

## Product Datasheet

### Human SAA Matched Antibody Pair Kit PSA885Hu01 ( 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 Serum Amyloid A (SAA) in ELISA, CLIA, ELISPOT, Luminex, Immunochromatography and other immunoassays. The Standard in the kit is recombinant SAA. Both capture and detection antibody are mouse monoclonal antibodies.

#### [ Components And Properties ]

Components	Quantity	Form
<b>Standard</b>	4µg	Lyophilized, 1 vial
<b>Capture Antibody</b>	125µg / 0.07mL	Liquid, 1 vial, contains 0.1% sodium azide
<b>Biotinylated Detection Antibody</b>	5µg / 0.115mL	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 Buffers and Solutions ]

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

## [ Recommended Range / Dilution ]

**Standard:** Reconstitute the Standard with 1.0mL of Standard Diluent (Cat # IS051). The recommended Range of Standard curve is 3.12-200ng/mL.

**Capture Antibody:** Dilute 1430 times with Coating Buffer (Cat # IS052). For example, to make enough for 1 plate, add 7uL capture antibody to 10.003mL Coating Buffer.

**Biotinylated Detection Antibody:** Dilute 870 times with Antibody Dilution Buffer (Cat # IS053). For example, to make enough for 1 plate, add 11uL Biotinylated Detection Antibody to 9.559mL 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.

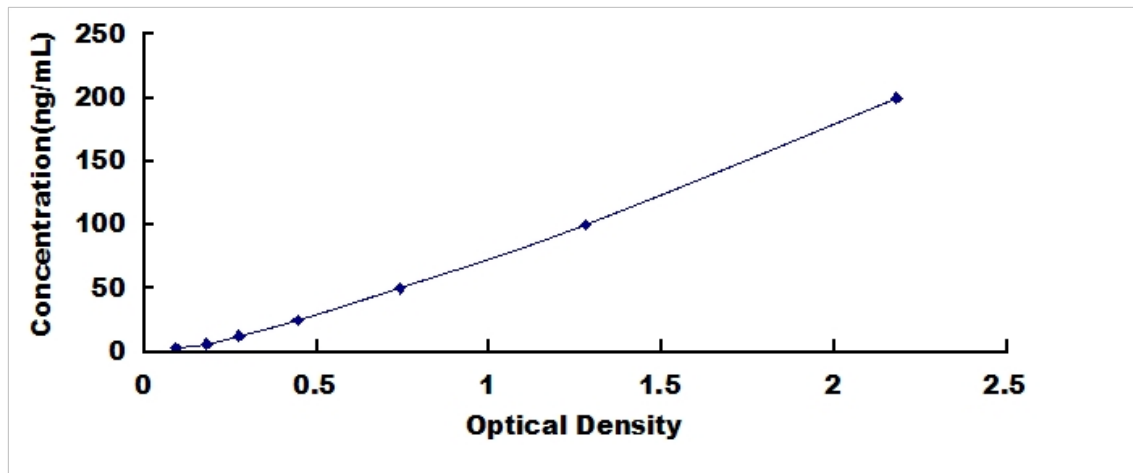
## [ Storage ]

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.

## [ 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 $\mu$ 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 $\mu$ 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 $\mu$ 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 $\mu$ L of different concentration of standards, samples 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 100 $\mu$ L 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 100 $\mu$ L 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 90 $\mu$ L of TMB Substrate to each well. Cover the wells, and incubate for 10-20 minutes at 37°C. Protect from light.
12. Add 50 $\mu$ L 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.