



3611 14TH AVE SUITE 102  
 BROOKLYN, NY 11218-3750  
 Tel. (718)851-5773 Fax. (718)851-3919  
 Laboratory Director: **Samar Roy, PhD**  
 CLIA#: 33D0985206



**Client: MASTERS, KEN MD** 99  
 3611 14th Avenue STE 220  
 BROOKLYN, NY 11218  
 (718) 851-5773 Fax: (718) 851-3919  
**Phys:**

**Patient: IDR-06, SAMPLE-01**  
**DOB:** 07/09/2024 **Age:** 0 **Sex:** M  
**Phone:** ( ) - **ID#:** A2407090011

<b>Accession:</b> 2407090011	<b>Coll. Date:</b> 07/09/24	<b>Recv. Date:</b> 07/09/24	<b>Print Date:</b> 08/06/24
<b>Chart#</b>	<b>Coll. Time:</b> 12:33	<b>Recv. Time:</b> 12:33	<b>Print Time:</b> 13:39
<b>First reported on:</b>	07/10/24	<b>Final report date:</b>	07/10/24

**Clinical Report:**

**Clinical Abnormalities Summary:** (May not contain all abnormal results; narrative results may not have abnormal flags. Please review entire report.)

**Coronavirus NL63** **POSITIVE**  
**Mycoplasma pneumoniae** **POSITIVE**

Test Name	Within Range	Out Of Range	Ref.Range	Units	Previous Result (Date)
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**GENETIC AND MOLECULAR TESTING**

SARS-CoV-2 RT-PCR **NEGATIVE**

The SARS-CoV-2 test is intended for the qualitative detection of nucleic acid from SARS-CoV-2( who meet COVID-19 clinical and/or epidemiological criteria. For lower respiratory tract specimens, the assay is submitted for authorization by FDA under the Emergency Use Authorization (EUA).

Test Methodology:  
 Real Time PCR Detection of SARS-CoV-2 using Qiagen QIAstat-Dx.

Disclaimer:  
 This test has not been Food and Drug Administration (FDA) cleared or approved and has been authorized by FDA under an Emergency Use Authorizatio (EUA). The test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of SARS-CoV-2 under Section 564(b)(1) of the Act, 21 U.S.C. section 360bbb-3(b)(1), unless the authorization is terminaed or revoked sooner. Alliance Laboratories is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C section 263a, to perform high complexity tests.

Fact Sheet for Healthcare Providers:  
<https://www.cdc.gov/coronavirus/2019-nCoV/hcp/index.html>

Fact Sheet for Patients:  
<https://www.cdc.gov/coronavirus/2019-ncov/your-health/index.html>

Adenovirus	NEGATIVE
Coronavirus 229E	NEGATIVE
Coronavirus HKU1	NEGATIVE
<b>Coronavirus NL63</b>	<b>POSITIVE</b>
Human Metapneumovirus A+B	NEGATIVE
Influenza A	NEGATIVE

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**GENETIC AND MOLECULAR TESTING (Continued)**

Influenza A H1	NEGATIVE
Influenza H1N1 pdm09	NEGATIVE
Influenza A H3	NEGATIVE
Influenza B	NEGATIVE
Parainfluenza virus 1	NEGATIVE
Parainfluenza virus 2	NEGATIVE
Parainfluenza virus 3	NEGATIVE
Parainfluenza virus 4	NEGATIVE
Resp. Syncytial Virus A+B	NEGATIVE
Rhinovirus/Enterovirus	NEGATIVE
Bordetella pertussis	NEGATIVE
Chlamydomphila pneumoniae	NEGATIVE
<b>Mycoplasma pneumoniae</b>	<b>POSITIVE</b>
Coronavirus OC43	NEGATIVE
SARS-CoV-2 Collection	NP SWAB

**Report Status: FINAL**

**END OF REPORT**