bs-0175R

[Primary Antibody]

TrkB Rabbit pAb



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— DATASHEET ———		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500)
GenelD: 4915	SWISS: Q16620	IF (1:100-500)
Target: TrkB		Flow-Cyt (1µg/Test)
Immunogen: KLH conjugated synthetic peptide derived from human NTRK2: 401-500/822.		K2: Reactivity: Human, Rat (predicted: Mouse)
Purification: affinity purified by	Protein A	
Concentration: 1mg/ml		Predicted
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.		MW.: ^{90-92 kDa}
freeze/thaw cycles		Location: Cell membrane, Cytoplasm
Background: This gene encodes kinase (NTRK) fam that, upon neurotr members of the M	a member of the neurotrophic tyrosine re ily. This kinase is a membrane-bound rece ophin binding, phosphorylates itself and APK pathway. Signalling through this kina.	eceptor eptor se

leads to cell differentiation. Mutations in this gene have been associated with obesity and mood disorders. Alternate transcriptional splice variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008].

- VALIDATION IMAGES



Sample: Lane 1: SH-SY5Y (Human) Cell Lysate at 30 ug Lane 2: U251 (Human) Cell Lysate at 30 ug Lane 3: U87MG (Human) Cell Lysate at 30 ug Primary: Anti-TrkB (bs-0175R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 130 kD Observed band size: 130 kD



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 (TrkB) Polyclonal Antibody, Unconjugated (bs-0175R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Tissue/cell: human nasopharyngeal carcinoma; 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-Trk-B Polyclonal Antibody, Unconjugated(bs-0175R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control: RSC96(blue), the cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with ice-cold 90% methanol for 30 min on ice. Isotype Control Antibody: Rabbit IgG(orange) ; Secondary Antibody: Goat antirabbit IgG-PE(white blue), Dilution: 1:200 in 1 X

- SELECTED CITATIONS -

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- **[IF=4.362]** Zhang Yan-hui. et al. α-Lipoic Acid Maintains Brain Glucose Metabolism via BDNF/TrkB/HIF-1α Signaling Pathway in P301S Mice. Front Aging Neurosci. 2020 Aug;12:262 IHC ;MOUSE. 32973490
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