bs-0200R

[Primary Antibody]

PACAP-38 Rabbit pAb



www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn

		400-901-9800	
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)	
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)	
GenelD: 116	SWISS: P18509	IF (1:100-500)	
Target: PACAP-38 Immunogen: KLH conjugated synthetic peptide derived from human PACAP-38: 31-38/38 (162-139/176).		Reactivity: Human, Mouse, Rat (predicted: Pig, Sheep, Cow)	
Purification: affinity purified by	Protein A		
Concentration: 1mg/ml		Predicted MW.: ^{5 kDa}	
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.		Subcellular Secreted ,Extracellular Location: matrix	
Background: This gene encodes into multiple matu cyclase and increa levels, resulting in The products of th stress responses.	a secreted proprotein that is further processed ire peptides. These peptides stimulate adenylate se cyclic adenosine monophosphate (cAMP) the transcriptional activation of target genes. is gene are key mediators of neuroendocrine Alternative splicing results in multiple transcript		

- VALIDATION IMAGES



variants. [provided by RefSeq, Feb 2013]

Sample: Lane 1: Mouse Pituitary Lysates Primary: Anti-PACAP-38 (bs-0200R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 5kDa Observed band size: 19kDa



Tissue/cell: human kidney tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-PACPA Polyclonal Antibody, Unconjugated(bs-0200R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-PACAP-38 Polyclonal Antibody, Unconjugated(bs-0200R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: mouse small intestine tissue;4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-PACAP-38 Polyclonal Antibody, Unconjugated(bs-0200R)

1:200, overnight at 4°C; The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C. DAPI(5ug/ml,blue,C-0033) was used to stain the cell nuclei

- SELECTED CITATIONS -

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- [IF=8.039] Yifan Zhu. et al. Discovery of Selective P2Y6R Antagonists with High Affinity and In Vivo Efficacy for Inflammatory Disease Therapy. J MED CHEM. 2023;XXXX(XXX):XXX-XXX WB ;MOUSE. 37078976
- [IF=0.7] Ming XU. et al. Underlying mechnism of electroacupuncture for treating detrusor hyperreflexia after suprasacral spinal cord injury through the PACAP-cAMP signaling pathway: 基于PACAP-cAMP. WORLD J ACUPUNCT-MOX. 2023 Jun;: IHC,WB;Rat. 10.1016/j.wjam.2023.06.002
- [IF=0] PENG Y et al. Effect of electroacupuncture on urodynamics, PACAP-38 and PAC1R of spinal cord in rats with neurogenic bladder ☆. World Journal of Acupuncture Moxibustion, 2018 28(1), 50–54. WB ;Rat. 10.1016/j.wjam.2018.03.014