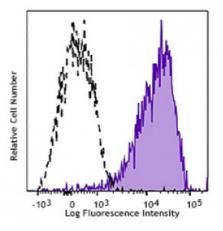
Product Data Sheet

PE/Cy7 anti-human CD11a/CD18 (LFA-1)

Catalog # / Size:	2417090 / 100 tests 2417085 / 25 tests
Clone:	m24
Isotype:	Mouse IgG1, κ
Immunogen:	Fibronectin-purified human monocytes
Reactivity:	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE/Cy7 under optimal conditions. The solution is free of unconjugated PE/Cy7 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



PMA-stimulated human peripheral blood granulocytes were stained with CD11/CD18 (clone m24) PE/Cy7 (filled histogram) or mouse IgG1, κ PE/Cy7 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 μ l per million cells or 5 μ l per 100 μ l of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.
Application Notes:	Clone m24 can be used as a reporter of the activation state of integrin receptor in response to exposure with Mg^{2+} or Mn^{2+} .
Application References:	1. Anderson D, <i>et al.</i> 1987. <i>Annu. Rev. Med.</i> 38:175. 2. Springer T. 1994. <i>Cell</i> 76:301.
Description:	CD11/CD18 belongs to the integrin family of proteins. It is heterodimeric cell surface receptor expressed on all leukocytes. CD18, in association with integrin α chain CD11a, CD11b, and CD11c forms LFA-1, Mac-1, and $\alpha_X\beta_2$, respectively, and plays an important role in leukocyte adhesion. CD11/CD18 complexes bind ICAM-1 (CD54), ICAM-2 (CD102), ICAM-3 (CD50), iC3b, and fibrinogen. Clone m24 binds the extended/open high affinity conformation of CD11a/CD18. The antibody can be used as a reporter of the activation state of the integrin receptor in response to exposure to Mg ²⁺ or Mn ²⁺ .
Antigen References:	1. Anderson D, <i>et al.</i> 1987. <i>Annu. Rev. Med.</i> 38:175. 2. Springer T. 1994. <i>Cell</i> 76:301.

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