SONY

Product Data Sheet

PerCP/Cyanine5.5 anti-mouse CD140a

Catalog # / Size:	1279570 / 100 µg 1279565 / 25 µg
Clone:	APA5
Isotype:	Rat IgG2a, κ
Reactivity:	Mouse
Concentration:	0.2

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤1.5 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application. * PerCP/Cyanine5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm.
Application Notes:	Additional reported (for relevant formats) applications include: Western Blot, blocking function2, and immunohistochemical staining of paraffin and frozen sections. The LEAF™ purified antibody is recommended for functional assays.
Application References:	1. Mukouyama YS, *et al.* 2006. *Proc Natl Acad Sci USA.* 103(5):1551 2. Miyawaki T, *et al.* 2004. *J Neurosci.* 24(37):8124 3. Takakura N, *et al.* 1997. *J Histochem Cytochem.* 45(6):883

Description:	Platelet-derived growth factor receptor-α (PDGFR-α), CD140a, is one of two receptors for platelet-derived growth factors (PDGFs) and binds to all isoforms of PDGFs: PDGF-AA, PDGF-AB, and PDGF-BB. PDGFRa is a receptor tyrosine kinase that forms homodimers or heterodimers on the surface upon ligand binding and phosphorylates substrates. PDGFRs consist of either homodimers of α/α, β/β, or heterodimers of α/β. PDGF receptors, α and β, are single glycoproteins with intracellular tyrosine kinase domain. Their ligand, PDGF, is a mitogen for connective tissue and glial cells. CD140a is expressed on embryonic tissues and mesenchymal-derived cells of adult mice. PDGF plays a role in wound healing and acts as a chemoattractant for fibroblasts, smooth muscle cells, glial cells, monocytes, and neutrophils.
Antigen References:	1. Mukouyama YS, *et al.* 2006. *Proc Natl Acad Sci USA.* 103(5):1551 2. Miyawaki T, *et al.* 2004. *J Neurosci.* 24(37):8124 3. Takakura N, *et al.* 1997. *J Histochem Cytochem.* 45(6):883

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