

## Technical Data Sheet

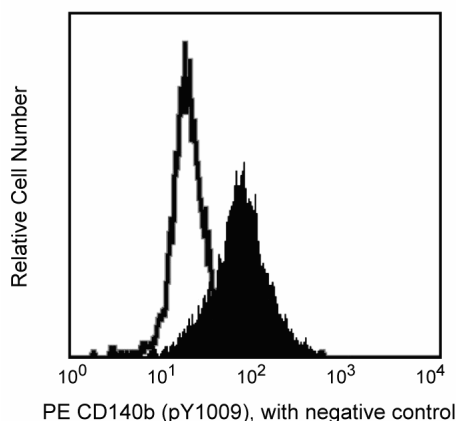
**PE Mouse anti-PDGFR $\beta$  (CD140b) (pY1009)****Product Information**

<b>Material Number:</b>	558322
<b>Alternate Name:</b>	CD140b (pY1009)
<b>Size:</b>	50 tests
<b>Vol. per Test:</b>	20 $\mu$ l
<b>Clone:</b>	J25-602
<b>Immunogen:</b>	Phosphorylated Human PDGFR $\beta$ (pY1009) Peptide
<b>Isotype:</b>	Mouse IgG2b, $\kappa$
<b>Reactivity:</b>	QC Testing: Mouse Confirmed by immunohistochemistry using purified, unconjugated antibody: Human
<b>Storage Buffer:</b>	Aqueous buffered solution containing BSA and $\leq 0.09\%$ sodium azide.

**Description**

Platelet-derived growth factor (PDGF) is a potent mitogen for cells of mesenchymal origin and exerts its effects by binding to the PDGF receptor (PDGFR), a transmembrane protein tyrosine kinase. PDGFR is composed of PDGFR $\alpha$  (CD140a) and/or PDGFR $\beta$  (CD140b) polypeptides. Both PDGF and PDGFR consist of subunits that form homo- or heterodimers with varying specificities: PDGF-AA binds only to  $\alpha\alpha$  PDGFR, PDGF-AB binds to both  $\alpha\alpha$  and  $\alpha\beta$  PDGFR, and PDGF-BB binds to all three PDGFRs. Ligand binding induces dimerization and activation of the receptor. Upon activation, CD140b is phosphorylated at multiple tyrosine sites and, in turn, an intracellular phosphorylation cascade is initiated. PDGFR localizes primarily to membrane invaginations termed caveolae, compartments that are enriched in several of its downstream effectors, including phosphatidylinositol 3'-kinase, Src, and phospholipase C- $\gamma$  (PLC- $\gamma$ ).

The J25-602 monoclonal antibody recognizes the phosphorylated tyrosine 1009 (pY1009) in the C-terminal noncatalytic region of CD140b, which interacts primarily with protein tyrosine phosphatase 1D and possibly with PLC- $\gamma$ . The orthologous phosphorylation site in mouse PDGFR $\beta$  is Y1008.



**Analysis of CD140b (pY1009) in activated mouse embryonic fibroblasts.** NIH/3T3 cells (ATCC CRL-1658) were serum starved overnight and either stimulated with Platelet-Derived Growth Factor-BB (Cat. No. 354051) at 37°C for 2 minutes (filled histogram) or unstimulated (open histogram). The cells were fixed with BD Phosflow™ Fix Buffer I (Cat. No. 557870) at 37°C for 10 minutes, permeabilized in BD Phosflow™ Perm Buffer III (Cat. No. 558050) on ice for at least 30 minutes, and stained with PE anti-CD140b (pY1009). Flow cytometry was performed on a BD FACSCalibur™ flow cytometry system.

**Preparation and Storage**

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated with R-PE under optimum conditions, and unconjugated antibody and free PE were removed.

**Application Notes****Application**

Intracellular staining (flow cytometry)	Routinely Tested
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## Suggested Companion Products

Catalog Number	Name	Size	Clone
557870	Fix Buffer I	250 ml	(none)
558050	Perm Buffer III	125 ml	(none)
554656	Stain Buffer (FBS)	500 ml	(none)

## Product Notices

1. This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use  $1 \times 10^6$  cells in a 100- $\mu$ l experimental sample (a test).
2. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at [www.bdbiosciences.com/colors](http://www.bdbiosciences.com/colors).
3. Please refer to [www.bdbiosciences.com/pharmlingen/protocols](http://www.bdbiosciences.com/pharmlingen/protocols) for technical protocols.
4. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
5. Source of all serum proteins is from USDA inspected abattoirs located in the United States.

## References

Claesson-Welsh L. Platelet-derived growth factor receptor signals. *J Biol Chem*. 1994; 269(51):32023-32026. (Biology)  
Liu J, Oh P, Horner T, Rogers RA, Schnitzer JE. Organized endothelial cell surface signal transduction in caveolae distinct from glycosylphosphatidylinositol-anchored protein microdomains. *J Biol Chem*. 1997; 272(11):7211-7222. (Biology)

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