



Tau (phospho Ser214) Polyclonal Antibody

Catalog NoBYab-03015IsotypeIgGReactivityHuman;Mouse;RatApplicationsWB;ELISAGene NameMAPTProtein NameMicrotubule-associated protein tauImmunogenThe antiserum was produced against synthesized peptide de Tau around the phosphorylation site of Ser214. AA range:50	rived from human
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Specificity Phospho-Tau (S214) Polyclonal Antibody detects endogenou protein only when phosphorylated at S214.	us levels of Tau
FormulationLiquid in PBS containing 50% glycerol, 0.5% BSA and 0.02%	
Source Polyclonal, Rabbit,IgG	
PurificationThe antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.	
DilutionWestern Blot: 1/500 - 1/2000. ELISA: 1/5000. Not yet tested	in other applications.
Concentration 1 mg/ml	
Purity ≥90%	
Storage Stability -20°C/1 year	
SynonymsMAPT; MAPTL; MTBT1; TAU; Microtubule-associated protein tangle protein; Paired helical filament-tau; PHF-tau	n tau; Neurofibrillary
Observed Band50-85kD	
	protein ; Cytoplasmic
Cell Pathway Cytoplasm, cytosol . Cell membrane ; Peripheral membrane j side . Cytoplasm, cytoskeleton . Cell projection, axon . Cell p Secreted . Mostly found in the axons of neurons, in the cytos with plasma membrane components (PubMed:10747907). C secretion is dependent on protein unfolding and facilitated by TMED10; it results in protein translocation from the cytoplasm (endoplasmic reticulum-Golgi intermediate compartment) follo entry and secretion (PubMed:32272059)	ol [°] and in association an be secreted; the the cargo receptor n into the ERGIC
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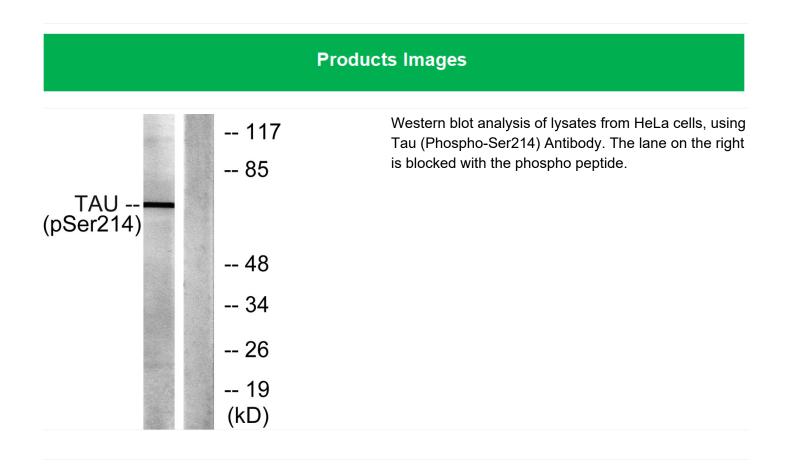
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	found in fetal brain, whereas three-repeat (type I) tau is found in both adult and fetal brain.,disease:Defects in MAPT are a cause of corticobasal degeneration (CBD). It is marked by extrapyramidal signs and apraxia and can be associated with memory loss. Neuropathologic features may overlap Alzheimer disease, progressive supranuclear palsy, and Parkinson disease.,disease:Defects in MAPT are a cause of frontotemporal dementia and parkinsonism linked to chromosome 17 (FTDP17) [MIM:600274, 172700]; also called frontotemporal dementia (FTD) or historically termed Pick complex. This form
Background	This gene encodes the microtubule-associated protein tau (MAPT) whose transcript undergoes complex, regulated alternative splicing, giving rise to several mRNA species. MAPT transcripts are differentially expressed in the nervous system, depending on stage of neuronal maturation and neuron type. MAPT gene mutations have been associated with several neurodegenerative disorders such as Alzheimer's disease, Pick's disease, frontotemporal dementia, cortico-basal degeneration and progressive supranuclear palsy. [provided by RefSeq, Jul 2008],
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.



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