Elabscience®

Bluetongue Competition Antibody ELISA Kit

REF E-AD-E116-96T*2

TEST PRINCIPLE

This kit is comprised by HRP conjugate, other auxiliary reagents, ELISA Microtiter plate pre-coated with the bluetongue antigen. Apply the principle of enzyme-linked immunoassay (ELISA) to detect bluetongue antibody in serum and plasma samples of livestock animals (cattle, goats or sheep etc.). During the experiment, add control serum and samples into the ELISA Microtiter plate. If bluetongue antibodies exist in the samples, it will compete with the antibody in the antibody working solution to bind with the antigen pre-coated on the Microplate. Then wash 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 become blue. The color shade is negative correlation with bluetongue 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 Microplate Reader with 450 nm wavelength, then we can judge whether BTV antibody exist in the sample.

KIT COMPONENTS

Item	Specification
ELISA Microtiter plate	96*2 wells
HRP Conjugate	25 mL
20×Concentrated Wash Buffer	60 mL
Antibody Working Solution	15 mL
Substrate Reagent A	12 mL
Substrate Reagent B	12 mL
Stop Solution	12 mL
Positive Control	100 µL
Negative Control	100 µL
Plate Sealer	3 pieces
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, EP tubes and disposable pipette tips
37℃ incubator or water bath
Deionized or distilled water
Absorbent paper
Physiological saline solution

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. The ELISA plate obtained from cold storage conditions should be adjusted to room temperature before opening the bag. The unused plate should be kept in a sealed bag with desiccant.
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-E116. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-AD-E116 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℃. Avoid freeze.

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

Expiry date: expiration date is on the packing box.

REAGENT PREPARATION

1. **Serum/plasma:** Use the conventional method to prepare serum/plasma, the serum / plasma must be clear, no hemolysis and no pollution. Samples can be conserved at 2-8℃ in 1 weeks, and it should be stored at - 20℃ for a long term storage.
2. **Wash Buffer:** The **20×Concentrated Wash Buffer** should be adjusted to room temperature before use, then dilute it with distilled or deionized water at 1:19.

ASSAY PROCEDURE

Restore all reagents and samples to room temperature (25℃) 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℃.

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/positive control respectively. **Samples need test in duplicate**.
2. Add 40 µL **Wash Buffer** to each well.
3. **Add sample:** add 10 µL of **positive/negative control** to positive/negative control well, add 10 µL of sample to each sample well.
4. Add 50 µL **Antibody Working Solution** to each well.
5. **Incubate:** Cover the plate sealer and mix thoroughly, incubate at 37℃ for 60 min in shading light.
6. **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, 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).
7. **HRP conjugate:** add 100 µL of **HRP Conjugate** into each well, cover the plate sealer and incubate at 37℃ for 30 min in shading light.
8. **Wash:** repeat step 6 for washing.
9. **Color Development:** add 50 µL of **Substrate Reagent A** and 50 µL of **Substrate Reagent B** into each well, Cover the plate sealer and mix thoroughly, incubate at 37℃ for 10 min in shading light.
10. **Stop reaction:** add 50 µL of **Stop Solution** into each well, mix thoroughly.
11. **OD Measurement:** Measure the absorbance value (A-value) of each well by using a Microplate Reader with 450 nm wavelength (use 630 nm as reference wavelength).

INTERPRETATION OF THE RESULT

1. The average OD value of negative control is OD_{NC}, average OD value of positive control is OD_{PC}, OD value of sample is OD_S. Percentage of inhibition of positive sample is PI_{PC}, Percentage of inhibition of sample is PI_S.
2. $PI_{PC} = (OD_{NC} - OD_{PC}) / OD_{NC} \times 100\%$
   $PI_S = (OD_{NC} – OD_s) / OD_{NC} \times 100\%$
3. Normally, $OD_{NC} > 0.5$, and $PI_{PC} > 60\%$. Otherwise, it should be retested.
4. Positive result: $PI_S \geq 50\%$, negative result: $PI_S < 50\%$.

LIMITATIONS OF THIS TEST METHOD

1. This test is only used as the qualitative detection of bluetongue antibody in serum and plasma samples. A rough estimate (high, general, low) of the antibody concentration can be calculated according to the PI values.
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.

BASIC INFORMATION

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