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PhosphoPlus[®] IGF-I Receptor β Antibody



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UniProt ID: Entrez-Gene Id: #P06213, #P08069 3643, 3480

Product Includes	Product #	Quantity	Mol. Wt	Isotype/Source
Phospho-IGF-I Receptor β (Tyr1135) (DA7A8) Rabbit mAb	3918	100 µl	95 kDa	Rabbit IgG
IGF-I Receptor β (D23H3) XP [®] Rabbit mAb	9750	100 µl	95 kDa	Rabbit IgG

Please visit cellsignal.com for individual component applications, species cross-reactivity, dilutions, protocols, and additional product information.

Description

PhosphoPlus[®] Duets from Cell Signaling Technology (CST) provide a means to assess protein activation status. Each Duet contains an activation-state and total protein antibody to your target of interest. These antibodies have been selected from CST's product offering based upon superior performance in specified applications.

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, $100 \mu g/ml$ BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

Background

Type I insulin-like growth factor receptor (IGF-IR) is a transmembrane receptor tyrosine kinase that is widely expressed in many cell lines and cell types within fetal and postnatal tissues (1-3). Receptor autophosphorylation follows binding of the IGF-I and IGF-II ligands. Three tyrosine residues within the kinase domain (Tyr1131, Tyr1135, and Tyr1136) are the earliest major autophosphorylation sites (4). Phosphorylation of these three tyrosine residues is necessary for kinase activation (5,6). Insulin receptors (IRs) share significant structural and functional similarity with IGF-I receptors, including the presence of an equivalent tyrosine cluster (Tyr1146/1150/1151) within the kinase domain activation loop. Tyrosine autophosphorylation of IRs is one of the earliest cellular responses to insulin stimulation (7). Autophosphorylation begins with phosphorylation at Tyr1146 and either Tyr1150 or Tyr1151, while full kinase activation requires triple tyrosine phosphorylation (8).

Background References

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- 2. Baserga, R. (2000) Oncogene 19, 5574-81.
- 3. Scheidegger, K.J. et al. (2000) J Biol Chem 275, 38921-8.
- 4. Hernández-Sánchez, C. et al. (1995) *J Biol Chem* 270, 29176-81.
- 5. Lopaczynski, W. et al. (2000) Biochem Biophys Res Commun 279, 955-60.
- 6. Baserga, R. (1999) Exp Cell Res 253, 1-6.
- 7. White, M.F. et al. (1985) J Biol Chem 260, 9470-8.
- 8. White, M.F. et al. (1988) J Biol Chem 263, 2969-80.

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