Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	45D1069391	06/17/2026
Name of Provider or Supplier	**Street Address, City, State**	
Vitalogy Skincare Dba (Georgetown)	4513 Williams Drive, Georgetown, TX	

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
D0000	The Vitalogy Skincare DBA (Georgetown) laboratory was found to be in compliance with the Conditions of the CLIA regulations found at 42 CFR 493.1 through 493.1780, CLIA requirements for laboratories as a result of a recertification survey on 06/17/2026 and recertification is recommended. Standard level deficiencies were cited.
D5805	TEST REPORT CFR(s): 493.1291(c) (c) The test report must indicate the following: (c)(1) For positive patient identification, either the patient's name and identification number, or a unique patient identifier and identification number. (c)(2) The name and address of the laboratory location where the test was performed. (c)(3) The test report date. (c)(4) The test performed. (c)(5) Specimen source, when appropriate. (c)(6) The test result and, if applicable, the units of measurement or interpretation, or both. (c)(7) Any information regarding the condition and disposition of specimens that do not meet the laboratory's criteria for acceptability. This STANDARD is not met as evidenced by: I. Based on review of the patient testing logs, test report, and interview, the laboratory failed to ensure the test result was recorded in the patient's chart for 1 of 3 Potassium Hydroxide (KOH) test reports reviewed from Jan 2026 - May 2026. Findings follow. A. Review of the KOH Test Accession Log showed accession #K26-006 tested on 05 /08/2026 tested negative by KOH prep for fungal elements. B. Review of the Visit Note which also served as the test report for accession #K26-006 on 05/08/2026 showed the KOH test was not reported. C. Interview with the Clinic Manager on June 17, 2026 at 1140 hours in the break room confirmed the KOH test was performed but not recorded in the Visit Note which also served as the test report. II. Based on review of the test reports, quality control (QC) log, slides, and interview, the laboratory failed to report the result of the Mart-1 Immunohistochemical (IHC) stain for one of two

	Mohs test reports with Mart-1 stains reviewed from March 2025 - April 2026. Findings follow. A. Review of two Mohs cases with the Mart-1 stain showed the Visit Note for Mohs case VG26-152 tested on 04/28/2026 was missing the result of the Mart-1 stain. B. Review of the Mohs IHC Daily Quality Control log showed case # VG26-152 QC was reported as satisfactory. The column for "Stain results in Patient Report/Visit Note" did not indicate it was reported in the Visit Note. C. Review of the slides for case VG26-152 showed slides with the Mart-1 stain. D. Interview with the Laboratory Manager on July 17, 2026 at 1200 hours in the break room confirmed the result of the Mart-1 stain was not in the Visit Note and the column on the QC log was not completed to show it was included in the Visit Note.