[Primary Antibody]

phospho-IRS-2(Ser731) Rabbit pAb

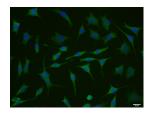


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- DATASHEE	Т ———		4(
	Rabbit	lsotype: IgG	Applicati
Clonality:	Polyclonal		
GenelD:	8660	SWISS: Q9Y4H2	Reactiv
Target:	IRS-2(Ser731)		
		sed phosphopeptide derived from huma orylation site of Ser731: AS(p-S)PA.	n Predie
Purification:	affinity purified by Protei	n A	M
Concentration:	1mg/ml		Subcell
	Glycerol.	% BSA, 0.02% Proclin300 and 50% 20°C for one year. Avoid repeated	Locat
	to play important roles for hormones. Four member Each IRS is believed to ha distinct physiological role gene encodes the insulin signaling molecule that n growth factor 1, and othe adaptor between diverse downstream effectors. Th	ptor substrates (IRSs) has been reported or signal transduction of various s of the IRS family have been described. we different functions; however, the es of each IRS are unclear. Summary: This receptor substrate 2, a cytoplasmic nediates effects of insulin, insulin-like er cytokines by acting as a molecular receptor tyrosine kinases and ne product of this gene is phosphorylated rosine kinase upon receptor stimulation,	s

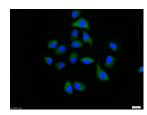
as well as by an interleukin 4 receptor-associated kinase in

- VALIDATION IMAGES

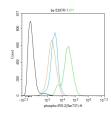


response to IL4 treatment.

SH-SY5Y cell; 4% Paraformaldehyde-fixed; 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 (phospho-IRS-2(Ser731)) polyclonal Antibody, Unconjugated (bs-5397R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Hela cell; 4% Paraformaldehyde-fixed; 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 (phospho-IRS-2(Ser731)) polyclonal Antibody, Unconjugated (bs-5397R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control (black line) :SH-SY5Y. Primary Antibody (green line): Rabbit Anti-phospho-IRS-2(Ser731) antibody (bs-5397R) Dilution:1ug/Test; Secondary Antibody (white blue line) : Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line) : Normal Rabbit IgG 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.

pplications: Flow-Cyt (1ug/Test) ICC/IF (1:100)

Reactivity: Human (predicted: Mouse, Rat, Dog, Horse)

Predicted MW.: 147 kDa

Subcellular Location: Cytoplasm

- SELECTED CITATIONS -

• [IF=7.376] Wei Yu. et al. Silencing TXNIP ameliorates high uric acid-induced insulin resistance via the IRS2/AKT and Nrf2/HO-1 pathways in macrophages. Free Radical Bio Med. 2022 Jan;178:42 WB ;Human. 34848368