Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	13D0520334	03/23/2026
Name of Provider or Supplier	**Street Address, City, State**	
Family Practice Group, Pa	1951 Bench Rd #B, Pocatello, ID	

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
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 review of laboratory temperature logs, a direct observation,review of manufacturer's instructions, and interview with the laboratory lead on 3/23/2026, the laboratory failed to monitor room temperature in the secondary laboratory room where Afinon and Quidel/Ortho Sofia testing was being performed since the last inspection (4/23/2024). The findings include: 1. A review of laboratory temperature logs identified that the laboratory documented room temperature for the main laboratory but failed to document room temperatures for the secondary laboratory room for eight (8) months in 2024, 12 months in 2025 and three (3) months in 2026. 2. A direct observation in a secondary laboratory room on 3/23/2026 at 3:03 pm identified an Afinon analyzer used for back-up A1C testing, a Quidel Sofia2 analyzer and all reagents for three (3) of three (3) tests performed on the Sofia2: Sofia Strep A+ test cartridges, Sofia Influenza A+ B test cartridges and Sofia SARS test cartridges. 3. A review of the Sofia Strep A+, Influenza A+ B and SARS manufacturer's instructions identified storage temperatures of 59- 86 F. A review of the Afinon manufacturer's instructions identified and operating temperature of 59-89 F. 4. An interview with the laboratory lead on 3/23/2026 at 3:03 pm confirmed that the laboratory failed to monitor temperature in the secondary laboratory room.

D5429	MAINTENANCE AND FUNCTION CHECKS CFR(s): 493.1254(a)(1) (a)(1) Maintenance as defined by the manufacturer and with at least the frequency specified by the manufacturer. This STANDARD is not met as evidenced by: Based on a review of laboratory policies, microscope maintenance logs, and an interview with the laboratory lead on 3/23/2026, the laboratory failed to perform microscope maintenance 37 of 64 weeks in 2025 and 2026. The findings include: 1. A review of the laboratory policy, "Microscope Maintenance Log Instructions" identified that the policy states, "Perform microscope maintenance weekly." 2. A review of the microscope maintenance logs identified that the laboratory failed to document microscope maintenance 31 of 52 weeks in 2025 and six (6) of 12 weeks in 2026. 3. An interview with the laboratory lead on 3/23/2026 at 2:50 pm confirmed that the laboratory failed to document microscope maintenance for the above weeks. 4. The laboratory reports performing 985 microscopic tests annually.