

Brilliant Violet 421™ anti-human CD158e1 (KIR3DL1, NKB1)

Catalog # / Size: 2163565 / 25 tests
2163570 / 100 tests

Clone: DX9

Isotype: Mouse IgG1, κ

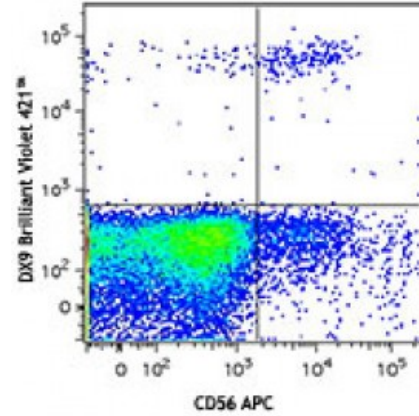
Immunogen: Human NK cell clone VL186-1.6

Reactivity: Human

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: Lot-specific

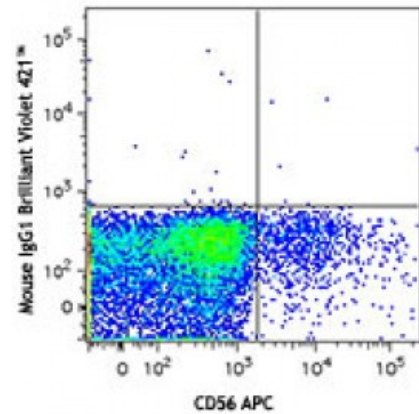


Human peripheral blood lymphocytes were stained with CD56 APC and CD158e1 (clone DX9) Brilliant Violet 421™ (top) or mouse IgG1 Brilliant Violet 421™ isotype control (bottom).

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 ≤5 microL per million cells or 5 microL per 100 microL of whole blood. 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.

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Application Notes: The DX9 antibody reacts with the KIR (killer cell inhibitory receptor) designated NKB1 or KIR3DL1. Additional reported applications (for the relevant formats) include: immunoprecipitation¹ and restoring the NK cell cytotoxicity^{4,8}. The LEAF™ purified antibody (Endotoxin <0.1 EU/μg, Azide-Free, 0.2 μm filtered) is recommended for functional assays (Cat. No. 312710).

Application References:

1. Litwin V, *et al.* 1994. *J. Exp. Med.* 180:537. (IP)
2. Gumperz J, *et al.* 1996. *J. Exp. Med.* 183:1817.
3. Gardiner CM, *et al.* 2001. *J. Immunol.* 166:2992.
4. Bakker ABH, *et al.* 1998. *J. Immunol.* 160:5239.
5. Goodier M, *et al.* 2000. *J. Immunol.* 165:139.
6. Kirwan SE and Burshtyn DN. 2005. *J. Immunol.* 175:5006. (FC)
7. Yawata M, *et al.* 2002. *Immunogenetics* 54:543.
8. Valiante NM, *et al.* 1997. *Immunity* 7:739.
9. Pascal V, *et al.* 2007. *J. Immunol.* 179:1625. (FC) [PubMed](#)
10. Lichterfeld M, *et al.* 2008. *J. Exp. Med.* 204:2813. (FC) [PubMed](#)
11. Terszowski G, *et al.* 2014. *J Immunol.* 192:5618. [PubMed](#)
12. Boudreau JE, *et al.* 2014. *PLoS One.* 9:99543. [PubMed](#)
13. Purdy AK, *et al.* 2014. *J Immunol.* 193:4675. [PubMed](#)
14. Lisovsky I, *et al.* 2015. *J Lukoc Biol.* 97:761. [PubMed](#)

Description: CD158e1, also known as NKB1, is a 70 kD member of the immunoglobulin superfamily that is expressed on a subset of natural killer cells and T cells at varying levels among individuals. NKB1 is a type I membrane protein containing two immunoglobulin C2-type domains. The interaction of NKB1 with specific HLA-B antigens on a target cell (the HLA-Bw4 allele, for example) inhibits cytotoxicity and prevents target cell lysis and death. The interactions between KIR and MHC class I are thought to be important in NK and T cell regulation following antigen stimulation. The absence of ligands for KIRs may lower the threshold for activation through activating receptors and increase inflammation and susceptibility to autoimmune disease.

Antigen References:

1. Colonna M, *et al.* 1995. *Science* 268:405.
2. D'Andrea A, *et al.* 1995. *J. Immunol.* 155:2306.
3. Uhrburg M, *et al.* 1997. *Immunity* 7:753.
4. Gumperz JE, *et al.* 1996. *J. E*