

African Swine Fever Antibodies ELISA Kit

Catalog No: E-AD-E143

96T/96T*3/96T*5

Version Number:	V1.4
Replace version:	V1.3
Revision Date:	2026.02.02

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.

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017

Email: techsupport@elabscience.com

Website: www.vetassay-elab.com

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 reagents and ELISA Microtiter plate pre-coated with recombinant African swine fever (ASFV) p30 and VP72 antigen. Apply the principle of enzyme-linked immunoassay (ELISA) to detect ASFV-Ab in serum or plasma of porcine. During the experiment, add control and samples into the ELISA Microtiter plate, ASFV-Ab will be bound with the antigen on the ELISA Microtiter plate. Then wash the plate to remove unbound components, horseradish peroxidase (HRP) conjugate is added to each ELISA Microtiter plate well. The unbound HRP Conjugate will be removed by washing and substrate reagent is added for color development. At last, end the reaction by adding Stop Solution to produce a yellow product. Measure the absorbance value of each well by using a microplate reader with 450 nm (630 nm) wavelength, then we can judge whether ASFV antibody exist in the sample.

Kit components

Item	Specification
ELISA Microtiter plate	96 wells
Positive Control	2 mL
Negative Control	2 mL
100×Concentrated HRP Conjugate	0.24 mL
HRP Conjugate Diluent	24 mL
Sample Diluent	12 mL
Substrate Reagent	12 mL
Stop Solution	15 mL
25×Concentrated Wash Buffer	50 mL
Plate Sealer	1 piece
Sealed Bag	1 piece
Manual	1 copy

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

Other materials required but not supplied

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

High-precision transferpettor (10µl-100µl, 100µl-1000µl), 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. Hemolysis, contaminated samples may lead to abnormal results.
7. The results shall depend on the readings of the microplate reader.
8. **Each reagent is optimized for use in the E-AD-E143. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-AD-E143 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.

Experimental preparation

Restore all reagents and samples to room temperature before use.

Open the microplate reader in advance, preheat the instrument, and set the testing parameters.

1. Sample pretreatment notice:

Experimental apparatus should be clean, and the pipette should be disposable to avoid cross-contamination during the experiment.

2. Solution preparation

Please prepare solution according to the number of samples. Don't use up all components in the kit at once!

Solution 1: Wash Buffer

Dilute **25×Concentrated Wash Buffer** with deionized water. (**25×Concentrated Wash Buffer** (V): Deionized water (V) = 1:24).

Solution 2: HRP Solution

Dilute **100×Concentrated HRP Conjugate** with **HRP Conjugate Diluent**. (**100×Concentrated HRP Conjugate** (V): **HRP Conjugate Diluent** (V) = 1:99).

Sample preparation

1. **Serum/plasma:** Use the conventional method to prepare serum/plasma. 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.

Assay procedure

Restore all reagents and samples to room temperature (25°C) 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 positive/negative control respectively. **Samples need test in duplicate.**
2. **Add sample diluent:** add 50 µL of **Sample Diluent** to each sample well.
3. **Add sample:** add 50 µL of **Positive/Negative control** to positive/negative control well, then add 50 µL **Serum/plasma** to the sample wells.
4. **Incubate:** cover the plate sealer and mix thoroughly, incubate at 37°C for 60 min in shading light.
5. **Wash:** remove the liquid in each well. Immediately add 300 µL of **Wash Buffer** (Solution 1) to each well and wash. Repeat wash procedure for 4 times, 30 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).
6. **HRP conjugate:** add 100 µL of **HRP Solution** (Solution 2) into each well, cover the plate sealer and incubate at 37°C for 30 min in shading light.
7. **Wash:** Repeat step 5 for washing.
8. **Color development:** Add 100 µL of **Substrate Reagent** into each well and mix thoroughly. Cover the plate sealer and mix thoroughly, incubate at 25°C for 10 min in shading light.
9. **Stop reaction:** add 50 µL of **Stop Solution** into each well, mix thoroughly.
10. **OD measurement:** measure the absorbance value (A-value) of each well by using a Microplate Reader with 450 nm/630 nm wavelength. **Note: Read the results within 5 min.**

Reference value

Normally, the A-value of negative control / A-value of positive control ≥ 3.0 , the test result is valid, otherwise the test should be conduct again.

Interpretation of the results

1. OD_{NC}: Average OD of negative control;
OD_{PC}: Average OD of positive control;
OD_S: Average OD of sample.
2. Inhibition rate of sample (X%)

$$X\% = \frac{OD_{NC} - OD_S}{OD_{NC} - OD_{PC}} \times 100\%$$

3. Positive result: X% $\geq 50\%$.
4. Negative result: X% $< 40\%$.
5. If the test results of the sample fall between the two ranges, making it a suspicious sample.

Limitations of this test method

1. This test is only used as the qualitative detection of ASFV antibodies in serum/plasma of porcine.
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.