

# OriGene Technologies Inc.

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# Monoclonal Antibody to T Cell Receptor (TCR) Vb 7 - Biotin

Alternate names: TCR V beta-7, TCR Vb7

Catalog No.: SM088B

Quantity: 0.1 mg

Concentration: 0.1 mg/ml

Host / Isotype: Rat / IgG2b

Recommended SM19B

Clone: TR310

**Isotype Controls:** 

Format: State: Liquid purified

Buffer System: PBS, 0.02% sodium azide (NaN3) and EIA grade BSA as a stabilizing protein

to bring total protein concentration to 4-5 mg/ml

Label: Biotin

**Applications:** Flow cytometry.

Immunoprecipitation.

Other applications not tested. Optimal dilutions are dependent on conditions and should

be determined by the user.

Specificity: This anti-mouse T cell receptor Vβ7 antigen monoclonal antibody reacts with TCR Vβ7

bearing T cells1. The TCR VB7 may be deleted in mouse strains expressing MIS-1a

haplotype. **Species:** Mouse.

Other species not tested.

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing. Shelf life: one year from despatch.

General Readings: 1. Okada CY, Holzmann B, Guidos C, Palmer E, Weissman IL. Characterization of a rat

monoclonal antibody specific for a determinant encoded by the V beta 7 gene segment. Depletion of V beta 7+ T cells in mice with Mls-1a haplotype. J Immunol. 1990 May

1;144(9):3473-7. PubMed PMID: 1691759.

2. Sugihara S, Fujiwara H, Shearer GM. Autoimmune thyroiditis induced in mice depleted of particular T cell subsets. Characterization of thyroiditis-inducing T cell lines and clones derived from thyroid lesions. J Immunol. 1993 Jan 15;150(2):683-94. PubMed PMID:

7678281.

3. Ignatowicz, L., Kappier, J. W. Marrack, P. and Scherer, M. T. 1994 identification of two

Vβ7-specific viral superantigens J. Immunol. 152:65-71.

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#### **Protocols:**

### **FLOW CYTOMETRY ANALYSIS:**

#### Method:

- 1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte®-M cell separation medium.
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of 2x10e7 cells/ml in media A. Add 50  $\mu$ l of this suspension to each tube (each tube will then contain 1 x 10e6 cells, representing 1 test).
- 4. To each tube, add ~1.0  $\mu$ g\* of this Ab per 10e6 cells.
- 5. Vortex the tubes to ensure thorough mixing of antibody and cells.
- 6. Incubate the tubes for 30 minutes at 4°C.
- 7. Wash 2 times at 4°C.
- 8. Add 100 µl of secondary antibody (Streptavidin-PE) at a 1:50 dilution.
- 9. Incubate tubes at 4°C for 30 60 minutes (It is recommended that tubes are protected from light since most fluorochromes are light sensitive).
- 10. Wash 2 times at 4°C.
- 11. Resuspend the cell pellet in 50 µl ice cold media B.
- 12. Transfer to suitable tubes for flow cytometric analysis containing 15  $\mu$ l of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

#### Media:

A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).

B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).

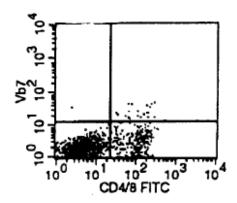
## Results - Tissue Distribution by Flow Cytometry Analysis:

(Representative dot blot)
Mouse Strain: C3H.SW

<u>Cell Concentration</u>: 1x10e6 cells per tests <u>Antibody Concentration Used</u>: 1.0 μg/10e6 cells

<u>Isotypic Control</u>: Biotin Rat IgG2b

### **Pictures:**



Cell source: Spleen
Percentage of cells stained above control: 1.6%