

EuroBioSciences

Data Sheet

anti-rat CD4 FITC-conjugated

Cat-No.: R32126F 1 ml

Clone: W3/25

Specificity: This anti-rat CD4 monoclonal antibody recognizes a determinant on the majority of thymocytes (90-95%), a subset of peripheral T cells and periteonal macrophages. The antigen recognized by this antibody is a surface glycoprotein of Mr 48,500-52,000and is the homologue of the human CD4 and the mouse L3/T4 antigen. This antibody labels the rat helper subset, which mediates the helper activity for B and T cells, graft vs. host (GVH) reactivity and produces IL-2 in the mixed lymphocyte reaction(MLR). Addition of it to the MLR, inhibeted proliferation and blocks the production of IL-2. T cells which mediate cytotoxicity and suppressor functions are not labelled (Thus, cells labelled by this antibody are not labelled by MRC OX-8(CD8a)). This antibody is invaluable for separating T cell subsets for functional studies and for labelling cells in tissue sections. It has been used in studying the role of T lymphocytes in graft rejections and in studying the subsets of T cells in the rat which mediate GVH disease. This particular antibody is also one of three antibodies which labels T lymphocytes populations in the rat. These clones include W3/13 which labels all T cells, as well as MRC OX8 and W3/25 which label non-overlapping T cell subpopulations. These monoclonal antibodies used in concert are being empolyed extensively to investigate cellular aspects of the immune response in rats and prove to be useful as markers for functionally distinct subpopulations of lymphocytes.

Isotype subclass: Mouse IgG 1

Form: Purified via Protein G chomatography. Conjugated with FITC.

Physical state: Liquid

Buffer/Additives/Preservative: PBS containing 1 % BSA and 0.09 % sodium azide (pH 7.4).

Expiration date: The reagent is stable until the expiry date stated on the vial label.

Storage conditions: Store at 4 °C. Do not freeze. Avoid prolonged exposure to light.

Application: Flow Cytometry

References:

1. Williams, A.F., Galfe and C. Milstein (1997), Analysis of cell surfaces by xenogenic myeloma-hybrid antibodies: Differentiation antigens of rat lymphocytes, Cell **12**, 633-673

2. Brideau, R.J. et al. (1980), Two subsets of rat T lymphocytes defined with monoclonal antibodies, Eur.J.of Immunol. 10, 609-615

3.Barclay, A.N. (1981), The localization of populations of lymphocytes defined with monoclonal antibodies in rat lymphoid tissues, Immunology **45**, 593-600

4. Cantrell, D.A. Robins, R.A. and R.W. Baldwin (1982), Rat lymphocyte subsets: Cellular requirements for the generation of T cell growth factors, Cell Immunol. **70**, 367-372

5. Mason, D.W., Pugh, C.W. and M.Webb (1981), The rat mixed lymphocyte reaction: roles of a dendritic cells in intestinal lymph and T cell subsets defined by monoclonal antibodies, Immunology **44**, 75-87

6. Webb, M., Mason, D.W. and A.F. Williams (1982), Inhibiton of mixed lymphocyte response by monoclonal antibody specific for rat T lymphocyte subset, Nature **282**, 841-843

7. Dallman, M.J., Mason, D.W. and M. Webb (1982), The role of host and donor cells in the rejection of skin allografts by T cell deprived rats injected with syngenic T cells, Eur.J.of Immunol. **12**, 511-518

8.Mason, D.W. (1981) subsets of T cells in the rat mediating lethal graft vs host disease, Transplantation **32**, 222-226 9. White R. A. H., Mason, D., Williams, A.F. and Galfe and C. Milstein (1978) T lymphocyte heterogeneity in the rat: separation iof functional sub-populations using a monoclonal antibody, J.Exp.Med. **148**, 644-673

10. Jefferies, W.A., Green, J.R. and A.F. williams (1985), Authentic T helper CD4 (W3/25) antigen on rat periteonal macrophages, J.Exp.Med. **162**, 117-127

Warning:

Sodium azide is harmful if swallowed (R22). Keep out of reach of children (S2). Keep away from food, drink and animal feeding stuff (S13). Wear suitable protective clothing (S36). If swallowed, seek medical advice immediately and show this container or label (S46). Contact with acids liberates very toxic gas (R32). Azide compounds should be flushed with large volumes of water during disposal to avoid deposits in lead or copper plumbing where explosive conditions can develop.

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