

Mouse Anti-SARS-CoV-2 (B.1.351) Antibody IgG Titer Serologic Assay Kit (Spike S1)

Pack Size: 96 tests

Catalog Number: RAS-T084

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 product is developed for titer measurement of Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike S1) in mouse serum. 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 mouse 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 S1 (B.1.351) on the microplate. Then add the samples, incubate and wash the wells. Next add Secondary antibody HRP-Anti-Mouse 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

		Size (96 tests)	Format	Storage	
Catalog	Components			Unopened	Opened
RAS084-C01	Pre-coated SARS-CoV-2 Spike S1 (B.1.351) Microplate	1 plate	Solid	2-8°C	2-8°C
RAS084-C02	Positive Control	100 μL	Liquid	2-8°C	2-8°C
RAS084-C03	Negative Control	100 μL	Liquid	2-8°C	2-8°C
RAS084-C04	HRP-Goat anti-Mouse IgG	10 μg	Powder	2-8°C, avoid light	-70°C, avoid light
RAS084-C05	10×Washing Buffer	50 mL	Liquid	2-8°C	2-8°C
RAS084-C06	Dilution Buffer	50 mL	Liquid	2-8°C	2-8°C
RAS084-C07	Substrate Solution	12 mL	Liquid	2-8°C, avoid light	2-8°C, avoid light
RAS084-C08	Stop Solution	7 mL	Liquid	2-8°C	2-8°C

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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.

MATERIALS REQUIRED BUT NOT PROVIDED

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

Centrifuge;

37 °C Incubator;

Single channel or multichannel pipettes with 10 μL, 200 μL and 1000 μL precision;

 $10~\mu L,\, 200~\mu L$ and $1000~\mu L$ pipette tips;

Test Tubes;

Graduated cylinder;

Deionized or distilled water for dilution;

REAGENT PREPARATION

- 1. Bring all reagents and samples to room temperature (20°C-25°C) before use.
- 2. As recommended in Table 2, the lyophilized materials of HRP-Goat anti-Mouse IgG will be diluted into a rehydrated solution with ultrapure water/deionized water. Before use, the rehydrated solution needs to be balanced at room temperature of 30 min, shake gently every 10 min. Do not shake or vortex violently. The rehydrated solution should be stored at -70°C, Do not thaw and freeze more than 3 times.

TABLE 2. RECONSTITUTION METHODS FOR 96 TESTS

Catalog	Components	Size	Stock Solution Con.	Reconstitution Buffer and Vol.
RAS084-C04 HRP-Goat anti-Mouse IgG		10 μg	50 μg/mL	200 μL water

RECOMMENDED SAMPLE PREPARATION

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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 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 1:100 with Dilution Buffer.

b. For determination of antibody titer:

It is recommended to dilute the samples, Positive Control and Negative Control from 1:100-1:51200 with Dilution Buffer.

2. Plate set up

Number the diluted samples corresponding to the wells of the Pre-coated SARS-CoV-2 Spike S1 (B.1.351) Microplate. Each experiment requires a set of Positive Control and Negative Control working fluid.

3. Add Samples

Add 100 µL diluted samples, Positive Control and Negative Control working fluid to the corresponding wells. Add 100 µL Dilution Buffer to blank control. Seal the plate with microplate sealing film and incubate at 37°C 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-Goat anti-Mouse IgG

Dilute HRP-Goat anti-Mouse IgG stock solution (50 μg/mL) to 0.08 μg/mL with Dilution Buffer to make a working solution. The prepared working fluid should be stored away from light.

For all wells, add 100 µL HRP-Goat anti-Mouse IgG working solution. Seal the plate with microplate sealing film and incubate at 37°C for 1.0 h, avoid light.

6. Washing

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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°C for 20 min, avoid light.

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, subtract the value read at $OD_{450 \text{ nm}}$ with the value read at $OD_{630 \text{ nm}}$.

CUT-OFF VALUE IDENTIFICATION

Cut-off value =0.1

Normal range of Negative control (1:100): $OD_{450 \text{ nm}}$ - $OD_{630 \text{ nm}}$ < 0.1

Normal range of Positive control (1:1600): $OD_{450 \text{ nm}}$ - $OD_{630 \text{ nm}} \ge 1.0$

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

INTERPRETION OF RESULTS

a. For qualitative detection of antibodies:

Positive reading: $OD_{450 \text{ nm}}$ - $OD_{630 \text{ nm}}$ of sample \geq Cut-off value means Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike S1) are detected.

Negative reading: OD450 nm-OD630 nm of sample < Cut-off value means Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike S1) are not detected.

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

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The kit cannot be used for quantitative detection.

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°C-25°C) 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 1xWashing Buffer, all prepared working solution is for one-time use and cannot be stored.

TYPICAL DATA

Note: The Typical data is for reference only.

a. For qualitative detection of antibodies:

	Value Result in units	Result	Test Result Interpretation	
	$OD_{450 \text{ nm}} - OD_{630 \text{ nm}} = 0.048$	Negative	Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike S1) are not detected	
$OD_{450 \text{ nm}}$ - $OD_{630 \text{ nm}} = 0.462$ Positive Anti-SARS-CoV-2(B.1.351) Anti-SARS-CoV-2(B.1.351)		Anti-SARS-CoV-2(B.1.351) Antibody IgG (Spike S1) are detected		

b. For determination of antibody titer:

Note: Quality control data between different plates should not be mixed, and negative and positive controls should be set for each test.

Ratio of Dilution	OD _{450 nm} - OD _{630 nm} (<u>Samples</u>)	Result
100	3.249	
200	3.200	
400	3.135	
800	2.916	
1600	2.501	
3200	1.670	
6400	0.878	The titer level of antibody is 51200
12800	0.548	
25600	0.322	
51200	0.162	
102400	0.078	
Blank	0.050	

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