

### 30-1574PB: Pacific Blue Conjugated Anti-CD65 Monoclonal Antibody (Clone:VIM8)

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| <b>Clonality :</b>             | Monoclonal   |
| <b>Clone Name :</b>            | VIM8   |
| <b>Application :</b>           | FACS   |
| <b>Reactivity :</b>            | Human  |
| <b>Gene :</b>                  | CD65   |
| <b>Alternative Name :</b>      | ceramide-dodecasaccharide, type II fucoganglioside |
| <b>Isotype :</b>               | Mouse IgM  |
| <b>Immunogen Information :</b> | THP-1 cell line                                    |

#### Description

CD65 is a fucosylated carbohydrate antigen (ceramide-dodecasaccharide, type II fucoganglioside), which serves as a ligand for CD62E (E-selectin). Its structure is Gal beta1-4 GlcNAc beta1-3 Gal beta1-4 GlcNAc (3-1 Fuc alpha) beta1-3 ceramide. Unlike CD65s, the CD65 antigen does not contain terminal sialic acid, the rest of their structure is identical. CD65 is expressed on granulocytes and monocytes and participates in cell adhesion. It has been reported as important for extravascular infiltration of acute monocytic leukemia cells.

Specificity: The mouse monoclonal antibody VIM8 recognizes human CD65, an asialo-fucoganglioside expressed on the surface of peripheral blood granulocytes (highly) and monocytes (weakly).

#### Product Info

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|----------------------------|--|
| <b>Amount :</b>            | 100 tests  |
| <b>Purification :</b>      | Purified antibody is conjugated with Pacific Blue NHS ester under optimum conditions and unconjugated antibody and free fluorochrome are removed by size-exclusion chromatography. |
| <b>Content :</b>           | Stabilizing MES buffered saline, pH 5.5, 15 mM sodium azide  |
| <b>Storage condition :</b> | 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.

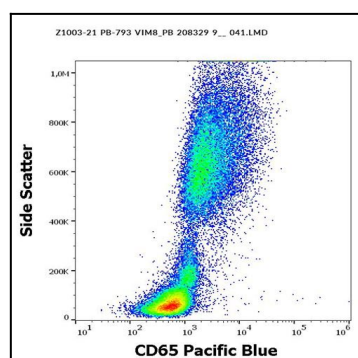


Figure 1: Flow cytometry surface staining pattern of human peripheral whole blood stained using anti-human CD65 (VIM8) Pacific Blue antibody (4 µl reagent / 100 µl of peripheral whole blood).

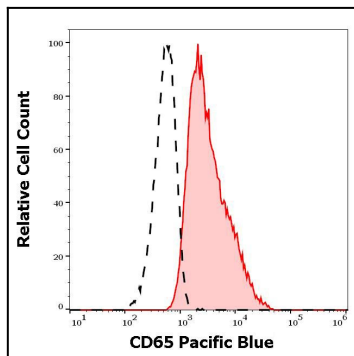


Figure 2: Separation of human neutrophil granulocytes (red-filled) from lymphocytes (black-dashed) in flow cytometry analysis (surface staining) of peripheral whole blood stained using anti-human CD65 (VIM8) Pacific Blue antibody (4 µl reagent / 100 µl of peripheral whole blood).