

Product Data Sheet

Recombinant Anti-STAU1 Rabbit mAb

| Catalog # | Source | Reactivity | Applications |
|----------------|----------------|-------------------------------|--|
| CMA4118 | Rabbit | H, M, R | WB, IF/IC |
| Description | Re | ecombinant rabbit monocl | onal antibody to STAU1 |
| Immunogen | KI | LH-conjugated synthetic pe | eptide encompassing a sequence within human STAU1. |
| | Tł | he exact sequence is propr | ietary. |
| Purification | Tł | he antibody was purified b | y immunogen affinity chromatography. |
| Specificity | Re | ecognizes endogenous leve | els of STAU1 protein |
| Clonality | Μ | Ionoclonal | |
| Conjugation | | | |
| Form | Li | quid in PBS containing 50% | 6 glycerol, 0.2% BSA and 0.01% sodium azide. |
| Dilution | Ŵ | /B (1⁄500 - 1⁄1000), IF/IC (1 | /50 - 1/200) |
| Gene Symbol | ST | TAU1 | |
| Alternative Na | ames ST | TAU; Double-stranded RNA | -binding protein Staufen homolog 1 |
| Entrez Gene | 67 | 780 (Human); 20853 (Mou | se) |
| SwissProt | 0 | 95793 (Human); Q9Z108 (| Mouse) |
| Storage/Stabi | lity Sł | hipped at 4°C. Upon delive | ry aliquot and store at -20°C for one year. Avoid |
| | fr | eeze/thaw cycles. | |

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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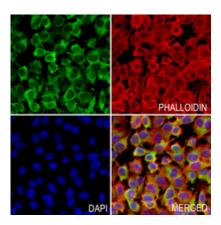
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Western blot analysis of STAU1 expression in A549 (A), MCF7 (B), K562 (C), mouse liver (D), mouse kidney (E), rat liver (F) whole cell lysates. (Predicted band size: 63 kD; Observed band size: 55 kD)



Immunofluorescent analysis of STAU1 staining in A375 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with an AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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