

(FOR RESEARCH USE ONLY. DO NOT USE IT IN DIAGNOSIS!)

Cat No.: EKF1068

Mabuterol (Mab) ELISA Kit

If you have any questions or need further help during experiment, please don't hesitate to contact us through the following methods:

- ✉ Email (Order) order@enkilife.com
- ✉ Email (Techsupport) techsupport@enkilife.com
- 📞 Tel: 0086-27-87002838
- 🌐 Website: www.enkilife.com

Shelf life: Please refer to the label on the outer package.

Techsupport: In order to provide you with better service, please inform us the lot number on the label of the outer package.

Product description

This ELISA kit is rapid, quantitative enzyme-linked immunosorbent assays (ELISA) for the determination of Mabuterol content in Food samples.

Key Features

- Sensitivity: 0.1 ppb (ng/mL)
- Detection range: 0.1 ppb~8.1 ppb
- Specificity:

Mabuterol	100%
Salbutamol, Terbutaline, Cimaterol, Ractopamine	< 0.1%

- Detection limits:

Pig urine, pork, beef, lamb, chicken, duck, pork liver, ham sausage	0.5 ppb
Cow urine, sheep urine	1.0 ppb
Pig feed	5.0 ppb

- Recovery rate: 90%±30%

Storage and Shelf Life

The kit should be stored at 2~8°C and must not be frozen. The shelf life of this product is 1 year. Do not use the kit beyond the expiration date.

Test Principle

This assay employs the competitive inhibition enzyme immunoassay technique. The kit consists of an enzyme-labeled plate pre-coated with coupled antigens, horseradish enzyme markers, antibodies, standard substances, and other supporting reagents. During detection, standard substances or sample solutions are added. Mabuterol in the samples competes with the pre-coated coupled antigens on the enzyme-labeled plate for Mabuterol specific antibody. After adding the enzyme markers, TMB substrate is used for color development. The absorbance value of the sample is negatively correlated with the content of Mabuterol it contains, and the residual amount of Mabuterol in the sample can be obtained by comparing with the standard curve.

Materials Supplied

Item	Quantity
Assay Plate	96T
Standard	6 x 1.5 mL
High concentration standard (100 ppb)	1 x 1 mL
HRP Conjugate dilution	1 x 9 mL
HRP Conjugate concentrate (11×)	1 x 1 mL
TMB Substrate Reagent (A\B)	2 x 7 mL
Stop Solution	1 x 7 mL
Concentrated Sample dilution (50×)	1 x 30 mL
Wash Buffer (20×)	1 x 25 mL
Plate Sealer	1 piece
Instruction Manual	1 piece
Sealed Bag	1 copy

Standard Concentration

Standard	1	2	3	4	5	6
Concentration (ppb=ng/mL=ng/g)	0	0.1	0.3	0.9	2.7	8.1

Materials Required, Not Supplied

- Instruments: Microplate reader, printer, homogenizer, nitrogen blow-drying device, vortex, centrifuge, constant temperature incubator, balance (sensitivity 0.01g)
- Single-channel micropipette (20 μ L-200 μ L, 100 μ L-1000 μ L)
- 300 μ L multichannel micropipette
- Reagents: KH₂PO₄, HCl, NaOH, Twain 20
- Distilled or deionized water

① Notes:

- Before using the kit, read the instructions carefully.
- Do not use expired kits, and do not mix reagents in kits with different batch numbers.
- Kindly use graduated containers to prepare the reagent.
- Bring all reagents to room temperature (20-25°C) before use for 30 min.
- Only the disposable tips can be used for the experiments and the tips must be changed when used for different reagents.
- Distilled water is recommended to be used to make the preparation for reagents. Contaminated water or container for reagent preparation will influence the detection

result.

- The stop solution is acidic. Wear eyes, hands, face, and clothing protection when using the product.
- EnkiLife is only responsible for the kit itself, but not for the samples consumed during the assay. The user should calculate the possible amount of the samples used in the whole test. Please reserve sufficient samples in advance.
- If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.
- Please predict the concentration before assaying. If values for these are not within the range of the standard curve, users must determine the optimal sample dilutions for their particular experiments.

Reagent Preparation

- **Wash Buffer (1x):** If crystals have formed in the concentrate, warm up to room temperature and mix gently until the crystals have completely dissolved. Dilute 10 mL of Wash Buffer (20x) into 190 mL deionized or distilled water to prepare 200 mL of Wash Buffer (1x). Keep it at 4 °C for one month.
- **Feed Extraction Solution:** Weigh 1.36 g of **KH₂PO₄** and add 1000 mL of deionized water, mix fully.
- **0.1 M HCl:** Dilute 8.6 mL of **HCl** to 1000 ml with deionized water.
- **0.5 M NaOH:** Dissolve 2 g of **NaOH** to 100 mL with deionized water, mix fully.
- **Sample diluent:** Dilute 1 part of the **Concentrated Sample dilution (50×)** and 49 parts of deionized water, mix well.
- **10% Twain 20:** Weigh 1 g of **Tween 20** and add 9 mL of deionized water, then mix well.
- **Cooked food extraction solution:** Take 400 ml of **0.1 M HCl** solution and add 6 ml of **10% Tween 20**, then mix well.
- **HRP Conjugate working solution:** Dilute 1 part of the **HRP Conjugate concentrate (11×)** and 10 parts of the HRP Conjugate diluent, mix well. Prepare and use immediately.

Sample Preparation and collection

The prepared sample may be stored for up to one day at 4°C.

Pig urine: Centrifuge the 200 μ L of sample at 4000 rpm for 5 min. Take 80 μ L of supernatant for analysis. Dilution factor of sample: 1.

