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Limited Viral Respiratory Panel (LVRP) Report

Patient Information

Name: Test Patient
DOB: 02.28.1996
Gender: M

Specimen Information

Accession Number: 000413
Date Collected: 03.13.2023
Date Received: 03.14.2023
Report Date: 03.14.2023
Sample Type: Anterior or Mid-Turbinate Nasal Swab

Facility Information

Facility Name: House Account
Provider Name: Test Provider
Address:

Panel: Limited Viral Respiratory Panel (LVRP)

Organism(s) / Gene(s) Detected	Results
MS2	Detected
Respiratory Syncytial Virus (RSV)	Positive

Tested Organisms / Genes and Results

Panel: Limited Viral Respiratory Panel (LVRP)

Organism / Gene	Results
Influenza A/B	Negative
MS2	**Detected**
Respiratory Syncytial Virus (RSV)	**Positive**

Processing and Detection Methodology:
Limitation: An absence of detection does not imply the absence of microorganisms other than that listed or does not exclude the possibility that the target sequence is present below the limit of detection.
Methodology: The Limited Respiratory Pathogen Panel utilizes Real-time PCR technology to detect the presence of genes associated with pathogens via TaqMan® chemistry.

Disclaimer: These tests were developed and characterized by Mokan Labs and interpreted by Mokan Labs. The tests in the Respiratory Pathogen Panel have not been approved by the Food and Drug Administration. The FDA has determined that such approval is not necessary, provided that the laboratory both (1) maintains its good standing as a clinical testing laboratory with all mandatory accrediting bodies, and (2) continually demonstrates that its testing protocols and procedures achieve a high degree of analytical accuracy.

This report, associated with order #000413, has been approved by the following reviewers:

Comment: Test samples, Reports are for fake patient

Report Reviewer:

Electronically signed and dated on 03.14.2023 10:11
Domenick Palmieri