

**CLE-RAC-SAL (Clenbuterol-Ractopamine-Salbutamol) Lateral Flow Assay Kit**

Catalog No: E-FS-C127

40T

**Version Number:** V1.4  
**Replace version:** V1.3  
**Revision Date:** 2026.04.29

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.

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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 Immunochromatography assay. It can detect Clenbuterol--Ractopamine-Salbutamol (CLE-RAC-SAL) in urine sample. After adding the sample solution into the sample well of detection card, CLE-RAC-SAL in the sample solution combine with the gold-labelled antibody, so as to prevent the combining of gold-labelled antibody with CLE-RAC-SAL conjugate on the cellulose membrane. When the concentration of CLE-RAC-SAL 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 CLE-RAC-SAL in the sample solution is less than the detection limit, the detect line show color and the result is negative.

## Technical indicator

**Detection limit:** (CLE /RAC /SAL)

Urine ---3 ppb /3 ppb /5 ppb

## Kits components

Item	Specifications
Detection card (with disposable dropper)	40 T
Manual	1 copy

## Other materials required but not supplied

**Instruments:** Centrifuge.

## Notes

1. FOR RESEARCH USE ONLY. Do not use product out of date or in a broken aluminum foil.
2. The detection card should be adjusted to room temperature after removed from the refrigerator before opening. The opening detection card should be used as soon as possible so as not to be invalid because of moisture.
3. Avoid of contacting the white membrane at the middle of the sample well.
4. The disposable dropper 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.
6. If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.
7. 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.
8. **Each reagent is optimized for use in the E-FS-C127. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-FS-C127 with different lot numbers.**

## Storage and expiry date

**Storage:** Store at 2-30°C. With cool and dry environment.

**Expiry date:** expiration date is on the packing box.

## Experiment procedure

1. Take the urine sample to be tested. If the sample is turbid, centrifuge at 3000 rpm for 5 minutes or filter it (Note: Before sample loading, ensure the sample temperature returns to room temperature of 20~25°C before testing!).
2. Tear the aluminum foil bag of the detection card and take out the detection card, and put it on a smooth, clean table.
3. Take the prepared clear sample supernatant with the matching disposable dropper, add 2 drops of sample to the each sample well (S) vertically and slowly (Avoid foaming).
4. Incubate for 3-5 minutes and then judge the results immediately.

## Judgment of result

1. **Negative:** The control line region (C) and the test line region (T) both show a line. It indicates the content of CLE-RAC-SAL in the sample is lower than detection limit or the sample doesn't contain CLE-RAC-SAL.
2. **Positive:** Only the control line region (C) show a line. It indicates the content of CLE-RAC-SAL in the sample is higher than detection limit.
3. **Invalid:** The control line region (C) does not show a line. It indicates operation process is wrong or the test card is invalid.

