

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 31D2133848	(X3) Date Survey Completed 05/19/2026
Name of Provider or Supplier Revive Spine And Pain Center	Street Address, City, State 1001 Lincoln Drive West, Marlton, NJ	
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
D5221	<p>EVALUATION OF PROFICIENCY TESTING PERFORMANCE CFR(s): 493.1236(d)</p> <p>All proficiency testing evaluation and verification activities must be documented.</p> <p>This STANDARD is not met as evidenced by: Based on surveyor review of the Proficiency Testing (PT) records and interview with the General Supervisor (GS), the laboratory failed to review coded results for Toxicology Testing performed with the American Proficiency Institute (API) in the first and second events of 2025. The findings include: 1. The laboratory received a coded result (Not Graded 1) "See Data Summary" for Specific Gravity 1st event 2025 sample UAD-01, 02, 03. 2. The laboratory received a coded result (Not Graded 1) "See Data Summary" for Specific Gravity 2nd event 2025 sample UAD-04, 05, 06. 3. There was no documented evidence that coded PT results was reviewed. 4. The GS confirmed on 5/19/26 at 11:15 am that the laboratory did not review coded PT results.</p>
D5469	<p>CONTROL PROCEDURES CFR(s): 493.1256(d)(10)(g)</p> <p>(d)(10) Establish or verify the criteria for acceptability of all control materials. (d)(10) (i) When control materials providing quantitative results are used, statistical parameters (for example, mean and standard deviation) for each batch and lot number of control materials must be defined and available. (d)(10)(ii) The laboratory may use the stated value of a commercially assayed control material provided the stated value is for the methodology and instrumentation employed by the laboratory and is verified by the laboratory. (d)(10)(iii) Statistical parameters for unassayed control materials must be established over time by the laboratory through concurrent testing of control materials having previously determined statistical parameters.</p>

This STANDARD is not met as evidenced by:

Based on surveyor review of Quality Control (QC) records and interview with General Supervisor (GS), the laboratory failed to verify commercially assayed QC material with each new lot and/or shipment of Toxicology QC used on the Indiko Plus analyzer from 4/8/24 to the 5/19/26. The findings include: 1. There was no documented evidence that QC material was verified for the following Lot numbers: a) MAS total control level 3 lot DAT27053A b) MAS total control level 2 lot DAT27052A c) MAS total control level 4 lot DAT27054A d) MAS total control level 5 lot DAT27055A 2. The GS confirmed on 5/19/26 at 11:20 am that all assayed QC material was not verified before putting in use. Note: This was previously cited