Product Data Sheet

Alexa Fluor® 488 anti-mouse/human CD45R/B220

Catalog # / Size: 1116125 / 100 μg

1116140 / 25 μg

Clone: RA3-6B2 Isotype: Rat IgG2a, κ

Immunogen: Abelson murine leukemia virus-induced

pre-B tumor cells

Reactivity: Human

Preparation: The antibody was purified by affinity

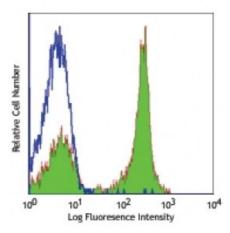
chromatography, and conjugated with Alexa Fluor® 488 under optimal

conditions.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.5



C57BL/6 mouse splenocytes stained with RA3-6B2 Alexa Fluor® 488

Applications:

Applications: Immunofluorescence

Recommended Usage:

Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤ 2.0 microg per million cells in 100 microL volume. For immunohistochemisty, a concentration range of 2.5-5 μ g/ml is suggested. For immunofluorescence microscopy, a concentration range of 1.25-10 μ g/ml is recommended. It is recommended that the reagent be titrated for optimal performance for each application.

* Alexa Fluor® 488 has a maximum emission of 519 nm when it is excited at 488 nm.

Application Notes:

Clone RA3-6B2 has been described to react with an epitope on the

extracellular domain of the transmembrane CD45 glycoprotein

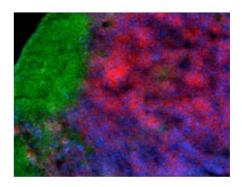
which is dependent upon the expression of exon A and specific carbohydrate residues. Additional reported

applications (for the relevant formats) include: immunoprecipitation1, *in vitro*

and *in vivo* modulation of B cell

responses²⁻⁴, and

immunohistochemistry of acetone-fixed frozen sections and formalin-fixed paraffin-embedded sections^{5,6}. The LEAF™ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered)



C57BL/6 mouse frozen lymph node section was fixed with 4% paraformaldehyde (PFA) for 10 minutes at room temperature and blocked with 5% FBS plus 5% rat serum for 1 hour at room temperature. Then the section was stained with 5 microg/ml of CD8 (clone 53-

is recommended for functional assays (Cat. No. 103216).

Application References:

- 1. Coffman RL. 1982. Immunol. Rev. 69:5. (IP)
- 2. George A, et al. 1994. J. Immunol. 152:1014. (Activ)
- 3. Asensi V, et al. 1989. Immunology 68:204. (Activ)
- 4. Domiati-Saad R, et al. 1993. J. Immunol. 151:5936. (Activ)
- 5. Hata H, et al. 2004. J. Clin. Invest. 114:582. (IHC)
- 6. Monteith CE, et al. 1996. Can. J. Vet. Res. 60:193. (IHC)
- 7. Shih FF, et al. 2006. J. Immunol. 176:3438. (FC)
- 8. Chang C L-T, et al. 2007. J. Immunol. 178:6984.
- 9. Fazilleau N, et al. 2007. Nature Immunol. 8:753.
- 10. Lang GL, et al. 2008. Blood 111:2158. PubMed
- 11. Charles N, et al. 2010. Nat. Med. 16:701. (FC) PubMed
- 12. del Rio ML, et al. 2011. Transpl. Int. 24:501. (FC) PubMed
- 13. Murakami R, et al. 2013. PLoS One. 8:73270. PubMed

Description:

CD45R, also known as B220, is an isoform of CD45. It is a member of the protein tyrosine phosphatase (PTP) family with a molecular weight of approximately 180-240 kD. CD45R is expressed on B cells (at all developmental stages from pro-B cells through mature B cells), activated B cells, and subsets of T and NK cells. CD45R (B220) is also expressed on a subset of abnormal T cells involved in the pathogenesis of systemic autoimmunity in MRL-Fas^{lpr} and MRL-Fas^{gld} mice. It plays a critical role in TCR and BCR signaling. The primary ligands for CD45 are galectin-1, CD2, CD3, and CD4. CD45R is commonly used as a pan-B cell marker; however, CD19 may be more appropriate for B cell specificity.

Antigen References:

- 1. Barclay A, et al. 1997. The Leukocyte Antigen FactsBook Academic Press.
- 2. Trowbridge IS, et al. 1993. Annu. Rev. Immunol. 12:85.
- 3. Kishihara K, et al. 1993. Cell 74:143.
- 4. Pulido R, <