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| <b>Statement of Deficiencies</b>   | <b>(X1) Provider/Supplier/CLIA Identification Number</b><br><br>21D0688709                       | <b>(X3) Date Survey Completed</b><br><br>03/27/2026 |
| <b>Name of Provider or Supplier</b><br><br>Anne Arundel Dermatology, Pa  | <b>Street Address, City, State</b><br><br>995 N Prince Frederick Blvd #204, Prince Frederick, 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>(b) 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: (b)(1) Water quality. (b)(2) Temperature. (b)(3) Humidity. (b)(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 define the acceptable ranges for temperature and humidity in the room where the cryostat was located. Findings: 1. The daily maintenance log did not state the acceptable temperature and humidity range for the room where the cryostat was located. 2. This was confirmed with the histotechnician on 3/27/26 at 12:00 pm.</p> |