

Product Data Sheet

Anti-TAU (Phospho-T522) Antibody

Catalog # Source Reactivity Applications

CPA4725 Rabbit H, M, R, B WB, IH

Description Rabbit polyclonal antibody to TAU (Phospho-T522)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

T522 of human TAU protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of TAU protein only when phosphorylated at T522.

Clonality Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/50 - 1/100)

Gene Symbol MAPT

Alternative Names MAPTL; MTBT1; TAU; Microtubule-associated protein tau; Neurofibrillary tangle

protein; Paired helical filament-tau; PHF-tau

Entrez Gene 4137 (Human); 17762 (Mouse)

SwissProt P10636 (Human); P10637 (Mouse); P19332 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/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

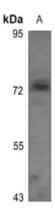
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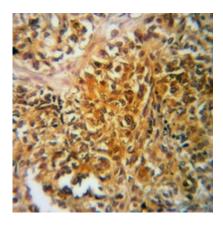




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Western blot analysis of TAU (Phospho-T522) expression in rat brain (A) whole cell lysates. (Predicted band size: 78 kD; Observed band size: 50-80 kD)



Immunohistochemical analysis of TAU (Phospho-T522) staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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