LabCorp

Patient Report

Specimen ID: 181-992-3201-0
Control ID:

SAMPLE REPORT, 164055

Acct #: 90000999 **Phone:** (336) 436-8645 **Rte:** 00
LabCorp Test Master
Test Account
5450 Millstream Road
MCLEANSVILLE NC 27301

Patient Details
DOB: 01/10/1980
Age(y/m/d): 040/05/19
Gender: F
Patient ID:

Specimen Details
Date collected: 06/29/2020 0000 Local
Date received: 06/29/2020
Date entered: 06/29/2020
Date reported: 06/29/2020 0000 ET

Physician Details
Ordering:
Referring:
ID:
NPI:

General Comments & Additional Information

Clinical Info: POSITIVE REPORT

Ordered Items

SARS-CoV-2 Antibody, IgG

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
SARS-CoV-2 Antibody, IgG [A]	Positive	Abnormal		Negative	01

Comments:

[A] This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner. This test has been authorized only for detecting the presence of antibodies against SARS-CoV-2, not for any other viruses or pathogens.

01	BN	LabCorp Burlington 1447 York Court, Burlington, NC 27215-3361	Dir: Sanjai Nagendra, MD

For inquiries, the physician may contact **Branch: 800-222-7566 Lab: 800-762-4344**

LabCorp

Patient Report

Specimen ID: 181-992-3202-0
Control ID:

SAMPLE REPORT, 164055

Acct #: 90000999 **Phone:** (336) 436-8645 **Rte:** 00
LabCorp Test Master
Test Account
5450 Millstream Road
MCLEANSVILLE NC 27301

Patient Details
DOB: 01/10/1980
Age(y/m/d): 040/05/19
Gender: F
Patient ID:

Specimen Details
Date collected: 06/29/2020 0000 Local
Date received: 06/29/2020
Date entered: 06/29/2020
Date reported: 06/29/2020 0000 ET

Physician Details
Ordering:
Referring:
ID:
NPI:

General Comments & Additional Information
Clinical Info: NEGATIVE REPORT

Ordered Items
SARS-CoV-2 Antibody, IgG

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
SARS-CoV-2 Antibody, IgG [A]	Negative			Negative	01

Comments:
[A] This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner. This test has been authorized only for detecting the presence of antibodies against SARS-CoV-2, not for any other viruses or pathogens.

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