SARS-CoV-2 Antibody Immunoassay Kit (GICA)

[Product Name]

SARS-CoV-2 Antibody Immunoassay Kit (GICA)

Catalog Number: XG200H1

Package Size:50T

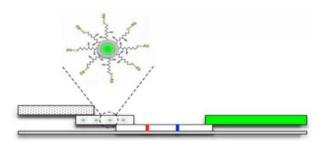
[Intended use]

The SARS-CoV-2 Antibody Immunoassay Kit (GICA) is used for the qualitatively determination of novel coronavirus (SARS-CoV-2) antibodies in human blood samples (plasma, serum) in vitro.

It is suitable for the rapid screening of SARS-CoV-2 virus antibodies for patients, employees, students and residents in medical institutions at all levels, which can realize rapid and accurate serological test.

[Principle]

This kit uses the principle of colloidal gold labeling technology, immunochromatographic technology and double antigen sandwich method, the binding pad is fixed with labeled colloidal gold SARS-CoV-2 antigen I, and the analysis membrane is coated with SARS-CoV-2 antigen II. During the test, if the sample contains SARS-CoV-2 virus antibody (including IgM and IgG), the virus antibody first binds to the colloidal gold-labeled antigen II in the binding pad, and then combines with the coated antigen II in analysis membrane to form a "colloidal gold-labeled antigen I- SARS-CoV-2 virus antibody-antigen II" in T-line and the "colloidal gold-labeled antigen I" that has not captured the virus in the binding pad will flow through the T-band and bind to the coated antibody in C-line. Test results are determined by reagent card color reaction.



【Kit components】

Size	Regents	Quantity
50T	Immunoassay card	50pieces
	Sample dilution buffer	30mL

[Storage conditions]

- 1. The kit remains stable for 12 months if stored at 4-30 $^{\circ}$ C. Please store the Sample dilution buffer at 2-8 $^{\circ}$ C after opening.
- 2. The unpacked Immunoassay card should be used as soon as possible.

[Sample requirements]

1. Sample type

The kit is applicable for human plasma and serum.

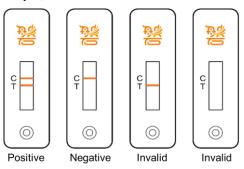
- 2. Sample collection and processing
- a) Plasma/serum collection: Serum/plasma samples are used right way or

stored at 2-8 $\,^{\circ}$ C for less than one week, or kept at -20 $\,^{\circ}$ C for no more than 1 month, and avoid repeated freeze-thaw cycles.

[Protocol]

- 1. Before testing, the sample, immunoassay card and sample dilution buffer must be equilibrated to room temperature (20–30 $^{\rm o}{\rm C}$) .
- $2.Mix\ 30\mu l$ of the plasma/serum sample with $70\mu l$ of sample dilution buffer. (The samples should be tested as soon as possible after processing)
- 3. Load $100\mu l$ of the diluted sample to the sample well of the immunoassay card;
- 5.After 15 minutes, test results were judged by the color reaction of the immunoassay card.

[Data analysis]



This result is for reference only, and the diagnosis result of the patient needs to be judged in conjunction with the clinical diagnosis.

【Limitations of the method】

- 1. Results beyond the measurement range of the kit are unreliable.
- Severe hemolysis, chyle, and bilirubin samples can cause abnormal test results.

[Caution]

- 1. This reagent is intended for in vitro diagnostic use only.
- Leakage and moisture absorption of the vacuum packaged immunoassay card may cause abnormal test results.
- 3. Reagents of different batches and varieties must not be mixed.
- 4.Please store and use reagents reasonably in strict accordance with the instructions.
- 5. Avoid direct sunlight.
- Testing must comply with the requirements of the Biosafety Code and strictly prevent cross-infection.
- 7. All samples, washing liquids and various wastes should be treated as potentially hazardous substances.

If You have Problems, please contact:

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