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- Trockeneiszuschlag
- Gefahrgutzuschlag
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PHE A (Phenylethanolamine A) ELISA Kit

Catalog No: E-FS-E015

96T/96T*3

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.elabscience.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 uses Competitive-ELISA as the method for the quantitative detection. It can detect Phenylethanolamine A (PHE A) in samples, such as muscle, feed. This kit is composed of ELISA Microtiter plate, HRP conjugate, antibody working solution, standard and other supplementary reagents. The microtiter plate in this kit has been pre-coated with coupled antigen. During the reaction, PHE A in the samples or standard competes with coupled antigen on the solid phase supporter for sites of anti-PHE A antibody. Then Horseradish Peroxidase (HRP) conjugate is added to each microtiter plate well, and substrate reagent is added for color development. There is a negative correlation between the OD value of samples and the concentration of PHE A. The concentration of PHE A in the samples can be calculated by comparing the OD of the samples to the standard curve.

Technical indicator

Reaction mode(Incubation time and temperature) **:** 25℃; 30 min,15 min

Detection limit: Urine ---0.1 ppb; Muscle ---0.1 ppb; Feed ---1 ppb

Cross-reactivity: PHE A ---100%, Clenbuterol ---＜1%, Salbutamol ---＜1%, Ractopamine ---＜1%

Sample recovery rate: Urine ---95%±15%, Muscle, Feed ---85%±15%,

Kits components

Item	Specifications
ELISA Microtiter plate	96 wells
Standard Liquid	1 mL each (ppb=ng/mL=ng/g) (0ppb, 0.1ppb, 0.3ppb, 0.9ppb, 2.7ppb, 8.1ppb)
HRP Conjugate	5.5 mL
Antibody Working Solution	5.5 mL
Substrate Reagent A	6 mL
Substrate Reagent B	6 mL
Stop Solution	6 mL
20×Concentrated Wash Buffer	40 mL
10×Reconstitution Buffer	50 mL
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

Instruments: Microplate reader, Printer, Homogenizer, Nitrogen evaporators, Water bath, Vortex mixer, Centrifuge, Graduated pipette, Balance (sensibility 0.01 g).

Micropipette: Single channel (20-200 μL, 100-1000 μL), Multichannel (30-300 μL).

Reagents: NaOH, Ethyl acetate, Concentrated HCl, Acetonitrile, Methanol, N-hexane, Na_2SO_4.

Experimental preparation

Restore all reagents and samples to room temperature before use.

Open the micro-plate 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: 0.1 M HCl Solution

   Dilute 0.86 mL of **Concentrated HCl** to 100mL with deionized water.

   Solution 2: 0.1 M NaOH Solution *(for muscle sample)*

   Dissolve 0.4 g of **NaOH** to 100mL with deionized water.

   Solution 3: Acetonitrile- **0.1 M HCl Solution** (Solution 1)

   Acetonitrile (V): 0.1 M HCl (V) =84:16 *(for tissue sample)*

   Solution 4: Reconstitution Buffer *(for muscle, feed sample)*

   Dilute the **10×Reconstitution Buffer** with deionized water. (10×Reconstitution Buffer (V): Deionized water (V)=1:9) .The Reconstitution buffer can be store at 4℃ for a month.

   Solution 5: Wash Buffer

   Dilute **20×Concentrated Wash Buffer** with deionized water. (20×Concentrated Wash Buffer (V): Deionized water (V) = 1:19).

3. Sample pretreatment procedure

3.1 Pretreatment of urine sample:

Take 50 μL of clear urine sample to detect (if the urine sample is turbid, it should be filtered or centrifuged at 4000 r/min for 5 min until the urine sample become clear). Temporarily used samples should be kept frozen to save.

Note: Sample dilution factor: 1, detection limit: 0.1ppb

3.2 Pretreatment of muscle sample:

(1) Remove fat from sample, homogenize the sample with homogenizer.

(2) Weigh 2±0.05 g of homogenate muscle sample, add 6mL of **Acetonitrile- 0.1 M HCl Solution (**Solution 3), vortex for 2 min, and centrifuge at 4000 r/min at room temperature for 10 min.

(3) Take 3 mL of the supernatant, add 2 mL of **0.1 M NaOH Solution (**Solution 2), add 6 mL of **Ethyl acetate**, vortex for 2 min, and centrifuge at 4000 r/min at room temperature for 10 min. Take all the supernatant to dry at 50-60℃with nitrogen evaporators or water bath.

(4) Add 1 mL of **Reconstitution Buffer (**Solution 4), mix and vortex for 30s, take 50 uL liquid for analysis

Note: Sample dilution factor: 1, detection limit: 0.1ppb

3.3 Pretreatment of feed sample:

(1) Homogenize the representative sample with a homogenizer and mix fully.

(2) Weigh 1.0±0.05 g of homogenate feed sample, add 10 mL of **Methanol**, then add 5 g of **Na_2SO_4**, vortex for 2 min, and centrifuge at 4000 r/min at room temperature for 10min.

(3) Take 1 mL of supernatant, dry at 50-60℃ with nitrogen evaporators or water bath. Dissolve the dried residual with 1 mL of **Reconstitution Buffer (**Solution 4), add 1 mL of **N-hexane** and mix for 30s. Centrifuge at 4000 r/min at room temperature for 5 min.

