

# Mouse Anti-SARS-CoV-2 (XBB.1.5) Antibody IgG Titer Serologic Assay Kit (Spike RBD)

Pack Size: 96 tests

Catalog Number: RAS-T142

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

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





# **INTENDED USE**

The kit is developed for titer measurement of Anti-SARS-CoV-2(XBB.1.5) Antibody IgG (Spike RBD) 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 RBD (XBB.1.5) on the microplate. Then add the samples, incubate and wash the wells. Next add Secondary antibody HRP-Conjugated Antibody 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

C	Components	Size (96 tests)	Format	Storage	
Catalog				Unopened	Opened
RAS142-C01	Pre-coated SARS-CoV-2 Spike RBD (XBB.1.5) Microplate	1 plate	Solid	2-8°C	2-8°C
RAS142-C02	SARS-CoV-2 Antibody Positive Control	100 μL	Liquid	2-8°C	2-8°C
RAS142-C03	42-C03 SARS-CoV-2 Antibody Negative Control		Liquid	2-8°C	2-8°C
RAS142-C04	HRP-Conjugated Antibody	50 μL	Liquid	2-8°C, avoid light	2-8°C, avoid light
RAS142-C05	10×Washing Buffer	50 mL	Liquid	2-8°C	2-8°C
RAS142-C06	142-C06 Dilution Buffer		Liquid	2-8°C	2-8°C
RAS142-C07	2-C07 Substrate Solution		Liquid	2-8°C, avoid light	2-8°C, avoid light
RAS142-C08	AS142-C08 Stop Solution		Liquid	2-8°C	2-8°C

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# STORAGE AND VALIDITY INSTRUCTIONS

The unopened kit is stable for 12 months from the date of manufacture if stored at 2°C to 8°C.

The opened kit should be stored per TABLE 1. The shelf life is 30 days from the date of opening.

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**

Bring all reagents and samples to room temperature  $(20^{\circ}\text{C}-25^{\circ}\text{C})$  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 SARS-CoV-2 Antibody Positive Control and SARS-CoV-2 Antibody 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.

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#### b. For determination of antibody titer:

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It is recommended to dilute the samples, SARS-CoV-2 Antibody Positive Control and SARS-CoV-2 Antibody

Negative Control from 1:100-1:102400 with Dilution Buffer.

2. Plate set up

Number the diluted samples corresponding to the wells of the Pre-coated SARS-CoV-2 Spike RBD (XBB.1.5)

Microplate. Each experiment requires a set of SARS-CoV-2 Antibody Positive Control and SARS-CoV-2 Antibody

Negative Control working fluid.

3. Add Samples

Add 100 µL diluted samples, SARS-CoV-2 Antibody Positive Control and SARS-CoV-2 Antibody 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-Conjugated Antibody

Dilute HRP-Conjugated Antibody solution at 1:1000 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-Conjugated Antibody working solution. Seal the plate with microplate sealing film and

incubate at 37°C for 1.0 h, 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°C for 20

min, avoid light.

8. Termination

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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<sub>450 nm</sub>-OD<sub>630 nm</sub>≥1.5

*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(XBB.1.5) Antibody IgG (Spike RBD) are detected.

Negative reading:  $OD_{450 \text{ nm}}$ - $OD_{630 \text{ nm}}$  of sample < Cut-off value means Anti-SARS-CoV-2(XBB.1.5) Antibody IgG (Spike RBD) 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

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.

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- 2. This kit should be used according to the provided instructions.
- 3. Do not mix reagents from different lots.

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- 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 10x Washing 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
$OD_{450 \text{ nm}} - OD_{630 \text{ nm}} = 0.053$	Negative	Anti-SARS-CoV-2(XBB.1.5) Antibody IgG (Spike RBD) are not detected
$OD_{450 \text{ nm}} - OD_{630 \text{ nm}} = 0.517$	Positive	Anti-SARS-CoV-2(XBB.1.5) Antibody IgG (Spike RBD) 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 <sub>450 nm</sub> - OD <sub>630 nm</sub> ( <u><b>Samples</b></u> )	Result	
100	3.163		
200	3.197		
400	3.037		
800	2.997		
1600	2.600		
3200	1.790		
6400	1.011	The titer level of antibody is 51200	
12800	0.592		
25600	0.292		
51200	0.149		
102400	0.071		
204800	0.036		
Blank	0.016		

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