Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	37D2177137	08/27/2025
Name of Provider or Supplier	**Street Address, City, State**	
Olympia Neurological Institute	6130 E 81st St, Tulsa, OK	

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
D0000	The recertification survey was performed on 08/27/2025. The laboratory was found in compliance with standard-level deficiencies cited. The findings were reviewed with testing person #3 and the regional manager during an exit conference performed at the conclusion of the survey.
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 a review of records, manufacturer's instructions, and interview with testing person #3, the laboratory failed to ensure the laboratory humidity was maintained as required by the manufacturer of the Horiba Yumizen C1200 analyzer for 29 of 54 days of patient testing. Findings include: (1) On 08/27/2025 at 11:00 am, testing person #3 stated that urine drug screens were performed using the Horiba Yumizen C1200 analyzer; (2) A review of the operator's manual for the test system identified the manufacturer required the analyzer be operated at a humidity of 40-70% with no condensation; (3) A review of humidity records from March through May 2025 identified the humidity readings were less than 40% for 29 of 54 days of patient testing; (4) The records were reviewed with testing person #3 who stated on 08/27 /2025 at 11:00 am, the laboratory humidity had not been maintained as required by the manufacturer.