

Tau (phospho-Ser416) rabbit pAb

Cat No.:ES12811

For research use only

Overview

Product Name	Tau (phospho-Ser416) rabbit pAb
Host species	Rabbit
Applications	WB
Species Cross-Reactivity	Human;Mouse;Rat
Recommended dilutions	WB 1:1000-2000
Immunogen	Synthesized phosho peptide around human Tau
	(Ser416)
Specificity	This antibody detects endogenous levels of Human
	Mouse Rat Tau (phospho-Ser416)
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and
	0.02% sodium azide.
Storage	Store at -20 $^\circ\!\mathrm{C}$. Avoid repeated freeze-thaw cycles.
Protein Name	Tau (Ser416)
Gene Name	MAPT MAPTL MTBT1 TAU
Cellular localization	Cytoplasm, cytosol . Cell membrane ; Peripheral
	membrane protein ; Cytoplasmic side . Cytoplasm,
	cytoskeleton . Cell projection, axon . Cell projection,
	dendrite . Secreted . Mostly found in the axons of
	neurons, in the cytosol and in association with
	plasma membrane components
	(PubMed:10747907). Can be secreted; the secretion
	is dependent on protein unfolding and facilitated by
	the cargo receptor TMED10; it results in protein
	translocation from the cytoplasm into the ERGIC
	(endoplasmic reticulum-Golgi intermediate
	compartment) followed by vesicle entry and
	secretion (PubMed:32272059)
Purification	The antibody was affinity-purified from rabbit
	antiserum by affinity-chromatography using
	epitope-specific immunogen.
Clonality	Polyclonal
Concentration	1 mg/ml
Observed band	50-85kD
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Human Gene ID Human Swiss-Prot Number Alternative Names

Background

4137 P10636

Microtubule-associated protein tau (Neurofibrillary tangle protein) (Paired helical filament-tau) (PHF-tau)

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],



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