Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	16D0864255	10/28/2025
Name of Provider or Supplier	**Street Address, City, State**	
Iowa Dermatology Clinic	6000 University Avenue, Suite #350, West Des Moines, IA	

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
D5801	TEST REPORT CFR(s): 493.1291(a) (a) The laboratory must have an adequate manual or electronic system(s) in place to ensure test results and other patient-specific data are accurately and reliably sent from the point of data entry (whether interfaced or entered manually) to final report destination, in a timely manner. This includes the following: (a)(1) Results reported from calculated data. (a)(2) Results and patient-specific data electronically reported to network or interfaced systems. (a)(3) Manually transcribed or electronically transmitted results and patient-specific information reported directly or upon receipt from outside referral laboratories, satellite or point-of-care testing locations. This STANDARD is not met as evidenced by: Based on review of patient test logs, patient electronic health records (EHR), laboratory policies and procedures, and confirmed by interview with Technical Supervisor #1 (TS #1) at 10:50 am on 10/28/2025, the laboratory failed to have a system in place to ensure the accurate and reliable transcription of manual test results into the laboratory's EHR for two out of three patients (patient identifiers A & B) reviewed having potassium hydroxide (KOH) examination testing performed in July and August 2025. The findings include: 1. Review of the laboratory's KOH log indicated that patient A had a KOH examination performed on 07/25/2025 with a positive result. 2. Patient A's EHR chart did not include a record of the KOH examination performed on 07/25/2025. 3. Review of the laboratory's KOH log indicated that patient B had a KOH examination performed on 08/01/2025 with a negative result. 4. Patient B's EHR chart did not include a record of the KOH examination performed on 08/01/2025. 5. The laboratory's Potassium Hydroxide Examination policy/procedure stated, "Results are reported by the physician and

	recorded on the patient's chart immediately by the physician." 6. At the time of the survey, TS #1 confirmed that the laboratory did not have a system in place to ensure the accurate and reliable transcription of KOH testing into the EHR.