FITC Anti-Mouse CD278 Monoclonal Antibody

Catalog NumberVial SizeMHR12781-02B50 μgMHR12781-02E500 μg



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Important Note: Centrifuge before opening to ensure complete recovery of vial contents. This product is guaranteed up to one year from purchase.

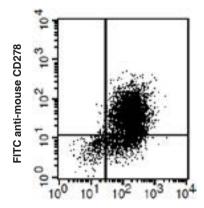
Purified Antibody Characterization

Clone	Isotype	Reactivity
17 G 9	Rat IgG2b	Mouse

Description

The 17G9 antibody reacts with the 47-57 kD ICOS protein, also known as inducible costimulatory molecule, and H4. This protein is homologous to the CD28/CTLA-4 proteins. ICOS is expressed on activated T cells and a subset of thymocytes and can costimulate T cells and induce proliferation. In addition ICOS has been shown to be involved in humoral immune responses (B cell germinal center formation). The ICOS ligand, B7h/B7RP-1 and B7-H2 is constitutively expressed in B cell areas of secondary lymphoid organs and can be induced in other tissues by LPS. ICOS stimulation has been shown to potentiate TCR-mediated IL-4 and IL-10 production and has been proposed to play a role in Th2 cell development. ICOS stimulation has been shown to be involved in airway tolerance and the downregulation of pulmonary inflammation.

Illustration of Immunofluorescent Staining



APC anti-mouse CD3

Con A-stimulated (3 days) C57BL/6 splenocytes stained with APC anti-mouse CD3 and FITC anti-mouse CD278

Product Information

Conjugation: FITC

Formulation: PBS pH 7.2, 0.09% NaN₃,

0.2% BSA

Concentration: 0.5 mg/ml

Storage: Keep as concentrated solution. Store at 4°C and protected from prolonged

exposure to light. Do not freeze.

Application: Recommended Application: FC

Usage: Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis (The amount of the reagent is suggested to be used ≤ 0.5 µg /10⁶ cells in 100 µl). Since applications vary, the appropriate dilutions must be determined for individual use.

References

- [1] Rudd CE, et al. 2003. Nat. Rev. Immunol. 3:544.
- [2] McAdam AJ, et al. 2000. J. Immunol. 165:5035.
- [3] Mak TW, et al. 2003. Nat. Immunol. 4:765.

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