Store at -200

Phospho-Tau (Ser199) Antibody



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Applications: WB	Reactivity:	Sensitivity: Endogenous	MW (kDa): 50-80	Source: Rabbit	UniProt ID: #P10636-8	Entrez-Gene Id: 4137	
	HMR						
Product Usage Information	Арр	Application			Dilution		
	We	Western Blotting			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	espe repe 25 s abilii (AD) In pa	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					

Ser199 of Tau is phosphorylated by various kinases such as GSK-3\(\beta\), 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

- 1. Johnson, G.V. and Stoothoff, W.H. (2004) J Cell Sci 117, 5721-9.
- 2. Hanger, D.P. et al. (1998) J Neurochem 71, 2465-76.
- 3. Bramblett, G.T. et al. (1993) Neuron 10, 1089-99.

diseases, collectively known as tauopathies (1,3).

- 4. Qian, W. et al. (2010) J Alzheimers Dis 19, 1221-9.
- 5. Leroy, A. et al. (2010) J Biol Chem 285, 33435-44.
- 6. Thornton, C. et al. (2011) Biochem J 434, 503-12.
- 7. Jin, N. et al. (2015) J Biol Chem 290, 15219-37.
- 8. Mondragón-Rodríguez, S. et al. (2008) Neuropathol Appl Neurobiol 34, 62-75.
- 9. Mondragón-Rodríguez, S. et al. (2008) Int J Exp Pathol 89, 81-90.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

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.

Applications Key

WB: Western Blotting

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus 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

GP: Guinea Pig Rab: rabbit All: all species expected

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Limited Uses

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