Alexa Fluor® 700 anti-human CD18

Catalog # / 2110615 / 25 tests

Size: 2110620 / 100 tests

Clone: TS1/18

Isotype: Mouse IgG1, κ **Reactivity:** Human, Other

Preparation: The antibody was purified by affinity

chromatography and conjugated with Alexa Fluor® 700 under optimal conditions. The solution is free of unconjugated Alexa Fluor® 700.

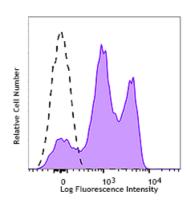
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

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

Workshop Number: V AS162

Concentration: Lot-specific



Human peripheral blood lymphocytes were stained with CD18 (clone TS1/18) Alexa Fluor® 700 (filled histogram) or Mouse lgG1, κ Alexa Fluor® 700 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.

* Alexa Fluor® 700 has a maximum emission of 719 nm when it is excited at 633 nm / 635 nm. Prior to using Alexa Fluor® 700 conjugate for flow cytometric analysis, please verify your flow cytometer's capability of exciting and detecting the fluorochrome.

Application

Additional reported applications (for the relevant formats) include:

Notes: inhibition of cell adhesion and migration^{3,4}.

Application

1. Anderson D, et al. 1987. Annu. Rev. Med. 38:175.

References: 2. Springer T. 1

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

Description:

CD18 is a 90-95 kD type I transmembrane protein also known as integrin β_2 subunit, LFA-1 β subunit, and β_2 integrin. CD18 non-covalently associates with CD11a, CD11b or CD11c. CD18 is expressed on all leukocytes. CD18 and associated α chains function in adhesion and signaling in

hematopoietic cells.

Antigen

1. Anderson D, et al. 1987. Annu. Rev. Med. 38:175.

References:

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