Brilliant Violet 711™ anti-mouse/human CD45R/B220

Catalog # / Size: 1116275 / 50 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 Brilliant Violet 711™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 711™ and

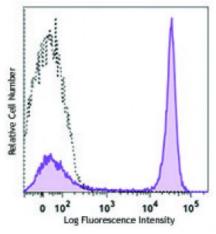
unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: 0.2



C57BL/6 mouse splenocytes were stained with CD45R/B220 (clone RA3-6B2) Brilliant Violet 711[™] (filled histogram) or rat IgG2a, κ Brilliant Violet 711[™] isotype control (open histogram).

Applications:

Applications: Flow Cytometry

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 ≤0.25 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

Brilliant Violet 711^{TM} excites at 405 nm and emits at 711 nm. The bandpass filter 710/50 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. **Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel.** Refer to your instrument manual or manufacturer for support. Brilliant Violet 711^{TM} is a trademark of Sirigen Group Ltd.

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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 LEAFTM purified antibody (Endotoxin <0.1 EU/ μ g, Azide-Free, 0.2 μ m filtered) is recommended for functional assays (Cat. No. 103216).

Application 1. Coffman RL. 1982. *Immunol. Rev.* 69:5. (IP)

References:

- 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)
- Hata H, et al. 2004. J. Clin. Invest. 114:582. (IHC)
 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. <u>PubMed</u>
- 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, <