

Phospho-Tau (Ser199) Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 50-80	Source/Isotype: Rabbit	UniProt ID: #P10636-8	Entrez-Gene Id: 4137
Product Usage Information	!	Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Phospho-Tau (Ser199) Antibodyrecognizes endogenous levels of Tau protein only when phosphorylated at Ser199. This antibody also detects a 110 kDa band of unknown origin.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser199 of human Tau protein. Antibodies are purified by protein A and peptide affinity chromatography.				
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, glycogen synthase kinase-3 (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 (AD); these tangles are bundles of paired helical filaments (PHFs) 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). Ser199 of Tau is phosphorylated by various kinases such as GSK-3β, AMP-activated protein kinase (AMPK), and dual specificity tyrosine-phosphorylation-regulated kinase 1A (Dyrk1A) (4-7). Phosphorylation of Ser199 of Tau is an early event in the pathogenesis of Alzheimer's disease (8,9).				
Background References		 Johnson, G.V. and Stoothoff, W.H. (2004) J Cell Sci 117, 5721-9. Hanger, D.P. et al. (1998) J Neurochem 71, 2465-76. Bramblett, G.T. et al. (1993) Neuron 10, 1089-99. Qian, W. et al. (2010) J Alzheimers Dis 19, 1221-9. Leroy, A. et al. (2010) J Biol Chem 285, 33435-44. Thornton, C. et al. (2011) Biochem J 434, 503-12. Jin, N. et al. (2015) J Biol Chem 290, 15219-37. Mondragón-Rodríguez, S. et al. (2008) Neuropathol Appl Neurobiol 34, 62-75. Mondragón-Rodríguez, S. et al. (2008) Int J Exp Pathol 89, 81-90. 				
Species Reacti	vity	Species reactivity is d	etermined by testir	ng in at least one approve	ed application (e.g.,	western blot).
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Western Blot Buffer

Applications Key

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

W: Western Blotting

Cross-Reactivity Key

H: Human M: Mouse R: Rat

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