

Store at
-20°C
#11834

Phospho-Tau (Ser202) Antibody

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100 µl (10 western blots)

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Entrez-Gene ID #4137
UniProt ID #P10636-8

rev. 06/19/14

For Research Use Only. Not For Use In Diagnostic Procedures.

Applications
W
Endogenous

Species Cross-Reactivity*
H, M, R

Molecular Wt.
50-80 kDa

Isotype
Rabbit**

Background: Tau is a heterogeneous microtubule-associated protein that promotes and stabilizes microtubule assembly, especially in axons. Six isoforms with different amino-terminal inserts and different numbers of tandem repeats near the carboxy terminus have been identified, and tau is hyperphosphorylated at approximately 25 sites by Erk, GSK-3, and CDK5 (1,2). Phosphorylation decreases the ability of tau to bind to microtubules. Neurofibrillary tangles are a major hallmark of Alzheimer's disease; these tangles are bundles of paired helical filaments composed of hyperphosphorylated tau. In particular, phosphorylation at Ser396 by GSK-3 or CDK5 destabilizes microtubules. Furthermore, research studies have shown that inclusions of tau are found in a number of other neurodegenerative diseases, collectively known as tauopathies (1,3).

Investigators have shown that Tau is phosphorylated during development and hyper-phosphorylated at Ser202 in Alzheimer's disease (4).

Specificity/Sensitivity: Phospho-Tau (Ser202) Antibody recognizes endogenous levels of Tau protein only when phosphorylated at Ser202.

Source/Purification: Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser202 of human Tau protein. Antibodies are purified by protein A and peptide affinity chromatography.

Background References:

- (1) Johnson, G.V. and Stoothoff, W.H. (2004) *J. Cell Sci.* 117, 5721-5729.
- (2) Hanger, D. P. et al. (1998) *J. Neurochem.* 71, 2465-2476.
- (3) Bramblett, G. T. et al. (1993) *Neuron* 10, 1089-1099.
- (4) Goedert, M. et al. (1993) *Proc Natl Acad Sci USA* 90, 5066-70.

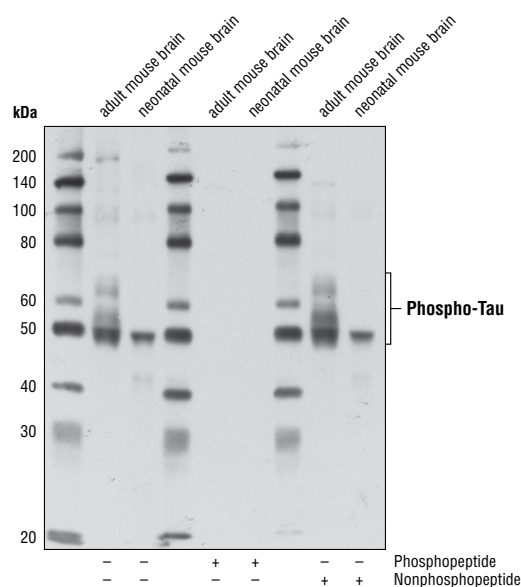
Storage: Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.

***Species cross-reactivity is determined by western blot.**

****Anti-rabbit secondary antibodies must be used to detect this antibody.**

Recommended Antibody Dilutions:

Western blotting 1:1000



Western blot analysis of extracts from adult and neonatal mouse brain using Phospho-Tau (Ser202) Antibody. The phospho-specificity of the antibody was verified by blocking with a phospho or nonphosphopeptide.

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v BSA, 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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Applications: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide **Species Cross-Reactivity:** H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—Horse All—all species expected *Species enclosed in parentheses are predicted to react based on 100% homology.