PDGF Receptor β (2B3) Mouse mAb



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Applications: W, IP	Reactivity: M R	Sensitivity: Endogenous	MW (kDa): 190	Source/Isotype: Mouse IgG1	UniProt ID: #P09619	Entrez-Gene Id: 5159
Product Usage Information		Application Western Blotting Immunoprecipitation		Dilution 1:1000 1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		PDGF Receptor β (2B3) Mouse mAb detects endogenous levels of total PDGF receptor β protein. The antibody does not cross-react with PDGF receptor α .				
Species predicted to react based on 100% sequence homology		Human				
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant human PDGF receptor $\boldsymbol{\beta}$ protein.				
Background		Platelet derived growth factor (PDGF) family proteins exist as several disulphide-bonded, dimeric isoforms (PDGF AA, PDGF AB, PDGF BB, PDGF CC, and PDGF DD) that bind in a specific pattern to two closely related receptor tyrosine kinases, PDGF receptor α (PDGFRα) and PDGF receptor β (PDGFRβ). PDGFRα and PDGFRβ share 75% to 85% sequence homology between their two intracellular kinase domains, while the kinase insert and carboxy-terminal tail regions display a lower level (27% to 28%) of homology (1). PDGFRα homodimers bind all PDGF isoforms except those containing PDGF D. PDGFRβ homodimers bind PDGF BB and DD isoforms, as well as the PDGF AB heterodimer. The heteromeric PDGF receptor α/β binds PDGF B, C, and D homodimers, as well as the PDGF AB heterodimer (2). PDGFRα and PDGFRβ can each form heterodimers with EGFR, which is also activated by PDGF (3). Various cells differ in the total number of receptors present and in the receptor subunit composition, which may account for responsive differences among cell types to PDGF binding (4). Ligand binding induces receptor dimerization and autophosphorylation, followed by binding and activation of cytoplasmic SH2 domain-containing signal transduction molecules, such as GRB2, Src, GAP, PI3 kinase, PLCγ, and NCK. A number of different signaling pathways are initiated by activated PDGF receptors and lead to control of cell growth, actin reorganization, migration, and differentiation (5). Tyr751 in the kinase-insert region of PDGFRβ is the docking site for PI3 kinase (6). Phosphorylated pentapeptides derived from Tyr751 of PDGFRβ (pTyr751-Val-Pro-Met-Leu) inhibit the association of the carboxy-terminal SH2 domain of the p85 subunit of PI3 kinase with PDGFRβ (7). Tyr740 is also required for PDGFRβ-mediated PI3 kinase activation (8).				
Background References		2. Bergsten, E. et al. (2 3. Betsholtz, C. et al. (4. Coughlin, S.R. et al. 5. Ostman, A. and Hel 6. Panayotou, G. et al 7. Ramalingam, K. et al	1988) Biofactors 1, 213-217. (2001) Nat. Cell Biol. 3, 512-516. . (2001) Bioessays 23, 494-507. al. (1988) Prog. Clin. Biol. Res. 266, 39-45. leldin, C.H. (2001) Adv. Cancer Res. 80, 1-38. al. (1992) EMBO J. 11, 4261-4272. tt al. (1995) Bioorg. Med. Chem. 3, 1263-1272. al. (1992) EMBO J. 11, 1373-1382.			

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

Cross-Reactivity Key M: Mouse R: Rat

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