

Name: Enwu Guan | DOB: 12/9/1969 | MRN: 103473972 | PCP: | Legal Name: Enwu Guan

CEPHEID COVID-19 RSV INFLUENZA A/B PCR - Details

Cepheid COVID-19 RSV Influenza A/B PCR

Status: Final result (Collected: 1/20/2023 9:50 AM)

Results

Patient Demographics

Patient Name	Legal	DOB	Address	Phone
Guan, Enwu	Sex	12/9/1969	400 Claremont Avenue	+8615194193689
(<E8304999>)	Male		Jersey City NJ 07304	(Mobile)

Cepheid COVID-19 RSV Influenza A/B PCR

Order: 451232307

Status: Final result Visible to patient: Yes (seen) Next appt: None

Dx: Encounter for screening for COVID-19

Component	Ref Range & Units	1/20/23 9:50 AM
Influenza A PCR	Negative	Negative
Influenza B PCR	Negative	Negative
RSV	Negative	Negative
SARS-CoV-2 PCR	Negative	Negative

Resulting Agency

DH RIVERSIDE LABORATORY, STD CLINIC
LABORATORY AT RIVERSIDE, CLIA:
33D0909985, CLEP/PFI: 7478

Narrative

Performed by: DH Riverside

DH RIVERSIDE C19

Specimen source is Nasopharyngeal Swab.
Results from Accession 42-23-018-00041

POSITIVE results indicate that viral RNA was detected. Positive results are indicative of active infection but do not rule out bacterial infection or co-infection with other viruses; clinical correlation with patient history and other diagnostic information is necessary to determine patient infection status. The agent detected may not be the definite cause of disease.

NEGATIVE results indicate that viral RNA was not detected. Negative results do not preclude SARS-CoV-2, influenza A virus, influenza B virus and/or respiratory syncytial virus (RSV) infection and should not be used as the sole basis for diagnosis, treatment or other patient management decisions. Negative results must be combined with clinical observations, patient history, and/or epidemiological information. Negative results in asymptomatic individuals cannot be used as definite evidence that an individual has not been exposed to SARS-CoV-2, influenza viruses, or RSV and has not been infected with any of these viruses.

INVALID result indicates that specimen was tested, but either did not meet acceptance criteria for quality control or could not be processed by the instrument. Please submit a new specimen for testing.

NOTE:

This test has been authorized by the FDA on September 10, 2021 for emergency use by authorized laboratories only for the simultaneous qualitative detection and differentiation of SARS-CoV-2, influenza A, influenza B, and respiratory syncytial virus (RSV) until emergency authorization is terminated or revoked.

Please refer to the following fact sheets for further information:

Fact Sheet for Healthcare providers -

<<https://www.fda.gov/media/152162/download>>

Fact Sheet for Patients - <<https://www.fda.gov/media/152166/download> >

Testing performed on Cepheid XpertXpress platform.

Specimen Collected: 01/20/23 9:50 AM

Last Resulted: 01/20/23 11:28 AM

Lab Information

Lab

DH RIVERSIDE LABORATORY, STD CLINIC LABORATORY AT RIVERSIDE, CLIA: 33D0909985, CLEP/PFI: 7478

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