Alexa Fluor® 647 anti-human CD11c

Catalog # / Size: 2286150 / 100 tests

2286145 / 25 tests

Clone:

Isotype: Mouse IgG1, κ

Reactivity: Human

The antibody was purified by affinity **Preparation:**

chromatography and conjugated with Alexa Fluor® 647 under optimal

conditions.

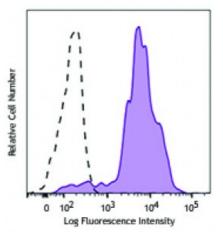
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

0.2% (w/v) BSA (origin USA).

Workshop Number: V S143

Concentration: Lot-specific



Human peripheral blood monocytes were stained with CD11c (clone Bu15) Alexa Fluor® 647 (filled histogram), or mouse IgG1, k Alexa Fluor® 647 isotype control (open histogram).

Applications:

Flow Cytometry **Applications:**

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 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

* Alexa Fluor® 647 has a maximum emission of 668 nm when it is excited at 633 nm / 635 nm.

Application Notes:

Clone Bu15 has a different binding epitope than clone 3.9. The binding of Bu15 with CD11c is divalent cation independent. Additional reported applications (for the relevant formats of this clone) include: inhibition of CD11c mediated adhesion and stimulation of chemokine production by monocytes.

Application References: 1. Sadhu C, et al. 2008. J. Immunoass. Immunoch. 29:42.

2. Rezzonico R. et al. 2001. Blood 97:2932.

3. Sadhu C, et al. 2007. J. Leukoc. Biol. 81:1395.

4. Yoshino N, et al. 2000. Exp. Anim. (Tokyo) 49:97. (FC)

Description: CD11c is a 145-150 kD type I transmembrane glycoprotein also known as integrin

> α_x and CR4. CD11c non-covalently associates with integrin β_2 (CD18) and is expressed on monocytes/macrophages, dendritic cells, granulocytes, NK cells, and subsets of T and B cells. CD11c has been reported to play a role in adhesion

and CTL killing through its interactions with fibrinogen, CD54, and iC3b.

Antigen References: 1. Petty H. 1996. Immunol. Today 17:209.

2. Springer T. 1994. Cell 76:301.

3. Ihanus E, et al. 2007. Blood 109:802-810.