Brilliant Violet 421™ anti-human HLA-E

Catalog # / Size: 2313055 / 25 tests

2313060 / 100 tests

Clone: 3D12

Isotype: Mouse IgG1, κ

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

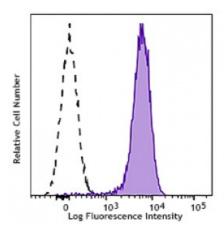
unconjugated antibody.

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 HLA-E (clone 3D12) Brilliant Violet 421[™] (filled histogram) or mouse IgG1, κ 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 μ l per million cells or 5 μ l per 100 μ l of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

Brilliant Violet 421^{TM} excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421^{TM} is a trademark of Sirigen Group Ltd.

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Application 1. Sullivan LC, et al. 2008. *Tissue Antigen.* 72:415. **References:** 2. Koller BH, et al. 1988. *J. Immunol.* 141:897.

Description: HLA-E is a non-classical MHC class I (MHC-Ib) molecule. It is characterized by

limited polymorphism and ubiquitous expression. HLA-E, as other MHC-I molecules, is heterodimerized with $\beta2\text{-microglobulin}.$ HLA-E interacts with

CD94/NKG2 receptor to regulate NK cell cytolysis activities.

Antigen 1. Sullivan LC, et al. 2008. Tissue Antigen. 72:415. References: 2. Koller BH, et al. 1988. J. Immunol. 141:897.

