Genome LLC
CLIA ID: 10D2192601
Laboratory Director: Jesus E. Viloria, M.D.

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PATIENT NAME	Crende Piko, Evelyn M	**ACCESSION**	22010700091
PATIENT DOB	Dec. 15, 1966	**ORDER CODE**	2022-0000655
PATIENT GENDER	F	**SAMPLE TYPE & SOURCE**	Anterior Nasal Swab
PATIENT PHONE	(407) 235-8483		
ACCOUNT	Hollywood - Medix Urgent Care	**COLLECTED**	01-07-2022, 03:51PM (EST)
PROVIDER	Yvette Fletcher-Prince, MD	**RECEIVED**	01-07-2022, 03:52PM (EST)
REPORT STATUS	FINAL	**REPORTED**	01-07-2022, 09:40PM (EST)

TEST INFORMATION Covid-19 PCR (rt-PCR) **DIAGNOSIS CODES** U07.1 COVID-19

QUALITATIVE RESULTS	
	RESULT
COVID	
SARS-CoV-2 qPCR Extracted N1	NOT DETECTED (NEGATIVE)

Covid-19 PCR (rt-PCR):
Positive/Detected results do not rule out other bacterial infections or co-infection with other viruses.
Laboratories within the United States and its territories are required to report all Positive/Detected results to the appropriate public health authorities.
Positive/Detected results may be due to past or present infection with non-SARS-CoV-2 coronavirus strains, such as coronavirus HKU1, NL63, OC43, or 229E.
Negative/Not Detected results do not preclude a SARS-CoV-2 infection and should not be used as the sole basis for patient management decisions.
This test is an In-Vitro Diagnostic (IVD) laboratory developed test (LDT) using FDA approved emergency use authorized (EUA) reagents for real-time reverse transcription polymerase chain reaction (RT-PCR or PCR) to detect the genetic presence of the Covid-19 SARS-CoV-2 virus and its variants in specimens obtained from nasopharyngeal or midturbinate (anterior) nasal swabs.
This test was performed by a laboratory certified under the Clinical Laboratory Improvement Amendments (CLIA) of 1988, 42 U.S.C. §263a, to be qualified to perform High Complexity Testing.
Covid-19 PCR (rt-PCR):
This test was performed by Pinnacle Genetics Lab CLIA ID 10D2226941

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These tests were developed and their performance characteristics determined by Genome LLC.
They have not been cleared or approved by the FDA.