

SAR (Sarafloxacin) ELISA Kit

Catalog No: E-FS-E155

96T/96T*3

Version Number:	V1.3
Replace version:	V1.2
Revision Date:	2026.04.24

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 Sarafloxacin (SAR) in muscle, egg samples. 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, SAR in the samples or standard competes with coupled antigen on the solid phase supporter for sites of anti-SAR 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 SAR. The concentration of SAR 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°C; 40 min, 10-15 min

Detection limit: Muscle (Pork, Beef, Mutton, Chicken, Duck, Fish, Shrimp), Eggs---3 ppb; Serum (Cattle, Sheep) ---10 ppb

Cross-reactivity: Sarafloxacin---100%; Flumequine---100%; Enrofloxacin---130%; Enoxacin---110%; Norfloxacin---150%; Pefloxacin---150%; Danofloxacin---140%; Ofloxacin---110%; Difloxacin--130%; Fleroxacin---150%; Pazufloxacin---70%; Marbofloxacin---70%; Lomefloxacin---140%; Sparfloxacin ---25%; Oxolinic Acid ---200%; Ciprofloxacin---150%.

Sample recovery rate: 90%±30%.

Kits components

Item	Specifications
ELISA Microtiter plate	96 wells
Standard Liquid	1.5 mL each (ppb=ng/mL=ng/g) (0 ppb, 0.05 ppb, 0.15 ppb, 0.45 ppb, 1.35 ppb, 4.05 ppb)
HRP Conjugate	7 mL
Antibody Working Solution	7 mL
Substrate Reagent A	7 mL
Substrate Reagent B	7 mL
Stop Solution	7 mL
20×Concentrated Wash Buffer	25 mL
4×Concentrated Sample Diluent	60 mL
Plate Sealer	1 piece
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 (sensitivity 0.01 g).

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

Reagents: Anhydrous methanol, Concentrated phosphoric acid, Acetonitrile.

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°C.
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. FOR RESEARCH USE ONLY. ELISA Microtiter plate should be covered by plate sealer. Avoid the kit to strong light.
5. **Each reagent is optimized for use in the E-FS-E155. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-FS-E155 with different lot numbers.**
6. Substrate Reagent should be abandoned if it turns blue color. When OD value of standard (concentration: 0) < 0.8 unit (A450nm < 0.8), it indicates the reagent 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°C. 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°C. After opening, the kit is stable for up to 1 month.

Expiry date: expiration date is on the packing box.

Experimental preparation

Restore all reagents and samples to room temperature before use.

Open the microplate reader in advance, preheat the instrument, and set the testing parameters.

1. Sample pretreatment

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: **H₃PO₄ Solution**

Dissolve 2 mL of **Concentrated phosphoric acid** to 98 mL with deionized water.

Solution 2: **Sample Diluent**

Dilute **4×Concentrated Sample Diluent** with deionized water (4×Concentrated Sample Diluent (V): Deionized water (V) = 1:3).

Solution 3: **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 muscle Method 1 (Pork, Beef, Mutton, Chicken, Duck, Fish, Shrimp) sample:

- (1) Remove fat from sample, homogenize the sample with homogenizer.
- (2) Weigh 2.0±0.05 g of homogenate muscle sample into 50 mL centrifuge tube
- (3) Add 0.1 mL of **H₃PO₄ Solution** (Solution 1) and 6 mL **Acetonitrile**, vortex for 2 min, centrifuge at 4000 r/min at room temperature for 5 min.
- (4) Take 2 mL of the supernatant into 4 mL centrifugal tube, dry at 60-70°C of nitrogen evaporators or water bath.
- (5) Add 1 mL of **N-hexane**, vortex for 30 s, then add 0.5 mL of **Sample Diluent** (Solution 2), vortex for 30 s, centrifuge at 4000 r/min for 5 min at room temperature.
- (6) Remove the upper N-hexane and the impurities in the middle layer, take 50 µL of the clear liquid and add 450 µL of **Sample Diluent** (Solution 2), then vortex for 30 s.
- (7) Take 50 µL liquid to analysis.

Note: Sample dilution factor: 10, detection limit: 3 ppb

3.2 Pretreatment of muscle Method 2(Pork, Beef, Mutton, Chicken, Duck, Fish, Shrimp) sample:

- (1) Weigh 1±0.05 g of homogenate sample into 50 mL centrifuge tube, add 10 mL of deionized water, vortex for 1 min and mix fully.
- (2) Centrifuge at 4000 r/min for 5 min at room temperature.
- (3) Take 50 µL of supernatant to analysis (*Note: Do not aspirate impurities from the upper layer*).

Note: Sample dilution factor: 10, detection limit: 3 ppb

3.3 Pretreatment of eggs sample:

- (1) Take 100 μL of homogenate sample into 4 mL centrifuge tube, add 2 mL of **Wash Buffer** (Solution 3), vortex for 30 s.
- (2) Take 50 μL of liquid to analysis.

Note: Sample dilution factor: 20, detection limit: 3 ppb

3.3 Pretreatment of serum (Cattle, Sheep) sample:

- (1) Add 0.5 mL of sample into 4 mL centrifuge tube.
- (2) Add 0.5 mL of **Anhydrous methanol**, vortex for 30 s to dissolve it fully, Centrifuge at 4000 rpm at room temperature for 5 min.
- (3) Take 50 μL of the supernatant and add it to 950 μL of the **Sample Diluent** (Solution 2), then vortex for 30 s.
- (4) Take 50 μL of liquid to analysis.

Note: Sample dilution factor: 40, detection limit: 10 ppb

Assay procedure

Restore all reagents and samples to room temperature (25°C) 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°C.

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 (The sequence of adding liquid cannot be changed)**, cover the plate with plate sealer. Oscillate for 10 s gently to mix thoroughly, incubate at 25°C for 40 min in shading light.
3. **Wash:** uncover the sealer carefully, remove the liquid in each well. Immediately add 260 μL of **Wash Buffer** (Solution 3) to each well and wash. Repeat wash procedure for 4 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 100 μL of **Substrate Reagent A** and **Substrate Reagent B** mixture. (**Substrate Reagent A** and **Substrate Reagent B** are mixed 1:1 according to volume, must be fully mixed, the mixture is used within 5 minutes, avoid the use of metal container, avoid stirring reagents.) Gently oscillate for 10 s to mix thoroughly. Incubate 25°C for 10-15 min at in shading light.
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 5 min after stop reaction.

Result analysis

1. Absorbance (%)= $A/A_0 \times 100\%$

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.

Sarafloxacin (E-FS-E155) Standard Curve

