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IHDLAB
Innovative Health Diagnostics

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|---|---|---|---|---|---|
| **Patient:** | **Test, Patient** | | | **Acc #:** | **2438234** |
| Patient #: | MDT-279063 | Birth: | 1/1/1990 | | |
| Doctor: | Test, Provider | Age: | 34 years | Collection Date: | 9/10/2024 8:00 AM |
| | | Gender: | Female | Received in Lab: | 9/10/2024 11:13 AM SB |

| Test Name | In Range | Out of Range | Flag | Units | Reference Range |
|---|---|---|---|---|---|
| **Respiratory Pathogen Panel (RPP), BioFire (RP2.1)** | | | | | *Run by: SB on 9/10/2024 11:15 AM* |
| Adenovirus | Not Detected | | | | Not Detected |
| Bordetella parapertussis (IS1001) | Not Detected | | | | Not Detected |
| Bordetella pertussis | Not Detected | | | | Not Detected |
| Chlamydia pneumoniae | Not Detected | | | | Not Detected |
| Human Coronavirus 229E | Not Detected | | | | Not Detected |
| Human Coronavirus HKU1 | Not Detected | | | | Not Detected |
| Human Coronavirus NL63 | Not Detected | | | | Not Detected |
| Human Coronavirus OC43 | Not Detected | | | | Not Detected |
| Human Metapneumovirus (hMPV) | Not Detected | | | | Not Detected |
| Influenza A (Pan) | Not Detected | | | | Not Detected |
| Influenza A/H1 | Not Detected | | | | Not Detected |
| Influenza A/H3 | Not Detected | | | | Not Detected |
| Influenza A (no subtype detected) | Not Detected | | | | Not Detected |
| Influenza B | Not Detected | | | | Not Detected |
| Human Parainfluenza Virus 1 | Not Detected | | | | Not Detected |
| Human Parainfluenza Virus 2 | Not Detected | | | | Not Detected |
| Human Parainfluenza Virus 3 | Not Detected | | | | Not Detected |
| Human Parainfluenza Virus 4 | Not Detected | | | | Not Detected |
| Mycoplasma pneumoniae | Not Detected | | | | Not Detected |
| Respiratory Syncytial Virus | Not Detected | | | | Not Detected |
| SARS-CoV-2 | Not Detected | | | | Not Detected |

***Notes:*** ***Methodology: Innovative Health Diagnostics' FilmArray Respiratory panel is a qualitative multiplexed nucleic acid amplification test (NAAT). It is in-vitro diagnostic test intended for detection of microbial nucleic acids in Nasopharyngeal specimens.***

***Limitations: This panel is FDA approved for Diagnostics Testing in Laboratories Certified to Perform Moderate-High Complexity Testing under CLIA. It does not provide a quantitative value for the organism(s) in the sample. The detection of viral and bacterial nucleic acid is dependent upon proper specimen collection, handling, transportation, storage and preparation. Failure to observe proper procedures can lead to incorrect results. Results from this test must be correlated with the clinical history, epidemiological data, and other data available to the clinician evaluating the patient.***

Innovative Health Diagnostics is an MD Tox Laboratory Service Line
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