

**PerCP/Cyanine5.5 anti-human CD49b**

**Catalog # /** 2396575 / 25 tests  
**Size:** 2396580 / 100 tests

**Clone:** P1E6-C5

**Isotype:** Mouse IgG1,  $\kappa$

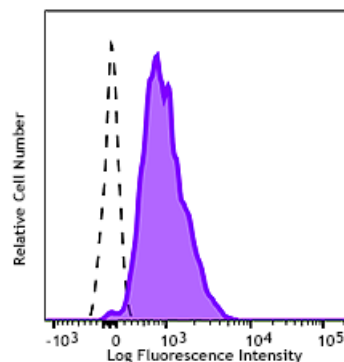
**Immunogen:** HT1080 cells

**Reactivity:** Human

**Preparation:** The antibody was purified by affinity chromatography and conjugated with PerCP/Cyanine5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cyanine5.5 and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).

**Concentration:** Lot-specific



Human peripheral blood platelets were stained with CD49b (clone P1E6-C5) PerCP/Cyanine5.5 (filled histogram) or mouse IgG1,  $\kappa$  PerCP/Cyanine5.5 isotype control (open histogram).

**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 5  $\mu$ L per million cells in 100  $\mu$ L staining volume or 5  $\mu$ L per 100  $\mu$ L of whole blood.

\* PerCP/Cyanine5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm.

**Application Notes:** Additional reported applications (for the relevant formats of this clone) include: *in vitro* blocking activity<sup>1,2</sup>, immunoprecipitation<sup>3</sup>, and ELISA<sup>4</sup>.

**Application References:**

1. Hirsch MS, *et al.* 1997. *Dev. Dyn.* 210:249. (Block)
2. Sawhney RS, *et al.* 2006. *J. Biol. Chem.* 281:8497. (Block)
3. Lee SA, *et al.* 2009. *Carcinogenesis*. 30:1872. (IP)
4. Zbrate S, *et al.* 2004. *J. Virol.* 78:10839. (ELISA)

**Description:** CD49b is a 170 kD transmembrane protein, also known as  $\alpha_2$  integrin, VLA-2  $\alpha$  chain, Integrin  $\alpha_2$  and GPIa. It associates with CD29 ( $\beta_1$  integrin) to form VLA-2, a collagen and laminin receptor on many cell types including monocytes, platelets, activated T cells, megakaryocytes, neuronal cells, epithelial cells, and osteoclasts. CD49b has been reported to interact with F-actin and matrix metalloproteinase 1. CD49b is a platelet alloantigen and has been associated with neonatal alloimmune thrombocytopenia. Deficiencies in this protein have been associated with hemorrhagic disorders.

**Antigen References:**

1. Kaplan C, *et al.* 1991. *Br. J. Haematol.* 78:425.
2. Kiefel V, *et al.* 1991. *Vox Sang.* 60:244.
3. Nieuwenhuis HK, *et al.* 1985. *Nature* 318:470.
4. Takada Y and Helmer ME. 1989. *J. Cell Biol.* 109:397.