

FITC anti-mouse CD244.2 (2B4 B6 Alloantigen)

Catalog # / Size: 1267520 / 500 µg
1267515 / 50 µg

Clone: m2B4 (B6)458.1

Isotype: Mouse IgG1, κ

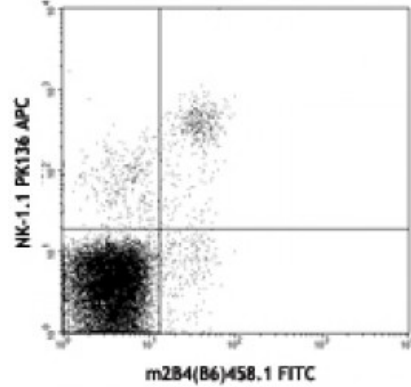
Immunogen: m2B4-Fc

Reactivity: Mouse

Preparation: The antibody was purified by affinity chromatography, and conjugated with FITC under optimal conditions. The solution is free of unconjugated FITC.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

Concentration: 0.5



C57BL/6 splenocytes stained with m2B4(B6)458.1 FITC and NK-1.1 APC

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 ≤ 0.25 microg per 10⁶ cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

Application Notes: The 2B4 antibody reacts with CD244.2, the 2B4 alloantigen which is expressed on C57BL/6 mice, but not on BALB/c mice.

Application References:

1. Lee KM, *et al.* 2006. *Blood*. 107:3181.
2. Schatzle JD, *et al.* 1999. *P. Natl Acad Sci USA*. 96:3870.
3. Stepp SE, *et al.* 1999. *Eur. J. Immunol.* 29:2392.
4. de la Luz Sierra M, *et al.* 2010. *Blood*. 115:3970. [PubMed](#)

Description: Mouse CD244, also known as 2B4, is a receptor belonging to the CD2 family of proteins involved in non- MHC-restricted cytotoxicity. It is expressed on all natural killer (NK) cells, IL-2 activated NK (LAK) cells, NKT cells, and a subset of T lymphocytes, including dendritic epidermal T cells. There are at least two isoforms of CD244.2 proteins that differ in the length of the cytoplasmic domain. The long form functions in an inhibitory manner, while the short form functions in an activating manner. The ligand of CD244 is CD48, which is expressed on all hematopoietic cells. It was reported that CD244 interaction with CD48 is essential for IL-2-driven expansion and activation of murine NK cells.

Antigen References:

1. Brown MH, *et al.* 1998. *J. Exp. Med.* 188:2083.
2. Davis SJ, *et al.* 1996. *Immunol Today*. 17:177.
3. Kumaresan PR, *et al.* 2000. *Immunogenetics*. 51:306.
4. Latchman Y, *et al.*