SAs (Sulfonamides) Lateral Flow Assay Kit

Catalog No: E-FS-C129

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.

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 the principle of Immunochromatography assay for the qualitative detection. It can detect SAs (Sulfonamides) in muscle sample. After adding the sample solution into the sample well of detection card, SAs in the sample solution combine with the gold-labelled antibody, so as to prevent the combining of gold-labelled antibody with SAs conjugate on the cellulose membrane. When the concentration of SAs in the sample solution is more than the detection limit, the detect line do not show color and the result is positive. When the concentration of SAs in the sample solution is less than the detection limit, the detect line show color and the result is negative.

Technical indicator

Sensitivity: 20 ppb (ng/mL)

Detection limit of sulfonamides:

Name	Sensitivity (ppb)
Sulfamethazine (SM2)	40
Sulfamonomethoxine (SMM)	40
Aristebon (SMD)	40
Sulfadimoxine (SDM')	40
Sulfamethoxypyridazine (SMP)	40
Sulfadimethoxine (SDT)	40
Sulfapirazinmetossina (SMZ)	40
Sulfanitran (SNT)	40
Sulfisomidine (SIM)	40
Sulfaquinoxaline (SQX)	40
Sulfamerazine (SMR)	40
Sulfadiazine (SD)	40
Sulfapyridine (SPD)	40

Detection limit (based on SM2 concentration): Muscle---40 ppb.

Kits components

Item	Specifications
Detection card (with disposable dropper)	50 T/kit
Sample extractant A	1 vial
Sample extractant B	1 vial
Reconstitution fluid	1 vial
Manual	1 copy

Note: All reagent bottle caps must be tightened to prevent evaporation and microbial pollution.

Other materials required but not supplied

Instruments: Homogenizer, Nitrogen Evaporators, Water bath, Centrifuge, Graduated pipette, Balance (sensibility 0.01g), Oscillators

High-precision transferpettor: Single channel (20-200 μL, 100-1000 μL)

Reagent: Ethyl acetate, N-hexane, NaOH, Concentrated HCl

Notes

1. FOR RESEARCH USE ONLY. Do not use product out of date or in a broken aluminum foil.
2. Bring detection card to room temperature before opening the aluminum foil. 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 and unobvious color 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-C129. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-FS-C129 with different lot numbers.

Storage and expiry date

Storage: Store at 2-30℃. With cool and dry environment.

Expiry date: expiration date is on the packing box.

Sample pretreatment

Restore all reagents and samples to room temperature before use.

1. Sample pretreatment Notice:

Experimental apparatus should be clean, and the disposable dropper should be disposable to avoid the experiment result be interfered by the contamination.

2. Sample pretreatment procedure:

Restore all reagents and samples to room temperature before use.

3.1 Pretreatment of muscle (Chicken, Pork, Fish, Shrimp, Duck) sample:

(1) Remove the skin and fat of sample, homogenize with homogenizer. Weigh 1±0.05 g of homogenized sample into a 10 mL centrifuge tube.

(2) Add 0.8 mL sample extractant A, mix vigorously by vortex mixer for 1 min.

(3) Add 4 mL sample extractant B, mix vigorously by vortex mixer for 3 min.

(4) Then centrifuge the sample for 5 min with 4000×g at room temperature.

(5) Transfer 0.5 mL sample supernatant into a new 5 mL centrifuge tube and evaporate it with nitrogen stream in 60C (140℉) water bath.

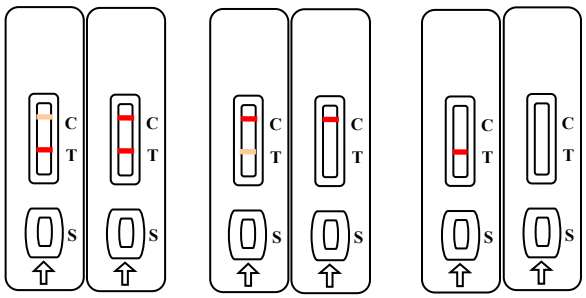
(6) Add 0.4 mL reconstitution fluid ， and then mix adequately by vortex mixer for 30 s.

Experiment procedure

1. Tear the aluminum foil bag of the detection card and take out the detection card, and put it on a smooth, clean table.
2. Take the prepared sample with the matching disposable dropper, add 120 μL of sample to the sample well (S) vertically and slowly (Avoid foaming).
3. Incubate for 5 minutes and then judge the results immediately.

Judgment of result

1. **Negative:** The control line region (C) show color, the test line region (T) shows equal or darker than line C. It indicates the content of SAs in the sample is lower than detection limit or the sample doesn't contain SAs.
2. **Positive:** The control line region (C) show color, the test line region (T) shows no color or lighter color than line C. It indicates the content of SAs in the sample is higher than detection limit.
3. **Invalid:** The control line region (C) shows no color. It indicates operation process is wrong or the test card is invalid.



Negative Positive Invalid