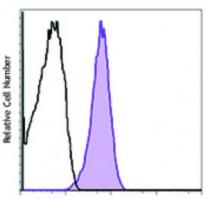
Product Data Sheet

PerCP/Cy5.5 anti-human HLA-E

| Catalog # / Size: | 2313050 / 100 tests 2313045 / 25 tests |
|--------------------|---|
| Clone: | 3D12 |
| Isotype: | Mouse IgG1, к |
| Reactivity: | Human |
| Preparation: | The antibody was purified by affinity chromatography and conjugated with PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cy5.5 and unconjugated antibody. |
| Formulation: | Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA). |
| Concentration: | 0.5 |



Log Fluorescence Intensity

Human peripheral blood lymphocytes were stained with HLA-E (clone 3D12) PerCP/Cy5.5 (filled histogram) or mouse lgG1, κ PerCP/Cy5.5 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. |
| | * PerCP/Cy5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm. |
| Application References: | 1. Lee N, <i>et al.</i> 1998. <i>Proc. Natl. Acad. Sci. USA</i> . 95:5199. <i>2.</i> Wooden SL, <i>et al.</i> 2005. <i>J. Immunol.</i> 175:1383. <i>3.</i> Monaco EL, <i>et al.</i> 2008. <i>J. Immunol.</i> 181:5442. 4. Corrah TW, <i>et al.</i> 2011. <i>J. Virol.</i> 85:3367. <u>PubMed</u> |
| 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 β 2-microglobulin. HLA-E interacts with CD94/NKG2 receptor to regulate NK cell cytolysis activities. |
| Antigen References: | 1. Sullivan LC, <i>et al.</i> 2008. <i>Tissue Antigen.</i> 72:415. 2. Koller BH, <i>et al.</i> 1988. <i>J. Immunol.</i> 141:897. |

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