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Adalimumab PK Quantitation Kit (ELISA)

Generic Name

Adalimumab PK Quantitation Kit (ELISA) Cat#: 17-0087

【Intended Use】

To detect the contents of Adalimumab antibody in the blood samples.

[Principle]

This product adopts the principle of Double Antibody Sandwich Method (Sandwich Elisa). The flat-bottom 96-well plates are coated with anti-Adalimumab antibody. After adding the samples, the anti-Adalimumab monoclonal antibody labeled with HRP is used for detection. The content of the Adalimumab antibody in the blood samples can be detected by the degree of TMB color development.

[Materials and Reagents]

- 1. Coated plate, 12 wells \times 8 strips
- 2. Enzyme conjugate, 120 µ L×1 tube (diluted 100 times for use)
- 3. BSA, 3G ×1 pack
- 4. Calf serum, 5ml×1 vial
- 5. 20× Washing Buffer, 50mL×1 vial
- 6. 20× PBST Washing Buffer, 50mL×1 vial
- 7. Substrate Solution A, 7mL×1 vial
- 8. Substrate Solution B, $7mL \times 1$ vial
- 9. Termination Solution, $7mL \times 1$ vial
- 10. Sealing plate film, 2 pieces
- 11. Instruction book

[Storage]

- 1. All components remain stable under the condition of $2-8^{\circ}$ C;
- 2. Avoid light. Valid for six months.

(Protocol for Detection **)**

1. Equilibration

Equilibrate the required reagents at room temperature (18~25°C) for 30 minutes.

- 2. Dosing solution: Please configure the reagents before use.
- 2.1 1× washing buffer: Take 1 vial of 20× PBST washing buffer, dilute it to 1000ml with deionized water, mix well.
- 2.2 $1 \times$ dilution buffer: Take 1 vial of $20 \times$ washing buffer, dilute it to 1000ml with deionized water, mix well.
- 2.3 Sample dilution buffer (20% CS): After Calculating the total volume of the sample dilution buffer, dilute the calf serum 5 times using the prepared 1×dilution buffer (step 2.2), mix well.
- 2.4 Enzyme dilution buffer: Dissolve BSA (3g/pack) completely into 100ml of the prepared 1× dilution buffer (step 2.2), mix well.
- 2.5 Enzyme solution: Dilute the enzyme conjugate 100 times with the enzyme dilution buffer prepared in step 2.4, mix well.
- 2.6 Standards: Dilute the Adalimumab by pure monkey negative serum by grads' multiple. the gradients are 5000ng/ml, 2000ng/ml, 1000ng/ml, 500ng/ml, 200ng/ml, 50ng/ml, and 10ng/ml. Dilute each gradient 100 times by the sample dilution buffer (step 2.3) respectively, mix well.
- 3. Adding standards

Remove the coated plate from the sealed bag and dilute the standard to the different concentrations. After adding 100µl of standard or sample to each well (including negative control), seal the plate with sealing film. Place the plate in a shaking incubator (37°C, 200 rpm) and incubate for 60 minutes

4. Washing

Discard the liquid in each well, fill the microwells $(350\mu l/well)$ with $1\times$ washing buffer, and discard the liquid in the wells after 30 seconds. Repeat these steps for 3 times, then pat the plate on the paper towel after the last wash.

5. Adding enzyme solution

After diluting the enzyme conjugate 100 times with the enzyme solution buffer, add the solution to the microplate (100µl per well). Seal the plate with sealing film. Place the plate in a shaking incubator (37°C, 200 rpm) and incubate for 60 minutes.

6. Washing

Repeat step 4.

7. Coloring

Add 50µl each of Substrate Coloring Solution A and B into each well. Mix well with gentle tapping. Then incubate the plate at room temperature for 10 minutes in the dark.

8. Termination

Terminate the reaction by adding 50μ l of 0.2M H₂SO₄ into each well and mix gently. Set the main wavelength of the microplate reader at 450nm and the reference wavelength at 630nm. Measure the absorbance (OD value) of each well.

【Data Analysis】

It is recommended to adopt the fitting method of double logarithm for fitting and calculation.

| Standard Curve Concentration | OD Value | | Mean Value |
|------------------------------|----------|-------|------------|
| 5000 | 2.175 | 2.084 | 2.1295 |
| 2000 | 1.564 | 1.42 | 1.492 |

| 1000 | 0.877 | 0.848 | 0.8625 |
|------|--------|-------|---------|
| 500 | 0. 451 | 0.472 | 0. 4615 |
| 200 | 0.217 | 0.222 | 0.2195 |
| 50 | 0.105 | 0.098 | 0.1015 |
| 10 | 0.074 | 0.069 | 0.0715 |
| NC | 0.069 | 0.064 | 0.0665 |

After double logarithm fitting of the concentrations and OD values, the curve equation is as follows:



[Product Performance Index]

- 1. Linear range:10~5000ng/ml
- 2. Detection limit: <50ng/ml
- 3. Accuracy: CV% ≤15% (n=10)

[Limitations]

- 1. This kit is only used to detect the content of Adalimumab antibody in blood samples.
- 2. Results out of the measurement range of the kit are unreliable.
- 3. Severe hemolysis, chyle, and bilirubin samples may cause abnormal test results.

Caution

- 1. Avoid cross contamination.
- 2. Follow reader measure as a standard.
- 3. All samples and buffers should be added or removed with pipette.
- 4. Do not mix reagents from different batches.