

### CLINIC INFORMATION

Name: Assurance Pathology Laboratory  
 Address: 2868 Acton Road Suite 207 BIRMINGHAM, AL 35243  
 Provider: Doctor Test

### PATIENT INFORMATION

Name: TEST, TEST  
 DOB: 5/12/1965  
 Sex: M  
 Phone: (405) 000-0000

### SPECIMEN INFORMATION

Lab Accession Number: ASL-104628  
 Date Collected: 4/27/2020  
 Date Received by Lab: 4/28/2020  
 Date Accessioned: 4/28/2020  
 Date Reported: 4/28/2020  
 Faxed to: 18777966185

### Controls

Patient Extraction Control	1	PASS
Endogenous Positive Control	2	PASS
Pathogen Positive Control	3	PASS
Pathogen Negative Control		PASS

- (1) Endogenous control confirms sample collection, DNA/RNA extraction, and assay enzyme activity
- (2) Positive control is synthetic inactive pathogen
- (3) Negative Control contains primers, probe, and enzymes with no DNA/RNA template
- (4) A "Detected" result indicates the presence of a pathogen (99.99% confidence) above the assay cutoff.  
 Assay cutoff is represented by CFU (bacteria), PFU (viruses) or Copy Number (DNA)

### Test Performed

Lab Result(4)  
 (Qualitative Low/Medium/High)

DNA Copy Number

Comments

Test Performed	Lab Result(4) (Qualitative Low/Medium/High)	DNA Copy Number	Comments
<b>COVID-19</b>			<b>Collection Type: Respiratory</b>
COVID-19	DETECTED	DETECTED	[4/28/2020 8:29 AM] The test has been internally validated, but FDA's independent review of this validation is pending. Contains Nonbinding Recommendations. Laboratory will immediately notify appropriate Federal, State, or local public health agencies of all positive results.
<b>SUMMARY COVID-19 Respiratory Panel</b>			<b>Collection Type: Respiratory</b>
Haemophilus influenzae	DETECTED - MEDIUM	1.00E+03	[4/28/2020 8:28 AM] Limitation: Test does not differentiate between patient with acute infection or an asymptomatic carrier.

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Lab Director: Ty Thomas, MD

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This test detects the presence of pathogen and must be evaluated with clinical symptoms to diagnose disease. All tests established and validated by Laboratory and not FDA approved.  
 Methodology statement: Real-time PCR assays are designed to detect pathogens with clinical significance and analytical sensitivity and specificity greater than 99%.  
 Limitations: While these assays are very sensitive and specific, theoretically these assays could detect pathogens not listed, resulting in a false positive.  
 In addition, while these assays are very specific, there may be target pathogen sequences with unknown sequence variability which may not be detected, resulting in a false negative result.

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Test Performed	Lab Result(4) (Low/Medium/High)	DNA Copy Number	Comments
<b>COVID-19 Respiratory Panel</b>			
Influenza A	Not Detected	Not Detected	
Influenza B	Not Detected	Not Detected	
Streptococcus pyogenes (Group A)	Not Detected	Not Detected	
Mycoplasma pneumoniae	Not Detected	Not Detected	
Moraxella catarrhalis	Not Detected	Not Detected	
<b>Haemophilus influenzae</b>	<b>DETECTED - MEDIUM</b>	<b>1.00E+03</b>	[4/28/2020 8:28 AM] Limitation: Test does not differentiate between patient with acute infection or an asymptomatic carrier.

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