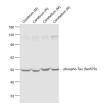
[Primary Antibody]

phospho-Tau (Ser579) Rabbit pAb

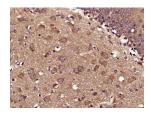


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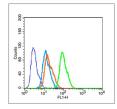
– DATASHEET –		400-901-9800
Host: Rabbit Clonality: Polyclon	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500)
GeneID: 4137	SWISS: P10636	IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (1µg/Test)
Target: phospho-Tau (Ser579) Immunogen: KLH conjugated Synthesised phosphopeptide derived from human Tau around the phosphorylation site of Ser579: IG(p-S)TE. Purification: affinity purified by Protein A		Reactivity: Human, Mouse, Rat (predicted: Rabbit, Cow, Dog, Horse)
Concentration: 1mg/ml 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.		Predicted 52/79 kDa MW.: ^{52/79} kDa Subcellular Location: ^{Cell} membrane ,Cytoplasm
Background: Tau proteins are important Promotes microtubule assembly and stability, and might be involved in the establishment and maintenance of neuronal polarity. The C-terminus binds axonal microtubules while the N-terminus binds neural plasma membrane components, suggesting that tau functions as a linker protein between both. Axonal polarity is predetermined by tau localization (in the neuronal cell) in the domain of the cell body defined by the centrosome. The short isoforms allow plasticity of the cytoskeleton whereas the longer isoforms may preferentially play a role in its stabilization. Tau proteins subcellular located in the axons of neurons, in the cytoso l and in association with plasma membrane components. It expressed in neurons. PNS-tau is expressed in the central nervous system.		
- VALIDATION IMAG	ies	1



Sample: Cerebrum (Mouse) Lysate at 40 ug Cerebrum (Rat) Lysate at 40 ug Cerebellum (Mouse) Lysate at 40 ug Cerebellum (Rat) Lysate at 40 ug Primary: Anti-phospho-Tau (Ser579) (bs-10108R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 50-70 kD Observed band size: 50 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 (Ser579)) Polyclonal Antibody, Unconjugated (bs-10108R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control (blue line): MCF 7 (fixed with 70% methanol (Overnight at $4^\circ C)$ and then permeabilized with 90% ice-cold methanol for 30 min on ice). Primary Antibody (green line): Rabbit Anti- phospho-Tau (Ser579) antibody (bs-10108R),Dilution: 1 μ g/10^5 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat antirabbit IgG-FITC,Dilution: 1µg /test.