Name: Guanglan Wei | DOB: 7/26/1972 | MRN: 103473968 | PCP: | Legal Name: Guanglan Wei

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

## Cepheid COVID-19 RSV Influenza A/B PCR

Results

Status: **Final result (Collected: 1/20/2023 9:51 AM)**

### Patient Demographics

| Patient Name | Legal Sex | DOB | Address | Phone |
|---|---|---|---|---|
| Wei, Guanglan (<E8304995>) | Female | 7/26/1972 | 400 Claremont Avenue Jersey City NJ 07304 | 646-403-7894 (Mobile) |

### Cepheid COVID-19 RSV Influenza A/B PCR

Order: 451233317

Status: **Final result** Visible to patient: **Yes (seen)**

Dx: **Encounter for screening for COVID-19**

| Component | Ref Range & Units | 1/20/23 9:51 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 42-23-018-00042

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 processes 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 9:51 AM Last Resulted: 01/20/23 11:22 AM

## Lab Information

Lab
DH RIVERSIDE LABORATORY, STD CLINIC LABORATORY AT RIVERSIDE, CLIA: 33D0909985, CLEP/PFI: 7478
160 West 100th Street
NEW YORK NY 10025