

Insulin Receptor β (phospho-Tyr1345) rabbit pAb

Cat No.:ES15451

For research use only

Overview

Product Name	Insulin Receptor β (phospho-Tyr1345) rabbit pAb
Host species	Rabbit
Applications	WB
Species Cross-Reactivity	Human;Rat;Mouse;
Recommended dilutions	WB 1:1000-2000
Immunogen	Synthesized phosho peptide around human Insulin
	Receptor β (Tyr1345)
Specificity	This antibody detects endogenous levels of Human
	Insulin Receptor β (phospho-Tyr1345)
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and
	0.02% sodium azide.
Storage	Store at -20 $^\circ \!\! \mathbb{C}$. Avoid repeated freeze-thaw cycles.
Protein Name	Insulin Receptor β (Tyr1345)
Gene Name	INSR
Cellular localization	Cell membrane ; Single-pass type I membrane
	protein . Late endosome . Lysosome . Binding of
	insulin to INSR induces internalization and lysosomal
	degradation of the receptor, a means for
	down-regulating this signaling pathway after
	stimulation. In the presence of SORL1, internalized
	INSR molecules are redirected back to the cell
	surface, thereby preventing their lysosomal
	catabolism and strengthening insulin signal
	reception
Purification	The antibody was affinity-purified from rabbit
	antiserum by affinity-chromatography using
	epitope-specific immunogen.
Clonality	Polyclonal
Concentration	1 mg/ml
Observed band	95kD
Human Gene ID	3643
Human Swiss-Prot Number	P06213
+86-27-59760950	ELKbio@ELKbiotech.com www.elkbiotech.com



23-2, No.388 Gaoxin 2nd Road, Wuhan East Lake Hi-tech Development Zone, Hubei , P.R.C



Alternative Names

Background

Insulin receptor (IR) (EC 2.7.10.1) (CD antigen CD220) [Cleaved into: Insulin receptor subunit alpha; Insulin receptor subunit beta] This gene encodes a member of the receptor tyrosine kinase family of proteins. The encoded preproprotein is proteolytically processed to generate alpha and beta subunits that form a heterotetrameric receptor. Binding of insulin or other ligands to this receptor activates the insulin signaling pathway, which regulates glucose uptake and release, as well as the synthesis and storage of carbohydrates, lipids and protein. Mutations in this gene underlie the inherited severe insulin resistance syndromes including type A insulin resistance syndrome, Donohue syndrome and Rabson-Mendenhall syndrome. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2015],



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ELKbio@ELKbiotech.com

www.elkbiotech.com

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