bs-3443R

[Primary Antibody]

Phospho-Tuberin (Tyr1571) Rabbit pAb



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– DATASHEET –		400-901-9800	400-901-9800	
Host: Rabbit	Isotype: IgG	Applications: Flow-Cyt (2ug/Tes	t)	
Clonality: Polyclonal		ICC/IF (1:100)		
GenelD: 7249	SWISS: P49815		Reactivity: Human (predicted: Mouse,	
Target: Phospho-Tuberin (Tyr1571)			Rat, Rabbit, Pig, Cow, Chicken, Dog, Horse)	
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human Tuberin around the phosphorylation site of Thr1571: R(p-Y)TE.		VITE	Predicted MW.: 180-220 kDa	
Purification: affinity purified by Protein A		MW.: ^{180-220 kDa}		
Concentration: 1mg/ml		Subcellular	Subcellular Location: ^{Cell} membrane ,Cytoplasm	
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.		J70		
Background: Mutations in this gene lead to tuberous sclerosis complex. Its gene product is believed to be a tumor suppressor and is able to stimulate specific GTPases. The protein associates with hamartin in a cytosolic complex, possibly acting as a chaperone for		to namartin		

– VALIDATION IMAGES



Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (Phospho-Tuberin (Tyr1571)) polyclonal Antibody, Unconjugated (bs-3443R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



hamartin. Alternative splicing results in multiple transcript variants

encoding different isoforms. [provided by RefSeq].

Blank control (black line) :Hela. Primary Antibody (green line): Rabbit Anti-Phospho-Tuberin (Tyr1571) antibody (bs-3443R) Dilution:2ug/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 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.