

**IGF-II Receptor/CI-M6PR (D3V8C) Rabbit mAb**

**Orders:** 877-616-CELL (2355)  
orders@cellsignal.com

**Support:** 877-678-TECH (8324)

**Web:** info@cellsignal.com  
cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

**For Research Use Only. Not for Use in Diagnostic Procedures.**

<b>Applications:</b> W, IP, IF-IC	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 275	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P11717	<b>Entrez-Gene Id:</b> 3482
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**Product Usage Information****Application**

Western Blotting  
Immunoprecipitation  
Immunofluorescence (Immunocytochemistry)

**Dilution**

1:1000  
1:50  
1:400

**Storage**

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

**Specificity/Sensitivity**

IGF-II Receptor/CI-M6PR (D3V8C) Rabbit mAb recognizes endogenous levels of total IGF-II Receptor/CI-M6PR protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala1675 of human IGF-II Receptor/CI-M6PR protein.

**Background**

Insulin-like growth factor II (IGF-II) receptor, also widely known as cation-independent mannose 6-phosphate receptor (CI-M6PR), is a multifunctional type I transmembrane glycoprotein that participates in the internalization of mannose-6-phosphate modified hydrolases and IGF-II from the plasma membrane (1,2). In the absence of ligands, IGF-II receptor is constitutively endocytosed from the cell surface to accumulate in the Golgi apparatus (3). In the presence of ligands, the receptor transports the mannose-6-phosphate modified hydrolases to acidified endosomes and lysosomes (4). The ligand-free receptor is then transported back to the Golgi compartment or the cell surface (4). In several research studies, IGF-II receptor has been recognized as a tumor suppressor in a number of cancers (5-7).

**Background References**

1. Lobel, P. et al. (1989) *Cell* 57, 787-96.
2. Kiess, W. et al. (1988) *J Biol Chem* 263, 9339-44.
3. York, S.J. et al. (1999) *J Biol Chem* 274, 1164-71.
4. Duncan, J.R. and Kornfeld, S. (1988) *J Cell Biol* 106, 617-28.
5. Oates, A.J. et al. (1998) *Breast Cancer Res Treat* 47, 269-81.
6. Martin-Kleiner, I. and Gall Troselj, K. (2010) *Cancer Lett* 289, 11-22.
7. Puxbaum, V. et al. (2012) *J Hepatol* 57, 337-43.

**Species Reactivity**

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

**Western Blot Buffer**

**IMPORTANT:** For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key**

**W:** Western Blotting **IP:** Immunoprecipitation **IF-IC:** Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key**

**H:** Human **M:** Mouse **R:** Rat **Mk:** Monkey

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