OriGene Technologies, Inc.
9620 Medical Center Drive, Ste 200
Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com
techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

Product datasheet for AM31862AC-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) Vα2 chains (1), and recognizes the majority of the TCR Vα2 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 and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml. Label: APC State: Liquid purified IgG fraction.
Concentration:	lot specific
Purification:	Protein G Affinity Chromatography.
Conjugation:	APC
Storage:	Store the antibody undiluted at 2-8°C. **DO NOT FREEZE!** 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 Vα2 is a distinct TCR subfamily found in mice having the a, b, and c haplotypes.

View online »

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 reparations, deplete the red blood cell population with Lympholyte®-M cell separation medium.
2. Wash 2 times.
3. Resuspend the cells to a concentration 2x10e7 cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain 1x10e6 cells, representing one test).
4. To each tube add 0.25 µg of this antibody AM31862AC-N per 1x10e6 cells.
5. Vortex the tubes to ensure thorough mixing of antibody and cells.
6. 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).
7. Wash 2 times at 4°C.
8. Resuspend the cell pellet in 50 µl ice cold Media B.
9. 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).