

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  03D2089517	<b>(X3) Date Survey Completed</b>  09/24/2024
<b>Name of Provider or Supplier</b>  Dermatology & Skin Surgery Specialists Of Arizona	<b>Street Address, City, State</b>  8415 N Pima Rd Ste 212, Scottsdale, AZ	
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>D5473</b>	<p>CONTROL PROCEDURES CFR(s): 493.1256(e)(2)(g)</p> <p>(e) For reagent, media, and supply checks, the laboratory must do the following: (e) (2) Each day of use (unless otherwise specified in this subpart), test staining materials for intended reactivity to ensure predictable staining characteristics. Control materials for both positive and negative reactivity must be included, as appropriate. (g) The laboratory must document all control procedures performed.</p> <p>This STANDARD is not met as evidenced by: Based on lack of Quality Control (QC) documentation for two out of five testing dates reviewed during the survey and interview with the facility personnel, the laboratory failed to document the acceptability of Hematoxylin &amp; Eosin (H&amp;E) staining materials each day of use, for intended reactivity to ensure predictable staining characteristics. Findings include: 1. No documentation of the H&amp;E stain acceptability was presented for review for the Mohs procedures that occurred on 3/31/23 and 9/26/23. 2. Nine Mohs cases were performed on 3/31/23 and six Mohs cases were performed on 9/26/23. 3. The facility personnel interviewed on 9/24/24 at 8:50 AM confirmed the laboratory failed to document the H&amp;E stain acceptability on the testing dates indicated above, for intended reactivity to ensure predictable staining characteristics.</p>