Cow urine, sheep urine: Take 600 μ L of urine sample, centrifuge at 4000 rpm for 5 min. Take 500 μ L of supernatant, add 500 μ L of **Wash Buffer** and vortex for 1 min. Take 80 μ L for analysis. Dilution factor of sample: 2.

Muscle (pork, beef, lamb, chicken, duck): Remove fat from fresh sample, homogenize the sample with homogenizer. Weigh 2 ± 0.05 g of homogenate sample into the 50 mL centrifuge tube. Add 6 mL of **Sample diluent**, vortex for 2 min, centrifuge at 4000 rpm for 5 min. Take 80 μ L of supernatant for analysis. Dilution factor of sample: 4.

Pork liver: Weigh 2 ± 0.05 g of homogenate sample into the 50 mL centrifuge tube. Add 6 mL of **0.1 M HCl**, vortex for 2 min, and centrifuge at 4000 rpm for 5 min. Remove 1 mL of the supernatant to another centrifuge tube, add 100 μ L of **0.5 M NaOH** to adjust the pH to 6-8. Vortex for 10s and centrifuge at 4000 rpm for 5 min. Take 80 μ L of supernatant for analysis. Dilution factor of sample: 4.

Pig feed: Weigh 0.5 ± 0.05 g of homogenate sample into the 50 mL centrifuge tube. Add 10 mL of **Extraction Solution**, vortex for 2 min, centrifuge at 4000 rpm for 10 min. Take 80 μ L of supernatant for analysis. Dilution factor of sample: 20.

Ham sausage: Weigh 2 ± 0.05 g of homogenate sample into the 50 mL centrifuge tube. Add 4 mL of **cooked food extraction solution**, vortex for 2 min, centrifuge at 4000 rpm for 5 min. Take 80 μ L of intermediate clear liquid for analysis. Dilution factor of sample: 20.

Assay Protocol

ⓘ Notes:

- Take the required reagent out of the refrigerated environment at 4°C and place it at room temperature for more than 30 min. If the reagent is crystalline, allow it to dissolve sufficiently at room temperature. Shake each liquid reagent well before use.
- It is recommended that all standards and samples be run at least in duplicate.
- Remove the required number of assay plates, put the unused assay plates in sealed bags, and store at 4°C.
- Avoid using metal packaging and stirring reagents.
- Mix the liquid well and complete removal of liquid at each step is essential to good performance.
- Discard the substrate with any color that indicates the degeneration of this solution. When the absorbance value of 0ppb standard less than 0.5 indicates its degeneration.
- The sample addition time for each step shall not exceed 3 min.

1. **Number:** The corresponding microwells of the sample and the standard are numbered in sequence, and each sample and standard are parallel to each well with 2 wells, and the location of the standard well and the sample well is recorded.

2. **Start the reaction:** Add 80 μ L of **Standard or Sample** per well to the respective

microwells, then add 70 μ L of **HRP Conjugate Working Solution** per well. Cover the assay plate with a plate sealer, gently shake for 5s to mix well, and react in the dark at 25 °C for 20 min.

3. Washing: Carefully remove the plate sealer, discard the liquid in the plate wells. Wash by filling each well with 350 μ L of **Wash Buffer (1X)** using a squirt bottle, multi-channel pipette, manifold dispenser, or auto washer, and let it stand for 30s. Aspirate or decant the liquid in the plate wells, pat it dry against clean absorbent paper, and complete one wash. Repeating the process 5 times.

Tips: After the plate wells is dried, the next step should be carried out immediately.

4. Add Substrate Reagent: Add 50 μ L of **Substrate Reagent A** to each well, then add 50 μ L of **Substrate Reagent B**, gently shake for 5s to mix well. React in the dark for 15 min at 25°C.

Tips: Adjust the incubation time according to the color change, but do not exceed 30 min. Once the standard wells show a clear gradient, the incubation can be stopped.

5. Stop the Reaction: Add 50 μ L of **Stop Solution** to each well, gently tap the plate to ensure thorough mixing.

6. Measure the OD value: Determine optical density (OD) result at 450 nm within 10 min. (Recommend reading the OD value at the dual-wavelength: 450/630 nm).

⌚ Calculation of results

1. Calculation of percent absorbance:

The percent absorbance of the standard or sample is equal to the average of the percent absorbance value of the standard or sample (double well) divided by the absorbance value of the first standard (0 ppb), multiplied by 100%.

$$\text{percent absorbance (\%)} = \frac{A}{A_0} \times 100\%$$

A—The average absorbance value of the standard or sample solution

A_0 —Average absorbance value of 0 ppb standard

2. Create of standard curve:

Take the percentage absorbance of the standard substance as the Y-axis and the logarithm of the corresponding standard substance concentration (ppb) as the X-axis to plot the semi-logarithmic curve of the standard substance. The percentage absorbance of the sample is substituted into the standard curve, and the concentration corresponding to the

sample is read out from the standard curve, and multiplied by its corresponding dilution factor to get the actual concentration of the analyte in the sample. Substitute the percentage absorbance of the sample into the standard curve, read the concentration corresponding to the sample from the standard curve, and multiply it by the corresponding dilution factor to obtain the actual concentration of the analyte in the sample.

3. Standard curve reference

Standard	1	2	3	4	5	6
Concentration (ppb=ng/mL=ng/g)	0	0.1	0.3	0.9	2.7	8.1
OD Standard	2.292	1.667	1.134	0.588	0.287	0.110

 We are always committed to providing high-quality products and thank you for your understanding and support. If you have any questions, please feel free to contact our technical support team.