Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	09D2177481	06/22/2023
Name of Provider or Supplier	**Street Address, City, State**	
Epiarx Pllc	4000 Albemarle St Nw Suite 208, Washington, DC	

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
D5821	TEST REPORT CFR(s): 493.1291(k) When errors in the reported patient test results are detected, the laboratory must do the following: (k)(1) Promptly notify the authorized person ordering the test and, if applicable, the individual using the test results of reporting errors. (k)(2) Issue corrected reports promptly to the authorized person ordering the test and, if applicable, the individual using the test results. (k)(3) Maintain duplicates of the original report, as well as the corrected report. This STANDARD is not met as evidenced by: Based on review of patient final reports and interview with the laboratory director (LD), The LD failed to ensure that patient reports were corrected in a timely manner prior to release of test results. Findings: 1.The laboratory performs Pathology reviews of patient slides to confirm diagnosis. 2.Review of six patient final reports showed that two of the six reports had errors that were not noticed until the day of the survey. 3.Review of Patient A final report showed that the specimen was received on July 14, 2021. The slide review was performed on 7/14/21 and test results were reported on 7 /19/21. Review of the patient electronic medical record showed that the patient's last name was spelled incorrect on the final report. 4.The LD stated that he was unaware that the last name was spelled incorrect on the report prior to the day of the survey. The LD performed a corrected report with the correct spelling of Patient A last name on the day of the survey. 5.Review of Patient B final report showed that the specimen was received on July 12, 2021. The slide review was performed on 7/ 6/2021 and the test results were reported on 7/6/2021. Review of the patient electronic medical record showed that the report date was 7/12/2021 and the receive date was 7/6/2021. 6.The LD stated that he was unaware that the receive and the report dates were incorrect on the report prior to the day of the survey. The LD performed a corrected report of the

	receive date and the report date on Patient B final report on the day of the survey. 7. The LD confirmed that he failed to ensure that patient reports were corrected in a timely manner prior to release of test results on the day of the survey.