manufacturer's instructions, if provided. These conditions must be monitored and documented and, if applicable, include the following: (1) Water quality. (2) Temperature. (3) Humidity. (4) Protection of equipment and instruments from fluctuations and interruptions in electrical current that adversely affect patient test results and test reports.</p> <p>This STANDARD is not met as evidenced by:<br/>Based on record review and interview, the laboratory did not record the humidity of the room where processing and staining of tissue for MOHS surgery was performed. Findings: 1. The laboratory reviewed the humidity readings for the room where processing and staining was performed, but did not document those reading on the log each day when MOHS surgery was performed. The log had a place to document these readings, and of four of four days reviewed, when MOHS surgery was performed in September 2024, the humidity was not documented. 2. This was confirmed with the histotechnologist at 10:30 a.m. on September 26, 2024.</p> |