Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	46D2084337	11/19/2025
Name of Provider or Supplier	**Street Address, City, State**	
Sacred Circle Healthcare Clinic	660 S 200 E, Ste 250, Salt Lake City, UT	

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
D5217	EVALUATION OF PROFICIENCY TESTING PERFORMANCE CFR(s): 493.1236(c)(1) 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. This STANDARD is not met as evidenced by: Based on record review and interview with the Technical Supervisor (TS), the laboratory failed to verify the accuracy of the tests twice annually for ten of ten analytes. The laboratory performs approximately 230,201 chemistry tests annually on the ABSciex 4500 Analyzer. Finding include: 1. A review of the laboratory Proficiency Test records revealed the laboratory failed to verify twice annually the accuracy of the analytes Cyclobenzaprine, Desipramine, Desmethyltramadol, Diphenhydramine, Doxepin, MDEA, Naltrexone, PCP, Nortriptyline and Tizanideine. 2. An interview on 11/19/2025, at 11:45 AM, with TS confirmed the laboratory failed to verify the accuracy of the tests twice annually for Cyclobenzaprine, Desipramine, Desmethyltramadol, Diphenhydramine, Doxepin, MDEA, Naltrexone, PCP, Nortriptyline and Tizanideine.
D5807	TEST REPORT CFR(s): 493.1291(d) (d) Pertinent "reference intervals" or "normal" values, as determined by the laboratory performing the tests, must be available to the authorized person who ordered the tests and, if applicable, the individual responsible for using the test results. This STANDARD is not met as evidenced by: Based on review of patient test reports, laboratory procedure manual, and an interview with the Technical Supervisor (TS), the laboratory failed to include the correct normal

	ranges for Hemoglobin (HGB) and Monocyte %. The laboratory performs approximately 19,630 hematology tests annually on the Sysmex XN-1000 Analyzer. Findings: 1. Review of patient reports revealed the HGB and Monocyte % reference range did not correctly match the reference range for HGB and Monocyte % in the ' Sysmex XN-1000 ' procedure manual. 2. The procedure manual showed the HGB reference interval was 12.0-16.0 g/dl and the Monocyte % reference interval was 4.1-12.7%. 3. The patient report showed the HGB reference interval was 11.8-15.1 g/dl and the Monocyte % reference interval was 4.1-12.4%. 4. An interview on 11/19 /2025, at 11:45 AM, with TS confirmed the laboratory failed to include the correct normal ranges for HGB and Monocyte %.