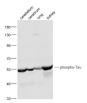
[Primary Antibody]

phospho-Tau (Ser422) Rabbit pAb

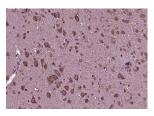


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- DATASHEET		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500)
Clonality: Polyclonal		IHC-F (1:100-500)
GenelD: 4137	SWISS: P10636	IF (1:100-500)
Target: Tau (Ser422)		Reactivity: Human, Mouse, Rat (predicted: Rabbit, Cow, Dog)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human Tau around the phosphorylation site of Ser422: VD(p-S)PQ.		
Purification: affinity purified by	Protein A	
Concentration: 1mg/ml		Predicted MW.: ^{52/79} kDa
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Cell membrane ,Cytoplasm Location:
stability, and migh maintenance of ne microtubules whil membrane compo protein between b localization (in the defined by the cer the cytoskeleton w play a role in its st the axons of neuro plasma membran is expressed in the	mportant Promotes microtubule assembly and at be involved in the establishment and euronal polarity. The C-terminus binds axonal e the N-terminus binds neural plasma onents, suggesting that tau functions as a linker both. Axonal polarity is predetermined by tau e neuronal cell) in the domain of the cell body attrosome. The short isoforms allow plasticity of whereas the longer isoforms may preferentially abilization. Tau proteins subcellular located in ons, in the cytoso I and in association with e components. It expressed in neurons. PNS-tau e peripheral nervous system while the others are entral nervous system.	
- VALIDATION IMAGES -		



Sample: Cerebellum (Mouse) Lysate at 40 ug Cerebrum (Rat) Lysate at 40 ug Lung (Mouse) Lysate at 40 ug Kidney (Mouse) Lysate at 40 ug Primary: Anti- phospho-Tau (Ser422) (bs-3489R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 83 kD Observed band size: 52 kD



Paraformaldehyde-fixed, paraffin embedded (Rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (phospho-Tau (Ser422)) Polyclonal Antibody, Unconjugated (bs-3489R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.

- SELECTED CITATIONS -----

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- [IF=6.2] Bo Zhang. et al. Arsenic exposure activates microglia, inducing neuroinflammation and promoting the occurrence and development of Alzheimer's disease-like neurodegeneration in mice. ECOTOX ENVIRON SAFE. 2025 Jun;297:118251 IHC, IF ;MOUSE. 40294499
- [IF=4] Wang Xiaoping. et al. S1PR2 Regulates Autophagy Through the AKT/mTOR Pathway to Promote Pathological Damage in Alzheimer's Disease. J ALZHEIMERS DIS. 2023 Nov;Preprint(Preprint):1-16 WB ;Human. 38007654