bs-0128R

[Primary Antibody]

PIK3R1 Rabbit pAb



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- DATASHEET -		400-901-9800
Host: Rab	bit Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Poly	<i>y</i> clonal	IHC-P (1:100-500) IHC-F (1:100-500)
GenelD: 529	5 SWISS: P27986	IF (1:100-500)
Target: PIKS	3R1	Flow-Cyt (1µg/Test) ICC/IF (1:100)
lmmunogen: KLH p85	conjugated synthetic peptide derived from human PI3 ki subunit alpha: 501-600/724.	nase Reactivity: Human, Mouse, Rat
Purification: affir	nity purified by Protein A	(predicted: Cow, Chicken, Dog. Horse)
Concentration: 1mg	g/ml	
Storage: 0.01 Glyc Ship free	M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% cerol. oped at 4°C. Store at -20°C for one year. Avoid repeated vze/thaw cycles.	Predicted MW.: ⁸⁰ kDa Subcellular Cell membrane ,Cytoplasm
Background: The kina resp path kDa is ar p11 kina inte neci	enzyme phosphatidylinositol 3 kinase (PI3 kinase) is a lip ise that generates phosphatidylinositol 3, 4, 5-triphospha ponse to receptor activation in many signal transduction hways. Class IA PI3Ks exist as a heterodimer of a catalytic (p110) and a regulatory p85 subunit (e.g. p85 alpha). p85 n adaptor molecule that regulates the activity of the cataly 0 subunit by binding to phosphorylated receptor tyrosine ases (RTKs) through its SH2 domain and mediating the reaction between p110 and the plasma membrane. p85 alp essary for insulin-stimulated increase in glucose untake a	id te in 110 alpha ytic e pha is nd

– VALIDATION IMAGES



Sample: Heart (mouse) Lysate at 40 ug Primary: Anti- PI3K p85 (bs-0128R)at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 80kD Observed band size: 85 kD



glycogen synthesis in insulin-sensitive tissues.

Paraformaldehyde-fixed, paraffin embedded (rat skeletal muscle); 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 (PI 3 Kinase p85 alpha) Polyclonal Antibody, Unconjugated (bs-0128R) 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 skeletal muscle); 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 (PI 3 Kinase p85 alpha) Polyclonal Antibody, Unconjugated (bs-0128R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Tissue/cell: HepG2 cell; 4% Paraformaldehydefixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum,



Tissue/cell: NIH/3T3 cell; 4% Paraformaldehydefixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum,



Positive control: H9C2(2% Paraformaldehydefixed) Isotype Control Antibody Antibody: Rabbit IgG; Dilution: 1µg in 100 µl 1 X PBS containing

C-0005) at 37°C for 20 min; Antibody incubation with (PI3K p85) polyclonal Antibody, Unconjugated (bs-0128R) 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. C-0005) at 37°C for 20 min; Antibody incubation with (PI 3 Kinase p85 alpha) polyclonal Antibody, Unconjugated (bs-0128R) 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. 0.5% BSA Secondary Antibody Antibody: Goat anti-rabbit IgG-FITC; Dilution: 1:200 in 1 X PBS containing 0.5% BSA Primary Antibody Supplier catalog number: bs-1297R; Dilution: 1µg in 100 µl 1X PBS containing 0.5% BSA

- SELECTED CITATIONS -

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- [IF=7.419] Hong Kyu Lee. et al. TGF-β2 antisense oligonucleotide enhances T-cell mediated anti-tumor activities by IL-2 via attenuation of fibrotic reaction in a humanized mouse model of pancreatic ductal adenocarcinoma. BIOMED PHARMACOTHER. 2023 Mar;159:114212 WB ;Mouse. 36610224
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- [IF=5.895] Bendong Yang. et al. Naringenin Ameliorates Hyperuricemia by Regulating Renal Uric Acid Excretion via the PI3K/AKT Signaling Pathway and Renal Inflammation through the NF-kB Signaling Pathway. J AGR FOOD CHEM. 2022;XXXX(XXX):XXX-XXX WB ;Mouse, Human. 36525382