

Foot and Mouth Disease Virus Type Asia I Antibodies ELISA Kit

Catalog No: E-AD-E020

96T/96T*2/96T*5

Version Number:	V1.2
Replace version:	V1.1
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This manual must be read attentively and completely before using this product.

If you have any problems, please contact our Technical Service Center for help.

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Please kindly provide us the lot number (on the outside of the box) of the kit for more efficient service.

Test principle

This kit is comprised by HRP conjugate, other auxiliary reagents and ELISA Microtiter plate pre-coated with recombinant Foot and Mouth Disease Virus Type Asia1 (FMD-Asia1) protein. Apply the principle of enzyme-linked immunoassay (ELISA) to detect porcine FMD-Asia1 antibody of swine serum samples. During the experiment, add control serum and samples into the ELISA Microtiter plate. If FMD-Asia1 antibodies exist in the samples, it will be bound with the recombinant protein on the microtiter plate after incubation. Then wash the plate to remove unbound antibodies and other components, add the HRP conjugate to specifically bind with the compound of antibody and antigen on the microtiter plate. The unbound HRP conjugate will be removed by washing. Substrate Reagent is added into the well, it will react with the enzyme and the products become blue. The color shade is of positive correlation with antibody levels in the samples. At last, end the reaction by adding stop solution to produce a yellow product. Measure the absorbance value of each well by using a microtiter plate Reader with 450 nm wavelength, then we can know whether there are porcine FMD-Asia1 antibody in the samples.

Kit components

Item	Specifications
ELISA Microtiter plate	96 wells
Dilution plate	96 wells
HRP Conjugate	12 mL
10×Concentrated Wash Buffer	50 mL
Sample Diluent	50 mL
Substrate Reagent A	6 mL
Substrate Reagent B	6 mL
Stop Solution	6 mL
Positive Control	0.8 mL
Negative Control	0.8 mL
Plate Sealer	3 pieces
Sealed Bag	1 pieces
Manual	1pieces

Note: All reagent bottle caps must be tightened to prevent evaporation and microbial pollution.

Other materials required but not supplied

Microplate Reader with 450nm wavelength filter or dual-wavelength (450/630nm)

High-precision transferpettor, EP tubes and disposable pipette tips

37°C incubator or water bath

Deionized or distilled water

Absorbent paper

Notes

1. Wear gloves and work clothes during experiment, and the disinfection and isolation system should be strictly performed. All the waste should be handled as contaminant.
2. The stop solution is corrosive, it should be avoided to contact with skin and clothing. Wash immediately with plenty of water if contacted carelessly.
3. FOR RESEARCH USE ONLY. ELISA Microtiter plate should be covered by plate sealer. Avoid the kit to strong light.
4. Concentrated wash buffer at low temperature condition is easy to crystallize, it should be adjusted to room temperature in order to dissolve completely before use.
5. Each well must be filled with liquid when washing in order to prevent residual free enzyme.
6. The tested sample should keep fresh.
7. The results shall depend on the readings of the Microplate Reader.
8. **Each reagent is optimized for use in the E-AD-E020. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-AD-E020 with different lot numbers.**
9. If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.

Storage and Expiry date

Store at 2-8°C. Avoid freeze.

Please store the opened plate at 2-8°C, the shelf life of the opened kit is up to 1 month.

Expiry date: expiration date is on the packing box.

Sample pretreatment

1. **Serum:** Use the conventional method to prepare serum, the serum must be clear, no hemolysis and no pollution. Samples can be conserved at 2-8°C in 1 week, and it should be stored at -20°C for a long term storage.
2. **Diluted serum:** Dilute the sample serum with the **Sample Diluent** at 1:50 (4 µL of sample serum and 200 µL of sample diluent, mix fully). The positive/negative control do not need to be diluted.
3. **Wash Buffer:** The **10×Concentrated Wash Buffer** should be adjusted to room temperature to make the sediment dissolved fully before use, then dilute it with deionized water at 1:9.

Assay procedure

Restore all reagents and samples to room temperature before use. All the reagents should be mixed thoroughly by gently swirling before pipetting. Avoid foaming. The unused ELISA Microtiter plate should be sealed as soon as possible and stored at 2-8°C.

1. **Number:** number the sample and control in order (multiple well), and keep a record of control wells and sample wells. Set 2 wells for negative control, 1 well for positive control and 1 well for blank control respectively. **Samples need test in duplicate.**
2. **Add sample:** add 100 µL of **positive/negative control** to positive/negative control well, and add 100 µL of **Sample Diluent** to blank control. Add 100 µL of **Diluted serum** to the sample wells, mix fully.
3. **Incubate:** cover the plate sealer and mix thoroughly, incubate at 37°C for 30 min in shading light.
4. **Wash:** remove the liquid in each well. Immediately add 300 µL of **Wash Buffer** to each well and wash. Repeat wash procedure for 3 times, 60 s intervals/time. Invert the plate and pat it against thick clean absorbent paper (If bubbles exist in the wells, clean tips can be used to prick them).
5. **HRP conjugate:** add 100 µL of **HRP Conjugate** into each well, and incubate at 37°C for 30 min.
6. **Wash:** Repeat step 4 for washing.
7. **Color Development:** add 50 µL of **Substrate Reagent A** and 50 µL of **Substrate Reagent B** into each well and mix thoroughly. Incubate for 10 min at 37°C in shading light.
8. **Stop reaction:** add 50 µL of **Stop Solution** into each well, mix thoroughly.
9. **OD Measurement:** Measure the OD-value of each well by using a microplate reader with 450 nm wavelength (use 630 nm as reference wavelength). This step should be finished in 10 min after stop reaction.

Reference value

Normally, the A-value of negative control < 0.1 and positive control ≥ 0.6 .

Interpretation of the results

1. Cut Off (C.O) = $0.20 + \text{Average of negative control ODs}$ (when average A450 of NC < 0.05 , calculate at 0.05)
2. Positive result: Sample OD $>$ Cut Off (C.O)
3. Negative result: Sample OD $<$ Cut Off (C.O)

Limitations

1. This kit can only be used to qualitatively detect the FMD-Asia1 antibody in serum of swine.
2. The detection results of this kit are only for reference. For confirmation of the result, please combine the symptoms and other methods of detection, this detection cannot be used as the only criteria for result.