

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 10D0958608	(X3) Date Survey Completed 02/24/2025
Name of Provider or Supplier Dermatology Group Of Florida, Pa	Street Address, City, State 5415 Park Central Ct, Naples, FL	
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	An announced CLIA recertification survey was conducted at DERMATOLOGY GROUP OF FLORIDA, PA from 02/21/2025 to 02/24/2025. The laboratory was not in compliance with 42 CFR Part 493, Requirements for Laboratories. The following is a description of the standard level deficiency:
D5805	<p>TEST REPORT CFR(s): 493.1291(c)</p> <p>(c) The test report must indicate the following: (c)(1) For positive patient identification, either the patient's name and identification number, or a unique patient identifier and identification number. (c)(2) The name and address of the laboratory location where the test was performed. (c)(3) The test report date. (c)(4) The test performed. (c)(5) Specimen source, when appropriate. (c)(6) The test result and, if applicable, the units of measurement or interpretation, or both. (c)(7) Any information regarding the condition and disposition of specimens that do not meet the laboratory's criteria for acceptability.</p> <p>This STANDARD is not met as evidenced by: Based on random patient reports review and staff interview, the laboratory failed to ensure that the patient report listed the name of the laboratory that performed the testing for four out of four patient's reports reviewed. Findings include: Review of four final patient reports: P#1(dated 11/03/2023), P#2 (dated 03/25/2024), P#3 (dated 10/15/2024) and P#4 (dated 02/10/2025), revealed that the reports failed to list the name of the laboratory that performed the histopathology testing. During an interview on 02/21/2025 at 11:30 AM, with the laboratory histotechnician, he confirmed that the final report reviewed did not include the name of the laboratory that performed the histopathology testing for the test reports reviewed.</p>