











INSR Antibody

| Product Code | CSB-RA155156A0HU |
|---------------------|--|
| Storage | Upon receipt, store at -20°C or -80°C. Avoid repeated freeze. |
| Uniprot No. | P06213 |
| Immunogen | A synthesized peptide derived from human Insulin Receptor |
| Species Reactivity | Human |
| Tested Applications | ELISA, IF; Recommended dilution: IF:1:20-1:200 |
| Relevance | Receptor tyrosine kinase which mediates the pleiotropic actions of insulin. Binding of insulin leads to phosphorylation of several intracellular substrates, including, insulin receptor substrates (IRS1, 2, 3, 4), SHC, GAB1, CBL and other signaling intermediates. Each of these phosphorylated proteins serve as docking proteins for other signaling proteins that contain Src-homology-2 domains (SH2 domain) that specifically recognize different phosphotyrosine residues, including the p85 regulatory subunit of PI3K and SHP2. Phosphorylation of IRSs proteins lead to the activation of two main signaling pathways: the PI3K-AKT/PKB pathway, which is responsible for most of the metabolic actions of insulin, and the Ras-MAPK pathway, which regulates expression of some genes and cooperates with the PI3K pathway to control cell growth and differentiation. Binding of the SH2 domains of PI3K to phosphotyrosines on IRS1 leads to the activation of PI3K and the generation of phosphatidylinositol-(3, 4, 5)-triphosphate (PIP3), a lipid second messenger, which activates several PIP3-dependent serine/threonine kinases, such as PDPK1 and subsequently AKT/PKB. The net effect of this pathway is to produce a translocation of the glucose transporter SLC2A4/GLUT4 from cytoplasmic vesicles to the cell membrane to facilitate glucose transport. Moreover, upon insulin stimulation, activated AKT/PKB is responsible for: anti-apoptotic effect of insulin by inducing phosphorylation of BAD; regulates the expression of gluconeogenic and lipogenic enzymes by controlling the activity of the winged helix or forkhead (FOX) class of transcription factors. Another pathway regulated by PI3K-AKT/PKB activation is mTORC1 signaling pathway which regulates cell growth and metabolism and integrates signals from insulin. AKT mediates insulin-stimulated protein synthesis by phosphorylating TSC2 thereby activating mTORC1 pathway. The Ras/RAF/MAPZK/MAPK pathway is mainly involved in mediating cell growth, survival and cellular differentiation of insulin, the insulin |

IGF2 and insulin. In contrast, PubMed:16831875 shows that hybrid receptors composed of IGF1R and INSR isoform Long and hybrid receptors composed of IGF1R and INSR isoform Short have similar binding characteristics, both bind

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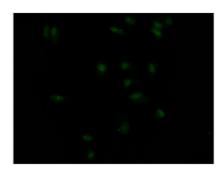




| IGF1 | and have | a low | affinity | for | insulin. |
|------|----------|-------|----------|-----|----------|
|------|----------|-------|----------|-----|----------|

| Form | Liquid |
|----------------------------|---|
| Conjugate | Non-conjugated |
| Storage Buffer | Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. |
| Purification Method | Affinity-chromatography |
| Isotype | Rabbit IgG |
| Clonality | Monoclonal |
| Product Type | Recombinant Antibody |
| Immunogen Species | Homo sapiens (Human) |
| Research Area | Neuroscience; Cancer; Cardiovascular; Metabolism; Signal transduction |
| Gene Names | INSR |
| Accession NO. | 4B2 |
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Image



Immunofluorescence staining of MCF7 Cells with CSB-RA155156A0HU at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG

Description

After binding to insulin, a conformational change is triggered that enables INSR to bind ATP and activate INSR tyrosine kinases. Insulin receptor substrate proteins (IRS) are then phosphorylated as a critical node, leading to the activation of downstream pathways of two other critical nodes, phosphatidylinositol 3-kinase (PI3K) and AKT/PKB, which control the metabolic action of insulin and regulate the expression of genes that mediate cell growth and differentiation. The major effects of insulin include stimulation of glucose utilization and synthesis of glycogen, lipids, and protein.

The main production processes of this recombinant INSR antibody included immunization, B cell harvest, antibody secreting cell enrichment, single cell sequencing, antibody expression and purification. The single B cell screening platform was used for the INSR antibody gene screening. And this INSR antibody was tested in ELISA, IF.