T030-EN.01

Anti-SARS-CoV-2 (B.1.351) Antibody IgG Titer Serologic Assay Kit (Spike Trimer)

Pack Size: 96 tests

Catalog Number： RAS-T030

IMPORTANT: Please carefully read this manual before performing your experiment.

For Research Use Only. Not For Use In Diagnostic Or Therapeutic Procedure

INTENDED USE

This kit is developed for detecting Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike Trimer) in human serum samples. It is intended for research use only (RUO).

PRINCIPLE OF THE ASSAY

The newly identified Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) has posed a serious threat to human health. A rapid and effective Assay kit detecting the levels of Anti-SARS-CoV-2 in human serum can facilitate research on characterization of antibodies produced in response to SARS-CoV-2 infection.

This assay kit is used to measure the titer of Anti-SARS-CoV-2 Antibody IgG by employing an indirect ELISA. Immobilize SARS-CoV-2 Spike Trimer(B.1.351) on the microplate. Then add the samples, incubate and wash the wells. Next add Secondary antibody HRP-Anti-Human IgG to the plate, incubate and wash the wells. Lastly load the substrate into the wells and monitor color development in proportion with the amount of antibody present. The reaction is stopped by the addition of a stop solution and the intensity of the absorbance can be measured at 450 nm and 630 nm. The OD Value reflects the amount of antibody bound.

MATERIALS PROVIDED

TABLE 1. MATERIALS PROVIDED

Catalog	Components	Size (96 tests)	Format	Storage	
				Unopened	Opened
RAS030-C01	Pre-coated SARS-CoV-2(B.1.351) Spike Trimer Microplate	1 plate	Solid	2-8℃	2-8℃
RAS030-C02	Positive Control	100 µL	Liquid	2-8℃	2-8℃
RAS030-C03	Negative Control	100 µL	Liquid	2-8℃	2-8℃
RAS030-C04	HRP-Anti-Human IgG	200 µL	Liquid	2-8℃,avoid light	2-8℃,avoid light
RAS030-C05	10xWashing Buffer	50 mL	Liquid	2-8℃	2-8℃
RAS030-C06	Dilution Buffer	50 mL	Liquid	2-8℃	2-8℃
RAS030-C07	Substrate Solution	12 mL	Liquid	2-8℃,avoid light	2-8℃,avoid light
RAS030-C08	Stop Solution	7 mL	Liquid	2-8℃	2-8℃

US and Canada: Tel: +1 800-810-0816
Asia and Pacific: Tel: +86 400-682-2521
Web: http://www.acrobiosystems.com
E-mail: order@acrobiosystems.com

REAGENTS/EQUIPMENT NEEDED BUT NOT SUPPLIED

Single or dual wavelength microplate reader with 450 nm and 630 nm filter;

Centrifuge;

37ºC Incubator;

10 µL, 200 µL and 1000 µL precision pipettes;

10 µL, 200 µL and 1000 µL pipette tips;

Multichannel pipettes;

Tubes;

Graduated cylinder to prepare Wash Solution;

Deionized or distilled water to dilute 10× Washing Buffer;

SHIPPING AND STORAGE

1. The unopened kit is stable for 12 months from the date of manufacture if stored at 2°C to 8°C.
2. The opened kit should be stored per TABLE 1. The shelf life is 30 days from the date of opening.
3. The kit shipped at room temperature that had been validated. Please contact us if you need blue ice shipping, but additional freight may be followed.

Note: *a. Do not use reagents past their expiration date.*

b. Find the expiration date on the outside packaging.

REAGENT PREPARATION

Bring all reagents and samples to room temperature (20℃-25℃) before use. If crystals have formed in buffer solution, warm until the crystals have completely dissolved and bring solution back to room temperature before use.

RECOMMENDED SAMPLE PREPARATION

1. Working fluid preparation

1.1 Preparation of 1×Washing Buffer:

Dilute 50 mL 10×Washing Buffer with ultrapure water/deionized water to 500 mL.

1.2 Preparation of HRP-Anti-Human IgG working fluid:

Dilute HRP-Anti-Human IgG at 1:100 with Dilution Buffer. The prepared working fluid should avoid light. Please

US and Canada: Tel: +1 800-810-0816 | Web: http://www.acrobiosystems.com
Asia and Pacific: Tel: +86 400-682-2521 | E-mail: order@acrobiosystems.com

prepare it for one-time use only.

