

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 46D2042238	(X3) Date Survey Completed 04/17/2023
Name of Provider or Supplier Utah Valley Dermatology	Street Address, City, State 680 E Main St Suite 201, Lehi, UT	
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
D2010	<p>TESTING OF PROFICIENCY TESTING SAMPLES CFR(s): 493.801(b)(2)</p> <p>The laboratory must test samples the same number of times that it routinely tests patient samples.</p> <p>This STANDARD is not met as evidenced by: A review of American Proficiency Institute (API) Proficiency Testing (PT) attestation statements and an interview with the clinical manager revealed that the laboratory failed to test PT samples in the Microbiology PT events for Potassium Hydroxide (KOH) testing the same number of times that it routinely tests patient samples. Approximately 125 KOH tests are performed annually. Findings include: 1. A review of the attestation statements for the API 2021 Microbiology 1st Event, 2022 Microbiology 2nd Event, and 2022 Microbiology 3rd event, at approximately 4:00 PM on 4/17/23, revealed that multiple providers attested to performing the testing on the KOH sample slides. 2. The API Microbiology events contain 2 slides per event. In the 2021 Microbiology 1st Event the attestation statement lists 5 Testing Personnel (TP) that each read both slides. The attestation statement for the 2022 Microbiology 2nd Event lists 4 TP that each read both slides. The 2022 Microbiology 3rd event attestation statement lists 6 TP that each read both slides. 3. In an interview with the clinical manager at approximately 4:30 PM on 4/17/23, it was stated that all the providers that participate in the PT event will each read the slides and come to a consensus before submitting the results for the PT testing. 4. In an interview with the clinical manager at approximately 4:30 PM on 4/17/23, it was confirmed that the laboratory PT testing was not performed the same number of times that it routinely tests patient samples.</p>