TEST CHANGE

COVID-19 IgG, Qualitative by CIA

3002776, COV19QUALG

Specimen Requirements:	
Patient Preparation:	
Collect:	Serum separator tube (SST) or EDTA plasma
Specimen Preparation:	Separate serum or plasma from cells ASAP or within 2 hours of collection. Transfer 0.5mL serum or plasma to an ARUP Standard Transport Tube. (Min: 0.25 mL)
Transport Temperature:	Refrigerated
Unacceptable Conditions:	Grossly hemolyzed, grossly icteric, or severely lipemic specimens. Postmortem specimens.
Remarks:	Preferred: ARUP Standard Transport Tube for specimen submission (ARUP Item# 15824).
Stability:	Refrigerated: 1 week; Frozen: 1 month
Methodology:	Qualitative Chemiluminescent Immunoassay (CLIA)
Performed:	Mon, Wed, Fri ~~Sun-Sat~~
Reported:	1-5 days
Note:	
CPT Codes:	86769
New York DOH Approval Status:	This test is New York DOH approved.

Interpretive Data:

The COVID-19 IgG, Qualitative by CIA test is for in vitro diagnostic use under an FDA Emergency Use Authorization (EUA). In compliance with this authorization, please visit https://www.aruplab.com/infectious-disease/coronavirus/testing for more information and to access the applicable information sheets. This test should not be used for screening of donated blood.
Use for the qualitative detection of IgG antibodies against the nucleocapsid protein of SARS-CoV-2 (COVID-19) that develop in response to natural infection with SARS-CoV-2. These antibodies do not develop as a result of a COVID-19 vaccination. There are no current recommendations for assessing COVID-19 vaccine response.

Reference Interval:

Negative