1.3 Preparation of Positive Control and Negative Control working fluid and pre-treatment of samples:

a. For qualitative detection of antibodies:

Dilute the samples,Positive Control and Negative Control at 1:800 with Dilution Buffer.

b. For determination of antibody titer:

It is recommended to dilute the samples,Positive Control and Negative Control from 1:800-1:25600 with Dilution Buffer.

2. Plate set up

Number the diluted samples corresponding to the wells of the Pre-coated SARS-CoV-2(B.1.351) Spike Trimer Microplate. Each experiment requires a set of Positive Control and Negative Control working fluid.

3. Add Samples

Add 100 µL diluted sample,Positive Control and Negative Control working fluid to the corresponding wells. Seal the plate with microplate sealing film and incubate at 37℃ for 1.0 h.

4. Washing

Remove the remaining solution by aspiration, add 300 µL of 1×Washing Buffer to each well, gently tap the plate for 1 min, remove any remaining 1×Washing Buffer: by aspirating or decanting, invert the plate and blot it against paper towels. Repeat the wash step above for three times.

5.Add HRP-Anti-Human IgG

For all wells, add 100 µL HRP-Anti-Human IgG working solution.Seal the plate with microplate sealing film and incubate at 37℃ for 1 hour, avoid light.

6. Washing

Repeat step 4.

7. Substrate Reaction

Add 100 µL **Substrate Solution** to each well. Seal the plate with microplate sealing film and incubate at 37℃ for 20 min, avoid light.

US and Canada: Tel: +1 800-810-0816 | Web: http://www.acrobiosystems.com
Asia and Pacific: Tel: +86 400-682-2521 | E-mail: order@acrobiosystems.com

8. Termination

Add 50 μL **Stop Solution** to each well, and tap the plate gently for 3 min to allow thorough mixing.

Note: *the color in the wells should change from blue to yellow.*

9. Data Recording

Read the absorbance at 450 nm and 630 nm using UV/Vis microplate spectrophotometer.

Note*: To reduce the background noise, substract the value read at* $OD_{450\ nm}$ *with the value read at* $OD_{630\ nm}$.

CUT-OFF VALUE IDENTIFICATION

Cut-off value =0.1.

Note：The cut-off value can be determined by the end user.

INTERPRETION OF RESULTS

a. For qualitative detection of antibodies:

Positive reading: Percent inhibition of sample ≥ Cut-off value means Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike Trimer) are detected.

Negative reading: Percent inhibition of sample ＜Cut-off value means Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike Trimer) are notdetected.

b. For determination of antibody titer:

Determination of antibody titer: the positive sample was diluted with a gradient, and the antibody titer of the sample corresponds to the highest dilution factor that still yields a positive reading.

LIMITATIONS OF THE PROCEDURE

This test is designed for detecting human serum of Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike Trimer). However, we do not have the LoQ (Limit of Quantitation) and ULMI (upper limit of measuring interval) and cutoff defined for semi-quantitative detection. Interested customer is recommended to establish the semi-quantitative detection procedure themselves.

US and Canada: Tel: +1 800-810-0816 | Web: http://www.acrobiosystems.com
Asia and Pacific: Tel: +86 400-682-2521 | E-mail: order@acrobiosystems.com

PERFORMANCE

Precision: Intra batch CV%<15%, Inter batch CV%<15%.

Specificity: 96.4% (two samples show false positive (n=56).

PRECAUTIONS

1. This kit is for research use only and is not for use in diagnostic or therapeutic applications.
2. This kit should be used according to the provided instructions.
3. Do not mix reagents from different lots.
4. Bring all reagents and samples to room temperature (20℃-25℃) before use. If crystals have formed in the buffer solution, incubate until the crystals have completely dissolved. Before use, bring the solution back to room temperature.
5. This kit should be stored at 2°C -8°C.
6. Please prepare the working solution of each component according to the needs of the experiment. Except for 10x Washing Buffer, all prepared working solution is for one-time use and cannot be stored.

US and Canada: **Tel**: +1 800-810-0816
Asia and Pacific: **Tel**: +86 400-682-2521
Web: http://www.acrobiosystems.com
E-mail: order@acrobiosystems.com