

Product datasheet for **AM31862PU-N**

T Cell Receptor (TCR) V alpha-2 Rat Monoclonal Antibody [Clone ID: B20.1]

Product data:

Product Type:	Primary Antibodies
Clone Name:	B20.1
Applications:	FC
Recommended Dilution:	Flow Cytometry (See Protocols). This clone has also been reported to work in Immunoprecipitation . (1,2)
Reactivity:	Mouse
Host:	Rat
Isotype:	IgG2a
Clonality:	Monoclonal
Specificity:	This antibody reacts with Mouse T-Cell Receptor (TCR) Va2 chains (1), and recognizes the majority of the TCR Va2 subfamily in mice carrying the a, b and c haplotypes 1,2. It also reacts with the products of T cell receptor, Vδ8 due to the high degree of homology (1).
Formulation:	PBS containing 0.09% Sodium Azide as preservative. State: Purified State: Liquid purified IgG fraction.
Concentration:	lot specific
Purification:	Protein G Affinity Chromatography.
Conjugation:	Unconjugated
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.
Stability:	Shelf life: one year from despatch.
Background:	The TCR alpha chain complexes with the TCR beta chain to form the T cell receptor in 95% of T cells, whereas the remaining 5% of T cells express gamma and delta chains (γ/δ). TCR Va2 is a distinct TCR subfamily found in mice having the a, b, and c haplotypes.



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Note:

Protocol: **Flow Cytometry Analysis:**

NOTE: Preblocking of Fc receptors for 10 minutes using 0.5 µg of purified anti-Mouse CD16/32 is recommended.

Method:

1. Prepare cell suspension in Media A. For cell replications, deplete the red blood cell population with Lympholyte®-M cell separation medium.
2. Wash 2 times.
3. Resuspend the cells to a concentration 2×10^7 cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain 1×10^6 cells, representing one test).
4. To each tube add 0.25 µg of this antibody AM31862PU-N per 1×10^6 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 (FITC Goat anti-Rat IgG (H+L) at a 1/500 dilution.
9. Incubate tubes at 4°C for 30-60 minutes. (It is recommended that the tubes are protected from light since most fluorochromes are light sensitive).
10. Wash 2 times at 4°C in Media B.
11. Resuspend the cell pellet in 50 µl ice cold Media B.
12. Transfer to suitable tubes for flow cytometric analysis containing 15 µl of propidium iodide at 0.5 mg/ml in phosphate buffered saline. This stains dead cells by intercalating DNA.

Media:

- A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 µl of 2 M sodium azide in 100 mls).
- B. Phosphate buffered saline (pH 7.2) + 0.5% bovine serum albumin + sodium azide (100 µl of 2 M sodium azide in 100 mls).