

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  16D2258627	<b>(X3) Date Survey Completed</b>  02/22/2023
<b>Name of Provider or Supplier</b>  Forefront Dermatology	<b>Street Address, City, State</b>  2769 Heartland Dr, Suite 303, Coralville, IA	
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>D5801</b>	<p><b>TEST REPORT</b> CFR(s): 493.1291(a)</p> <p>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.</p> <p>This STANDARD is not met as evidenced by: Based on review of the Mohs surgery patient log, patient specimen slides, Mohs maps, and test reports, and confirmed by laboratory personnel identifier #1 (refer to the Laboratory Personnel Report) at approximately 10:00 am on 02/22/2023, the laboratory failed to have a system in place to ensure the accuracy and reliability of Mohs surgery findings/results manually transcribed into patient electronic health records (EHR) for one out of three patients (Patient identifier A) reviewed from January 2023. The findings include: 1. Patient identifier A had Mohs surgery performed on 01/05/2023. 2. Review of the Mohs surgery patient log, patient specimen slides, and Mohs map for patient identifier A all showed the surgeon performed Mohs surgery in two stages. 3. The test report for patient identifier A listed one stage for Mohs surgery performed on 01/05/2023. Furthermore, documentation of stage one of the surgery did not include the findings of the slide review. 4. At the time of the survey, personnel identifier #1 confirmed that the test report for Mohs surgery performed on 01/05/2023 for patient identifier A should have included two stages total and the findings from the first stage of the surgery.</p>