

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  10D2111958	<b>(X3) Date Survey Completed</b>  05/19/2026
<b>Name of Provider or Supplier</b>  Pain Relief Solutions, Llc	<b>Street Address, City, State</b>  1500 N University, Ste 101, Coral Springs, FL	
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>D0000</b>	An announced CLIA recertification survey was conducted at PAIN RELIEF SOLUTIONS, LLC from May 05 , 2026 to May 19, 2026. The laboratory was surveyed under 42 CFR Part 493 CLIA requirements. Standard deficiency cited as follows:
<b>D5217</b>	<p><b>EVALUATION OF PROFICIENCY TESTING PERFORMANCE</b> CFR(s): 493.1236(c)(1)</p> <p>At least twice annually, the laboratory must verify the accuracy of any test or procedure it performs that is not included in subpart I of this part.</p> <p>This STANDARD is not met as evidenced by: Based on record review and staff interview, the laboratory failed to verify the accuracy of testing methods at least twice annually for two out of eight analytes tested in urine toxicology screening test at least from 09/01/2024 to 05/01/2026. Findings included: 1-Review of the test menu provided with the form CMS-116 Clinical Laboratory Improvement Amendment (CLIA) Application for Certification signed by the Laboratory Director on 04/12/2026 indicated the following urine screening analyte tests: Amphetamine, Barbiturates, Benzodiazepines, Cocaine metabolites, Methadone, Opiates, Phencyclidine, Creatinine and pH. 2-Review of procedure "PROFICIENCY TESTING", under GENERAL COMPLIANCE: stated -Enroll in PT for every regulated analyte performed in-house. -Enroll in PT or perform alternative split sample testing as an external validation of all non-regulated analytes at least twice a year. 3-Review of American Proficiency Institute (API) Proficiency Testing (PT) attestation for first event of 2025 with date 05/25/2025, revealed that the laboratory failed to include the test of pH and Creatinine that is part of the urine adulteration package and failed to perform the twice a year accuracy verification for the Creatinine and pH. 4-The laboratory tested 7,828 patients from September 1, 2024 to May 1, 2026 for Creatinine and pH. 5-During an interview on 05/05/2026 at 12:00 PM the Laboratory Director confirmed that the laboratory failed to do twice a year</p>

verification for the screening test for the Creatinine and pH for the period of reference.