bs-3633R

[Primary Antibody]

Synaptopodin Rabbit pAb



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- DATASHEFT		400-901-9800	
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)	
Clonality: Polyclonal		IHC-P (1:100-500)	
GenelD: 11346	SWISS: Q8N3V7	IF (1:100-500)	
Target: Synaptopodin		Flow-Cyt (1µg /test)	
Immunogen: KLH conjugated synthetic peptide derived from human Synaptopodin: 601-700/903.		Reactivity: Human, Mouse, Rat	
Purification: affinity purified by	Protein A		
Concentration: 1mg/ml		Prodicted	
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		Subcellular	
freeze/thaw cycles.		Location:	
actin-based cell shape and motility. May be essential for the formation of spine apparatuses in spines of telencephalic neurons, involved in synaptic plasticity. The name synaptopodin derives from the protein's associations with postsynaptic densities and		,	

- VALIDATION IMAGES

dendritic spines and with renal podocytes.



Sample: Cerebrum (Rat) Lysate at 40 ug Primary: Anti-Synaptopodin (bs-3633R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 102 kD Observed band size: 76 kD



Sample: Brain (Mouse) Lysate at 40 ug Primary: Anti-Synaptopodin (bs-3633R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 102 kD Observed band size: 102 kD



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by microwave in sodium citrate buffer (pH6.0) ; Block endogenous peroxidase by 3% hydrogen peroxide for 30 minutes; Blocking buffer (3% BSA) at RT for 30min; Antibody incubation with (Synaptopodin) Polyclonal Antibody, Unconjugated (bs-3633R) at 1:400 overnight at 4°C, followed by conjugation to the secondary antibody (labeled with HRP) and DAB staining.



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 (Synaptopodin) Polyclonal Antibody, Unconjugated (bs-3633R) at 1:400 overnight at 4°C, followed by operating



Blank control (black line) :SH-SY5Y. Primary Antibody (green line): Rabbit Anti-Synaptopodin antibody (bs-3633R) Dilution:1ug/Test; Secondary Antibody (white blue line) : Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line) : Normal Rabbit IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for



Positive control: RSC96 Isotype Control Antibody: Rabbit IgG; Secondary Antibody: Goat anti-rabbit IgG-FITC; Dilution: 1:200 in 1 X PBS containing 0.5% BSA Primary Antibody catalog number: bs-3633R; Dilution: 1µg in 100 µl 1X PBS containing 0.5% BSA 20 min at -20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

- [IF=3.73] Zhang, Xueming, et al. "Resolvin D1 Protects Podocytes in Adriamycin-Induced Nephropathy through Modulation of 14-3-3β Acetylation." PLOS ONE 8.6 (2013): e67471. IP ;="Mouse". 23840712
- [IF=0] Garovic, Vesna D., and Muthuvel Jayachandran. "DETECTING PODOCYTE INJURY IN DIABETIC NEPHROPATHY AND GLOMERULONEPHRITIS." U.S. Patent No. 20,170,003,299. 5 Jan. 2017. Other ;="Human". U.S.PatentNo.20,170,003,299.5