## [ Primary Antibody ]

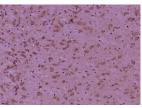
## phospho-Tau (Ser396) Rabbit pAb



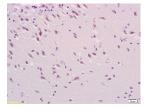
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- DATASHEET		400-901-9800
Host: Rabbit	<b>Isotype:</b> IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		<b>IHC-P</b> (1:100-500)
<b>GenelD:</b> 4137	SWISS: P10636	IHC-F (1:100-500) IF (1:100-500)
	500551 10050	<b>IF</b> (1.100-500)
Target: Tau (Ser396)		Reactivity: Mouse, Rat
<b>Immunogen:</b> KLH conjugated Synthesised phosphopeptide derived from human Tau around the phosphorylation site of Ser396: KT(p-S)T.		(predicted: Human, Dog)
Purification: affinity purified	by Protein A	
Concentration: 1mg/ml		Predicted MW.: <sup>46/79</sup> 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 mi maintenance of microtubules wi membrane com protein betweer localization (in t defined by the c the cytoskeletor play a role in its the axons of neu plasma membra is expressed in the	e important Promotes microtubule assembly and ght be involved in the establishment and neuronal polarity. The C-terminus binds axonal hile the N-terminus binds neural plasma ponents, suggesting that tau functions as a linker n both. Axonal polarity is predetermined by tau the neuronal cell) in the domain of the cell body entrosome. The short isoforms allow plasticity of n whereas the longer isoforms may preferentially stabilization. Tau proteins subcellular located in urons, in the cytoso l and in association with ane components. It expressed in neurons. PNS-tau he peripheral nervous system while the others are e central nervous system.	
- VALIDATION IMAGES -		I

Sample: Cerebrum (Rat) Lysate at 40 ug Primary: Anti- phospho-Tau (Ser396) (bs-3446R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 52/79 kD Observed band size: 52 kD



Paraformaldehyde-fixed, paraffin embedded (Mouse 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 (Ser396)) Polyclonal Antibody, Unconjugated (bs-3446R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-phospho-Tau protein(Ser396) Polyclonal Antibody, Unconjugated(bs-3446R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining

## - SELECTED CITATIONS -

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- [IF=3.5] Singh Nidhi Anand K.. et al. Ellagic Acid Reverses Alterations in the Expression of AMPA Receptor and Its Scaffolding Proteins in the Cerebral Cortex and Memory Decline in STZ-sporadic Alzheimer's Disease Mouse Model. PSYCHOPHARMACOLOGY. 2024 Jun;:1-15 WB ;MOUSE. 38842699
- [IF=2.65] Cai, Zhiyou, Yong Yan, and Yonglong Wang. "Minocycline alleviates beta-amyloid protein and tau pathology via restraining neuroinflammation induced by diabetic metabolic disorder." Clinical Interventions in Aging 8 (2013): 1089-1095. WB ;="Rat". 23983461
- [IF=3.499] Ding J et al. The Effect of α-Synuclein and Tau in Methamphetamine Induced Neurotoxicity in Vivo and in Vitro. Toxicol Lett. 2020 Feb 1;319:213-224. IF ;Mouse. 31783120
- [IF=1.89] Ku et al. Synergistic effects of particulate matter (PM2.5) and sulfur dioxide (SO2) on neurodegeneration via the microRNA-mediated regulation of tau phosphorylation. (2017) Toxicol.Res.(Camb). 6:7-16 WB ;MOUSE. 30090473