

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 03D2188616	(X3) Date Survey Completed 08/21/2026
Name of Provider or Supplier Regency Specialties	Street Address, City, State 14725 W Mountain View Blvd, Suite 100, Surprise, AZ	
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
D5413	TEST SYSTEMS, EQUIPMENT, INSTRUMENTS, REAGENT CFR(s): 493.1252(b) (b) The laboratory must define criteria for those conditions that are essential for proper storage of reagents and specimens, accurate and reliable test system operation, and test result reporting. The criteria must be consistent with the manufacturer's instructions, if provided. These conditions must be monitored and documented and, if applicable, include the following: (b)(1) Water quality. (b)(2) Temperature. (b)(3) Humidity. (b)(4) Protection of equipment and instruments from fluctuations and interruptions in electrical current that adversely affect patient test results and test reports. This STANDARD is not met as evidenced by: Based on review of temperature records from 2025, review of laboratory policy and interview with the facility personnel on 8/21/26 at 1:08 PM, the laboratory failed to monitor and document the room temperature where patient testing occurred on 11 out of 11 testing dates between October 28, 2025 and November 26, 2025. Findings include: 1. The laboratory performs the microscopic interpretation of Mohs specimens in the subspecialty of histopathology. The laboratory's reported annual test volume is 3,200. 2. The laboratory's established room temperature range is 70-75 degrees Fahrenheit. It is the policy of the laboratory to monitor and document the room temperature of the area where Mohs specimens are processed each day of patient testing. 3. The laboratory failed to monitor and document the room temperature of the area where Mohs specimens are processed on 11 out of 11 testing dates between October 28, 2025 and November 26, 2025. 4. Approximately 220 patients were tested between October 28, 2025 and November 26, 2025. 5. The facility personnel interviewed on 8/21/26 at 1:08 PM confirmed that the laboratory failed to monitor and document the room temperature of the laboratory on the testing dates referenced above.