(4) Take 50 μL of lower liquid for analysis.

Note: Sample dilution factor: 10, detection limit: 1ppb

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 standard in order (multiple well), and keep a record of standard wells and sample wells. **Standard and Samples need test in duplicate.**
2. **Add Sample:** add 50 μL of **Standard or Sample** per well, then add 50 μL of **HRP Conjugate** to each well, then add 50 μL of **Antibody Working Solution**, cover the plate with plate sealer. Oscillate for 5s gently to mix thoroughly, incubate at 25℃ for 30 min in shading light.
3. **Wash:** uncover the sealer carefully, remove the liquid in each well. Immediately add 300 μL of **Wash Buffer** (Solution 5) to each well and wash. Repeat wash procedure for 5 times, 30s 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).
4. **Color Development:** add 50 μL of **Substrate Reagent A** to each well, and then add 50 μL of **Substrate Reagent B**. Gently oscillate for 5s to mix thoroughly. Incubate shading light for 15 min at 25℃.
5. **Stop Reaction:** add 50 μL of **Stop Solution** to each well, oscillate gently to mix thoroughly.
6. **OD Measurement:** determine the optical density (OD value) of each well at 450 nm (reference wavelength 630 nm) with a microplate reader. This step should be finished in 10 min after stop reaction.

Result analysis

1. **Absorbance (%) =A/A_0 ×100%**

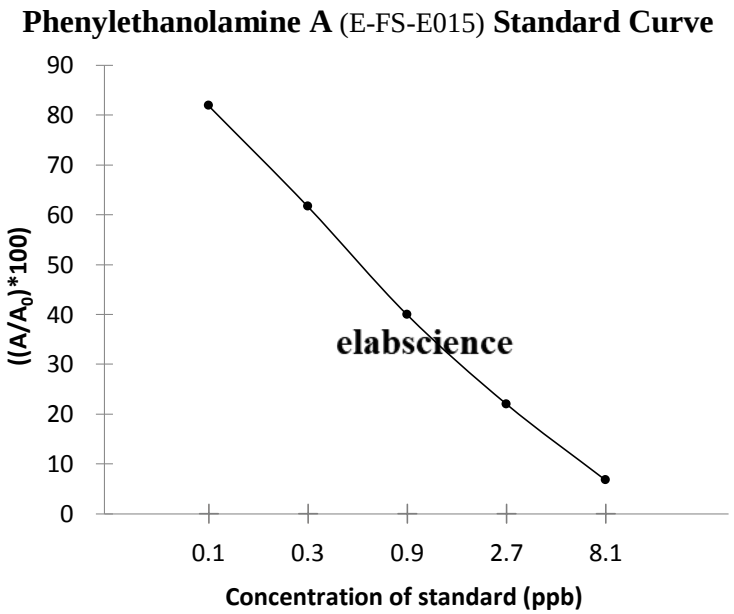
   A: Average absorbance of standard or sample

   A_0: Average absorbance of 0 ppb Standard

2. **Drawing and calculation of standard curve**

   Create a standard curve by plotting the absorbance percentage of each standard on the y-axis against the log concentration on the x-axis to draw a semi-logarithmic plot. Add average absorbance value of sample to standard curve to get corresponding concentration. **If samples have been diluted, the concentration calculated from the standard curve must be multiplied by the dilution factor.**

   For this kit, it is more convenient to use professional analysis form for accurate and fast analysis on a large number of samples.



Notes

1. The overall OD value will be lower when reagents have not been brought to room temperature before use or room temperature is below 25℃.
2. If the wells turn dry during the washing procedure, it will lead to bad linear standard curve and poor repeatability. Operate the next step immediately after wash.
3. Mix thoroughly and wash the plate completely. The consistency of wash procedure can strongly affect the reproducibility of this ELISA kit.
4. ELISA Microplate should be covered by plate sealer. Avoid the kit to strong light.
5. **Each reagent is optimized for use in the E-FS-E015. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-FS-E015 with different lot numbers.**
6. Substrate Reagent should be abandoned if it turns blue color. When OD value of standard (concentration: 0)＜0.5 unit (A450nm＜0.5), it indicates the reagent may be deteriorated.
7. Stop solution is caustic, avoid contact with skin and eyes.
8. As the OD values of the standard curve may vary according to the conditions of the actual assay performance (e.g. operator, pipetting technique, washing technique or temperature effects), the operator should establish a standard curve for each test.
9. Even the same operator might get different results in two separate experiments. In order to get reproducible results, the operation of every step in the assay should be controlled.
10. **For mentioned sample fast and efficient extraction methods are included in the kit description. Please consult technical support for the applicability if other sample need to be tested.**
11. The kit is used for rapid screening of actual samples. If the test result is positive, the instrument method such as HPLC, LC/MS, etc. can be used for quantitative confirmation.

Storage and expiry date

Store the kit at 2-8℃. Do not freeze any test kit components.

Return any unused microwells to their original foil bag and reseal them together with the desiccant provided and further store at 2-8℃.

Expiry date: expiration date is on the packing box.