bs-0028R

- DATASHEET -

[Primary Antibody]

GSK-3 Beta Rabbit pAb



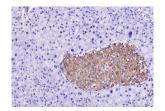
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

bbit Isotype: IgG	Appli
yclonal	
SWISS: P49841	
K-3 Beta	
H conjugated synthetic peptide derived from human GSł ta: 341-420/420.	K-3 Re
nity purified by Protein A	
g/ml	P
1M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% cerol.)
	yclonal 2 SWISS: P49841 K-3 Beta H conjugated synthetic peptide derived from human GSH a: 341-420/420. nity purified by Protein A g/ml IM TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

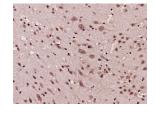
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: The protein encoded by this gene is a serine-threonine kinase, belonging to the glycogen synthase kinase subfamily. It is involved in energy metabolism, neuronal cell development, and body pattern formation. Polymorphisms in this gene have been implicated in modifying risk of Parkinson disease, and studies in mice show that overexpression of this gene may be relevant to the pathogenesis of Alzheimer disease. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Sep 2009]

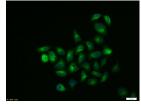
VALIDATION IMAGES



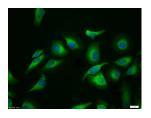
Paraformaldehyde-fixed, paraffin embedded (rat pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (GSK-3 Beta) Polyclonal Antibody, Unconjugated (bs-0028R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



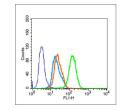
Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (GSK-3 Beta) Polyclonal Antibody, Unconjugated (bs-0028R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



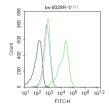
Tissue/cell:Hela cell; 4% Paraformaldehydefixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (GSK-3 Beta) polyclonal Antibody, Unconjugated (bs-0028R) 1:100, 90 minutes at 37°C; followed by a FITC conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Tissue/cell:Hela cell; 4% Paraformaldehydefixed; 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 (GSK-3 Beta) polyclonal Antibody, Unconjugated



Blank control (blue line): MCF 7 (blue). Primary Antibody (green line): Rabbit Anti- GSK-3 Beta (CT) antibody (bs-0028R) Dilution: 1µg /10^5 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue



Blank control: A431. Primary Antibody (green line): Rabbit Anti-GSK-3 Beta antibody (bs-0028R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF488

pplications: IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (1µg /test) ICC/IF (1:100)

Reactivity: Human, Mouse, Rat

Predicted MW.: 47 kDa

Subcellular Cell membrane ,Cytoplasm Location: ,Nucleus (bs-0028R) 1:100, 90 minutes at 37°C; followed by a FITC conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei. line): Goat anti-rabbit IgG-FITC Dilution: 1µg /test. Protocol The cells were fixed with 70% methanol (Overnight at 4°C) and then permeabilized with 90% ice-cold methanol for 30 min on ice. Cells stained with Primary Antibody for 30 min at room temperature. The cells were then incubated in 1 X PBS/2%BSA/10% goat serum to block nonspecific protein-protein interactions followed by the antibody for 15 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed. Dilution: 1µg /test. 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.

- SELECTED CITATIONS -

- [IF=15.304] Yao Lei. et al. Phytochemical natural killer cells reprogram tumor microenvironment for potent immunotherapy of solid tumors. BIOMATERIALS. 2022 Jun;:121635 WB ;MOUSE. 10.1016/j.biomaterials.2022.121635
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- [IF=5.895] Yu-Sheng Shi. et al. Pteris laeta Wall. and Its New Phytochemical, Pterosinsade A, Promote Hippocampal Neurogenesis via Activating the Wnt Signaling Pathway. J AGR FOOD CHEM. 2023;XXXX(XXX):XXX-XXX WB ;Murine. 36892329
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- [IF=5.108] Yuan C et al. OAB-14, a bexarotene derivative, improves Alzheimer's disease-related pathologies and cognitive impairments by increasing β-amyloid clearance in APP/PS1 mice. (2019) Biochim Biophys Acta Mol Basis Dis.Jan;1865(1):161-180. WB ;MOUSE. 30389579