bs-4995R

[Primary Antibody]

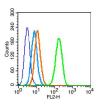
Insulin receptor subunit beta Rabbit pAb



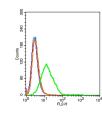
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Clonality: Polyclonal GenelD: 3643 SWISS: P06213	Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (0.2µg/Test) ICC/IF (1:100) Reactivity: Human (predicted: Mouse,
GenelD: 3643 SWISS: P06213	IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (0.2µg/Test) ICC/IF (1:100)
	ICC/IF (1:100)
 Target: Insulin receptor subunit beta Immunogen: KLH conjugated synthetic peptide derived from human Insulin receptor subunit beta: 801-900/1382. < Extracellular > Purification: affinity purified by Protein A Concentration: 1mg/ml 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. Background: The human insulin receptor is a heterotetrameric membrane glycoprotein consisting of disulfide linked subunits in a beta-alpha-alpha-beta configuration. The beta subunit (95 kDa) possesses a single transmembrane domain, whereas the alpha subunit (135 kDa) is completely extracellular. The insulin receptor exhibits receptor tyrosine kinase (RTK) activity. RTKs are single pass transmembrane receptors that possess intrinsic cytoplasmic enzymatic activity, catalyzing the transfer of the gamma phosphate of ATP to tyrosine residues in protein substrates. RTKs are essential components of signal transduction pathways that affect cell proliferation, differentiation, migration and metabolism. Included in this large protein family are the insulin receptor and the receptors drow factors such as epidermal growth factor. Receptor activation occurs through ligand binding, which facilitates receptor dimerization and autophosphorylation of specific tyrosine residues in the cytoplasmic portion. The interaction of insulin with the alpha subunit of the insulin receptor activates the protein tyrosine kinase of the beta subunit, which then undergoes an autophosphorylation that increases its tyrosine kinase activity. Three adapter proteins, IRS1, IRS2 and Sbc, become phosphorylated on tyrosine residues following insulin receptor activation. These three phosphorylated proteins then 	Rat) Predicted MW.: 68/152 kDa Subcellular Location: Cell membrane
interact with SH2 domain containing signaling proteins.	

— VALIDATION IMAGES



Blank control (blue line): HL60(fixed with 70% ethanol Overnight at 4°C). Primary Antibody (green line): Rabbit Anti-Insulin receptor subunit beta antibody (bs-4995R),Dilution: 0.2µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-PE,Dilution: 1µg /test.



Blank control: RSC96(blue). Primary Antibody:Rabbit Anti-Insulin receptor subunit beta antibody(bs-4995R), Dilution: 5µg in 100 1µL 1X PBS containing 0.5% BSA; Isotype Control Antibody: Rabbit IgG(orange) ,used under the same conditions); Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA.

Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

- SELECTED CITATIONS -

• [IF=2.369] Martinez-Rachadell L et al. Cell-specific expression of insulin/insulin-like growth factor-I receptor hybrids in the mouse brain. Growth Horm. IGF Res. 2019 Feb 22;45 IHC ;MOUSE. 30825797