

Post – Test Report COVID-19 Virus Antigen Test Results available in 15/20 minutes

Full Name:	Date of Birth:
Signature:	Email:
Home Address:	
City:	State: Zip Code:
Phone Number:	Current Date:
Sex: Race:	Ethnicity:
 I authorize Dripp IV Therapy to notify me of my results by phone is necessary Results by Premier Biotech COVID-19 IgG/IgM Rapid Test Cassette Results by CareStart™ Rapid Diagnostic Test for Detection of SARS-CoV-2 Antigen 	
Positive for COVID-19. Negative for COVID-19.	Positive for IgG and IgM Antibodies Negative for IgG and IgM Antibodies
Nurse:	
Signaturo	



Require Disclosure: Laboratory test results should always be considered in the context of clinical observations and epidemiological data (such as local prevalence rates and current outbreak/epicenter locations) in making a final diagnosis and patient management decisions. Patient management should be made by a healthcare provider and follow current CDC guidelines. In the USA, this test has not been FDA cleared or approved; this test has been authorized by FDA under an EUA for use by authorized laboratories; use by laboratories certified under the CLIA,42 U.S.C. \Box 253a,that meet requirements to perform moderate, high or waived complexity tests. This test is authorized for use at the Point of Care(POC),i.e, in patient care settings operating under a CLIA certification of Waiver, Certificate of Compliance, or Certificate of Accreditation. This test has been authorized only for detection of protein from SARS-CoV-2, not for any other viruses or pathogens. In the USA,- this test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of the vitro diagnostics for detection and/or diagnosis of the virus that causes COVID-19 under Section 564(b)(1) of the Federal Food, Drug and Cosmetic Act,21 U.S.C. \Box 360bbb-3(b)(1), unless the authorization is termination or revoked sooner.