

Product Datasheet

Human RNASE3 Matched Antibody Pair Kit PSB758Hu01 (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 potentially be used for quantifying natural and recombinant Antibody Pair for Ribonuclease A3 (RNASE3) in ELISA, CLIA, ELISPOT, Luminex, Immunochromatography and other immunoassays. The Standard in the kit is recombinant RNASE3 . Both capture and detection antibody are mouse monoclonal antibodies.

[Components And Properties]

Components	Quantity	Form
Standard	5µg	Lyophilized, 1 vial
Capture Antibody	400µg / 0.8mL	Liquid, 1 vial, containing 0.05% Proclin-300, 50% glycerol
Biotinylated Detection Antibody	100µg / 0.5mL	Liquid, 1 vial, containing 0.05% Proclin-300, 50% glycerol

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 Standard Diluent (Cat # IS077). The recommended Range of Standard curve is 0.312-20ng/mL.

Capture Antibody: Dilute 125 times with Coating Buffer (Cat # IS077). For example, to make enough for 1 plate, add 80uL capture antibody to 9.92mL Coating Buffer.

Biotinylated Detection Antibody: Dilute 200 times with Detection Antibody Diluent (Cat # IS077). For example, to make enough for 1 plate, add 50uL Biotinylated Detection Antibody to 9.95mL 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.

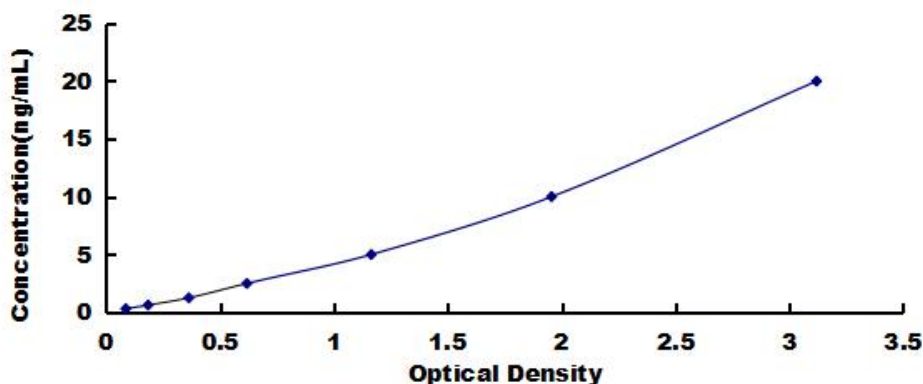
[Storage]

Avoid repeated freeze/thaw cycles. Store at 2-8°C for one month. Aliquot and store at -20°C for 12 months. Please make all solutions fresh before the experiment.

Notes: Please avoid contamination.

[Typical Data]

Typical standard curve below is provided for reference only. A standard curve should be generated for each 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 (Cat # IS077) 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 (Cat # IS077) 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 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 μ 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 100µL of the working solution of Streptavidin-HRP (Cat # IS077) 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 Substrate Solution (Cat # IS077) to each well. Cover the wells, and incubate for 10-20 minutes at 37°C. Protect from light.
12. Add 50µL of Stop Solution (Cat # IS077) to each well. Mix the liquid by tapping the side of the plate.
13. Run the microplate reader and conduct measurement at 450nm immediately.