

Statement of Deficiencies	(X1) Provider/Supplier/CLIA Identification Number 21D2149623	(X3) Date Survey Completed 05/08/2026
Name of Provider or Supplier Advanced Spine And Pain	Street Address, City, State 3720 Washington Boulevard Suite 100, Halethorpe, MD	
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
D5403	<p>PROCEDURE MANUAL CFR(s): 493.1251(b)</p> <p>(b) The procedure manual must include the following when applicable to the test procedure: (b)(1) Requirements for patient preparation; specimen collection, labeling, storage, preservation, transportation, processing, and referral; and criteria for specimen acceptability and rejection as described in 493.1242. (b)(2) Microscopic examination, including the detection of inadequately prepared slides. (b)(3) Step-by-step performance of the procedure, including test calculations and interpretation of results. (b)(4) Preparation of slides, solutions, calibrators, controls, reagents, stains, and other materials used in testing. (b)(5) Calibration and calibration verification procedures. (b)(6) The reportable range for test results for the test system as established or verified in 493.1253. (b)(7) Control procedures. (b)(8) Corrective action to take when calibration or control results fail to meet the laboratory's criteria for acceptability. (b)(9) Limitations in the test methodology, including interfering substances. (b)(10) Reference intervals (normal values). (b)(11) Imminently life-threatening test results, or panic or alert values. (b)(12) Pertinent literature references. (b)(13) The laboratory's system for entering results in the patient record and reporting patient results including, when appropriate, the protocol for reporting imminently life threatening results, or panic, or alert values. (b)(14) Description of the course of action to take if a test system becomes inoperable.</p> <p>This STANDARD is not met as evidenced by: Based on record review and interview, the laboratory's written procedure failed to provide step-by-step instructions for toxicology testing. Findings: 1. The Procedure for monthly review of toxicology test screening referred to the use of the AB Sciex analyzer, but the laboratory screened patient specimens with the Diatron Pictus analyzer. 2. The written procedure for the toxicology screen did not identify the calibrators used for each analyte and frequency of use. The technical supervisor stated</p>

that calibration was performed each week of patient testing, during interview on 5/15/26 at 12:00 PM.