

30-1069F: FITC Conjugated Anti-CD102 / ICAM-2 Monoclonal Antibody (Clone:CBR-IC2/2)

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| Clonality : | Monoclonal |
| Clone Name : | CBR-IC2/2 |
| Application : | FACS |
| Reactivity : | Human |
| Conjugate : | FITC |
| Gene : | CD102 |
| Gene ID : | 3384 |
| Uniprot ID : | P13598 |
| Alternative Name : | ICAM2; CD102 |
| Isotype : | Mouse IgG2a |
| Immunogen Information : | Human CD102 cDNA transfected COS cells |

Description

CD102 / ICAM-2 (intracellular cell adhesion molecule-2), a counter receptor of LFA-1 (CD11a/CD18), is a transmembrane glycoprotein with two extracellular IgC-like domains and intracellular C-terminal tail. It is involved in lymphocyte recirculation and homing to the sites of inflammation. Through interaction with integrins it provides to the immune cells costimulatory signals. Expression of CD102 on blood cells (lymphocytes, monocytes, thrombocytes) is lower than on endothelium and follicular dendritic cells. CD102 levels are upregulated in lymph nodes with malignant infiltration.

Specificity: The mouse monoclonal antibody CBR-IC2/2 recognizes an extracellular epitope of CD102 (ICAM-2), an approximately 55 kDa type I transmembrane glycoprotein expressed mainly on vascular endothelial cells and follicular dendritic cells, in lower amount on lymphocytes, monocytes and platelets.

Product Info

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| Amount : | 100 tests |
| Purification : | Purified antibody is conjugated with fluorescein isothiocyanate (FITC) under optimum conditions and unconjugated antibody and free fluorochrome are removed by size-exclusion chromatography. |
| Content : | Formulation: Stabilizing phosphate buffered saline (PBS), pH 7.4, 15 mM sodium azide |
| Storage condition : | Store at 2-8°C. Protect from prolonged exposure to light. Do not freeze. |

Application Note

Flow cytometry: The reagent is designed for analysis of human blood cells using 4 µl reagent / 100 µl of whole blood or 10⁶ cells in a suspension. The content of a vial (0.4 ml) is sufficient for 100 tests.

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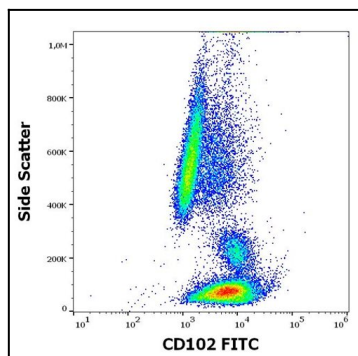


Figure 1: Flow cytometry surface staining pattern of human peripheral whole blood stained using anti-human CD102 (CBR-IC2/2) FITC antibody (4 μ l reagent / 100 μ l of peripheral whole blood).

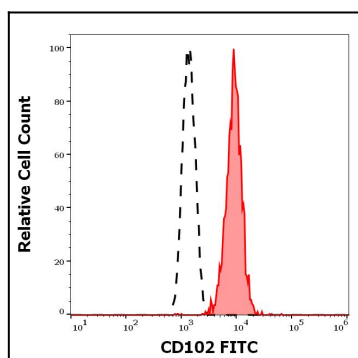


Figure 2: Separation of human monocytes (red-filled) from CD102 negative neutrophil granulocytes (black-dashed) in flow cytometry analysis (surface staining) of human peripheral whole blood stained using anti-human CD102 (CBR-IC2/2) FITC antibody (4 μ l reagent / 100 μ l of peripheral whole blood).