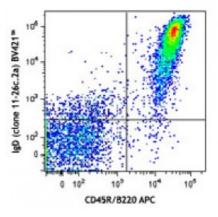
Product Data Sheet

Brilliant Violet 421[™] anti-mouse IgD

Catalog # / Size:	2628625 / 50 μg
Clone:	11-26c.2a
Isotype:	Rat IgG2a, к
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 421 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 421 [™] and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).
Concentration:	0.2

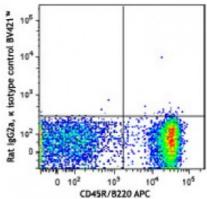


C57BL/6 splenocytes were stained with CD45R/B220 APC and IgD (clone 11-26c.2a) Brilliant Violet 421[™] (top) or rat IgG2a, κ Brilliant Violet 421[™] isotype control (bottom).

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 421 [™] excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421 [™] is a trademark of Sirigen Group Ltd.
	This product is subject to proprietary rights of Sirigen Inc. and is made and sold under license from Sirigen Inc. The purchase of this product conveys to the buyer a non-transferable right to use the purchased product for research purposes only. This product may not be resold or incorporated in any manner into another product for resale. Any use for therapeutics or diagnostics is strictly prohibited. This product is covered by U.S. Patent(s), pending patent applications and foreign equivalents.

Application The 11-26c.2a antibody reacts with



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Notes:	immunoglobulin D in all tested mouse haplotypes. The antibody binds membrane IgD expressed on most B cells. The 11-26c.2a antibody neither induces proliferation of splenic B cells nor induces B cell activation. Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen sections ^{2,3} .
Application References:	 Nitschke L, <i>et al.</i> 1993. <i>P. Natl. Acad. Sci. USA</i> 90:1887. (FC) Weih D, <i>et al.</i> 2001. <i>J. Immunol.</i> 167:1909. (IHC) Koni PA, <i>et al.</i> 2001. <i>J. Exp. Med.</i> 193:741. (IHC) Ahuja A, <i>et al.</i> 2007. <i>J. Immunol.</i> 179:3351. (FC) <u>PubMed</u> Haynes NM, <i>et al.</i> 2007. <i>J. Immunol.</i> 179:5099. (FC) Good-Jacobson KL, <i>et al.</i> 2010. <i>Nat. Immunol.</i> 11:535. (FC) <u>PubMed</u> Tomayko MM, <i>et al.</i> 2010. <i>J. Immunol.</i> 185:7146. <u>PubMed</u> Park SY, <i>et al.</i> 2013. <i>J. Immunol.</i> 190:1094. <u>PubMed</u>

Description: Surface IgD is an important B cell differentiation marker.