

7th Edition, revised inApril, 2017

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

PHEA(Phenethylamine A)Rapid Test Kit

Catalog No: E-FS-C011 50T

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.

Phone: 240-252-7368(USA)240-252-7376(USA) Email: <u>techsupport@elabscience.com</u> Website: <u>www.elabscience.com</u>

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 the principle of competitive-inhibition-GICA. It can detect PHEA (Phenethylamine A) in urine sample.After adding the sample solution into the sample well of detect card, PHEA of the sample solution combine with the gold-labelled antibody, so as to prevent the combining of gold-labelled antibody with PHEA conjugate on the cellulose membrane. When the concentration of PHEA in the sample solution is more than the detection limit, the detect line do not show color reaction and the result is positive. When the concentration of PHEA in the sample solution is less than the detection limit, the detect line shows purple and the result is negative.

Technical indicator

Sensitivity:3 ppb (ng/mL)

Note: The final detection limit of sample equal to the result of sensitivity multiply by dilution ratio of sample pretreatment.

Detection limit:Urine---3ppb

Kits components

Item	Specifications
Detect card	50 T/kit
Manual	1 copy

Other supplies required

Instruments:Centrifuge

High-precision transferpettor: Single channel (20-200µL)

Sample pretreatment

1. Sample pretreatment Notice:Experimental apparatus should be clean, and the pipette should be disposable to avoid the experiment result be interfered by the contamination.

2. Sample pretreatment procedure:

Take clear upper urine sample to determine, the sample needs to be centrifuged at 4000 r/min for 10 min if turbid.

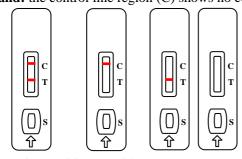
Note: Sampledilution factor: 1Detection limit:3ppb

Experiment procedure

- 1. Tear the aluminum foil bag of the detect card and take out the detect card, and put it on a smooth, clean table.
- 2. Take the prepared clear sample supernatant with the matching straw, add 2-3 drops (about 60uL) of sample to the sample well (S) vertically and slowly.
- 3. Keep the detect card at room temperature for 8-10 min, then judge the result. The result can only be considered as a reference if lasts for more than 10 min.

Judgment of result

Negative: the control line region (C) and the test line region (T) both show purple. . **Positive:** the control line region (C) shows purple, the test line region (T) shows no color. **Invalid:** the control line region (C) shows no color.



Negative Positive Invalid

Notes

- 1. Do not use product out of date or in a broken aluminum foil.
- 2. The detect card should be adjusted to room temperature after removed from the refrigerator before opening. The opening detect card should be used as soon as possible so as not to be invalid because of moisture.
- 3. Avoid of contacting the whitemembrane at the middle of the sample well.
- 4. The droplets cannot be mixing to avoid the cross-contaminant.
- 5. The tested sample should be clear, no turbidity particle and no bacterial pollution, otherwise it is easy to result in abnormal phenomena such as obstruction, unobvious color, etc., which affect the judgment of the experiment result.

Storage and valid period

Storage: Store at 2-30 °C withdry condition. **Valid Period:** 1 year, production date is on the packing box. Copyright ©2017-2018Elabscience Biotechnology Inc.All Rights Reserved