ASPIRA LABS®

Patient Info
Last Name: SAMPLE
First Name: PATIENT
MRN: 12345
DOB: 07/01/1980
Age: 40
Gender: FEMALE
Ethnicity: African American

Provider Info
Ordering Provider: PROVIDER
Practice Name: OBGYN OFFICE
Street Address: 123 Main Street
City, State, Zip: ANYTOWN, ST 01234
Tel: 555-555-5555
Fax:
Copy-to-Physician:
Fax:

Laboratory Info
Accession No: A0000789
Client No: 1234A
Collection Date: 06/02/2020
Received Date: 06/03/2020
Final Report Date: 06/04/2020

Clinical Information:

Menopausal Status: ✓ Premenopausal

Test Name	Result	Status	Reference Range
SARS-CoV-2 Antibody	Negative	**Negative**	n/a

Interpretation: This sample does not contain detectable SARS-CoV-2 IgG antibodies. This negative result does not rule out SARS-CoV-2 infection. Clinical correlation is recommended. Results should not be used as the sole basis to diagnose or exclude recent SARS-CoV-2 infection.

Notes:

Methods and Limitations

Elecsys® Anti-SARS-CoV-2 is an immunoassay intended for qualitative detection of antibodies to SARS-CoV-2 in human serum and plasma (K2-EDTA, K3-EDTA, Li-heparin). The test is intended for use as an aid in identifying individuals with an adaptive immune response to SARS-CoV-2, indicating recent or prior infection. The electrochemiluminescence immunoassay "ECLIA" is intended for use on cobas e immunoassay analysers. The assay uses a recombinant protein representing the nucleocapsid (N) antigen for the determination of antibodies against SARS-CoV-2.

Disclaimer

Serum or plasma samples from the early (pre-seroconversion) phase of illness can yield negative findings. Therefore, this test cannot be used to diagnose an acute infection. Over time, titers may decline and eventually become negative. Testing with a molecular diagnostic should be performed to evaluate for active infection in symptomatic individuals. It is not known at this time if the presence of antibodies to SARS-CoV-2 confers immunity to reinfection. False positive results for the Elecsys Anti SARS CoV 2 assay may occur due to cross reactivity from pre-existing antibodies or other possible causes. For Emergency Use Authorization only.

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