

Product Datasheet

Human TFF1 Matched Antibody Pair Kit PSB049Hu01 (96T x 10)

[Products overview]

Matched Antibody Pair Kit is composed of unlabeled capture antibody, Biotinylated detection antibody and a calibrated protein standard. The Matched Antibody Pair Kit can be used for quantifying natural and recombinant human Trefoil Factor 1 (TFF1) in ELISA, CLIA, ELISPOT, Luminex, Immunochromatography and other immunoassays. The Standard in the kit is recombinant TFF1 . Both capture and detection antibody are rabbit polyclonal antibodies.

[Components And Properties]

Components	Quantity	Form
Standard	200ng	Lyophilized, 1 vial
Capture Antibody	220µg / 0.39mL	Liquid, 1 vial, contains 0.1% sodium azide
Biotinylated Detection Antibody	20µg / 0.11mL	Liquid, 1 vial, contains 0.1% sodium azide

Notes: The kit contains raw materials for approximately 96 Tests x 10 plates. However, individual results may vary depending on the researcher's assay protocol and other variables.

[**Recommended Range / Dilution**]

Standard: Reconstitute the Standard with 1.0mL of Standard Diluent (Cat # IS051). The recommended Range of Standard curve is 0.156-10ng/mL.

Capture Antibody: Dilute 260 times with Coating Buffer (Cat # IS052). For example, to make enough for 1 plate, add 40uL capture antibody to 10.36mL Coating Buffer.

Biotinylated Detection Antibody: Dilute 910 times with Antibody Dilution Buffer (Cat # IS053). For example, to make enough for 1 plate, add 10uL Biotinylated Detection Antibody to 9.09mL Antibody Dilution Buffer.

Notes: The recommended Cloud-Clone's products of diluents and buffers are validated in the lab, other reagents selected for use can alter the performance of an immunoassay.

[**Storage**]

Avoid repeated freeze/thaw cycles. Store at 2-8°C for one month. Aliquot and store at -80°C for 12 months. Please make all solutions fresh before the experiment.

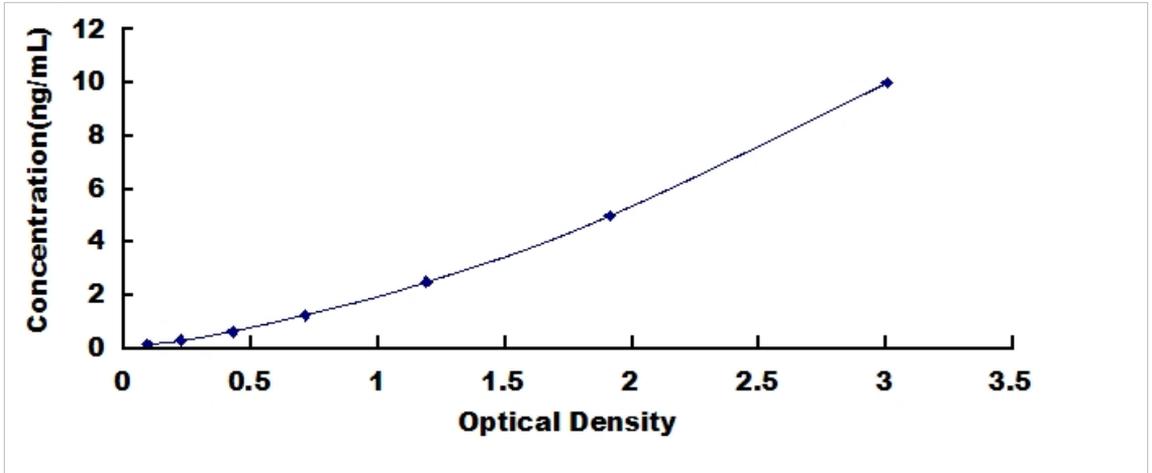
Notes: Please avoid pollution.

[**Recommended Buffers and Solutions**]

Cloud-Clone's product of Assay Kit DIY Support Pack 1 (Cat # IS077), which includes Coating Buffer, Blocking Buffer, Standard Diluent, Antibody Dilution Buffer, Streptavidin-HRP, HRP Dilution Buffer, Substrate, Solution, Wash Buffer, Stop Solution is highly recommended for reagent preparation.

[Typical Data]

Typical standard curve below is provided for reference only. A standard curve should be generated from each set of experiment.



[Recommended Assay Protocol]

1. Dilute the Capture Antibody to working concentration in Coating Buffer. Immediately coat the 96-well microplates with 100 μ L per well of the diluted Capture Antibody. Seal the plate and incubate overnight at 4°C or incubate at 37°C for 2 hours.
2. Aspirate wells and wash with 350 μ L of Wash Buffer per well, and let it sit for 1~2 minutes. Remove the remaining liquid by inverting and tapping the plate on absorbent paper.
3. Block plate with 200 μ L per well of Blocking Buffer for 1.5 hours at 37°C.
4. Repeat the aspiration/wash process as in step 2.
5. Add 100 μ L of different concentration of standards, samples and diluent into the appropriate wells. Cover with the Plate sealer. Incubate for 1 hour at 37°C.
6. Repeat the aspiration/wash process as in step 2.

7. Add 100µL of the working Biotinylated Detection Antibody working solution to each well, cover the wells, and incubate for 1 hour at 37°C.
8. Repeat the aspiration/wash process for 3 times as in step 2.
9. Add 100µL of the working solution of Streptavidin-HRP to each well, cover the wells, and incubate for 30 minutes at 37°C.
10. Repeat the aspiration/wash process for total 5 times as in step 2.
11. Add 90µL of TMB Substrate to each well. Cover the wells, and incubate for 10-20 minutes at 37°C. Protect from light.
12. Add 50µL of Stop Solution to each well. Mix the liquid by tapping the side of the plate.
13. Run the microplate reader and conduct measurement at 450nm immediately.