

**Brilliant Violet 421™ anti-mouse CD69**

**Catalog # / Size:** 1122635 / 125 µl  
 1122640 / 500 µl  
 1122725 / 50 µg

**Clone:** H1.2F3

**Isotype:** Hamster IgG

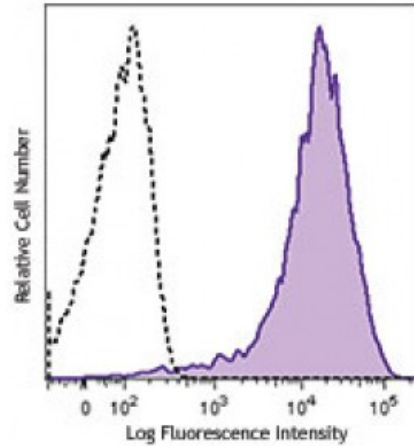
**Immunogen:** Mouse dendritic epidermal T cell line Y245

**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:** Lot-specific



PMA+ionomycin-stimulated (6 hours) C57BL/6 mouse splenocytes were stained with CD69 (clone H1.2F3) Brilliant Violet 421™ (filled histogram) or Armenian hamster IgG Brilliant Violet 421™ 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 ≤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 H1.2F3 antibody has been reported to augment T cell activation. Additional reported applications (for the relevant formats) include: *in vitro* T cell and NK cell activation<sup>1-3</sup>, immunohistochemistry<sup>4,5</sup>, and immunoprecipitation<sup>1</sup>.

This antibody has been characterized in the literature as containing a λ (λ) light chain.

- Application References:**
1. Yokoyama WM, *et al.* 1988. *J. Immunol.* 141:369. (IP)
  2. Sobel ES, *et al.* 1993. *J. Immunol.* 150:673.
  3. Karlhofer FM, *et al.* 1991. *J. Immunol.* 146:3662.
  4. Zhou X, *et al.* 2005. *J. Biol. Chem.* 280:31240. (IHC)

5. Podd BS, *et al.* 2006. *J. Immunol.* 176:6532. (IHC)
  6. Lawson BR, *et al.* 2007. *J. Immunol.* 178:5366.
  7. Lee JW, *et al.* 2006. *Nature Immunol.* 8:181.
  8. Epardaud M, *et al.* 2008. *Cancer Res.* 15:2972. [PubMed](#)
  9. Jordan JM, *et al.* 2008. 76:3717. [PubMed](#)
  10. Kenna TJ, *et al.* 2008. *Blood* 111:2091. [PubMed](#)
  11. Ishikawa C, *et al.* 2013. *Biochim Biophys Acta.* 167:99. [PubMed](#)
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**Description:** CD69 is a 60 kD type II membrane protein composed of a 27/33 kD disulfide-linked homodimer, also known as Very Early Activation Antigen (VEA), AIM, EA1, MLR3, and gp34/28. It is expressed on a subset of thymocytes and platelets. CD69 is rapidly induced on activated T and B cells, neutrophils, and NK cells. It is a C-type lectin, closely related to the NKR-P1 and Ly-49 NK cell activation molecules. CD69 is involved in the early events of cell activation and thymocyte positive selection.

**Antigen**  
**References:**

1. Barclay AN, *et al.* 1997. *The Leukocyte Antigen FactsBook* Academic Press.
2. Testi R, *et al.* 1994. *Immunol. Today* 15:479.
3. Moretta A, *et al.* 1991. *J. Exp. Med.* 174:1393.
4. Yokoyama WM,