Brilliant Violet 421[™] anti-human CD71

Catalog # / Size:	2270605 / 25 tests 2270610 / 100 tests	
Clone:	CY1G4	
Isotype:	Mouse IgG2a, к	Jagen
Immunogen:	NALM-6 pre-B cell line	
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).	PHA-stimulated (3 days) human peripheral blood lymphocytes were stained with CD71 (clone CY1G4)
Workshop Number:	A015	Brilliant Violet 421â"¢ (filled histogram) or mouse IgG2a, κ Brilliant Violet 421â"¢ isotype control (open histogram).
Concentration:	Lot-specific	

Applications:

Applications: Recommended Usage:	Flow Cytometry 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 [™] 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 References:	1. Hentze M, <i>et al.</i> 1996. <i>P. Natl. Acad. Sci. USA</i> 93:8175. 2. Trowbridge I, <i>et al.</i> 1993. <i>Annu. Rev. Cell Biol.</i> 9:129.	
Description:	CD71 is a 95 kD type II homodimeric transmembrane glycoprotein also known as T9 and transferrin receptor. It is expressed on proliferating cells, reticulocytes, and erythroid precursors. CD71 plays a role in the control of cellular proliferation by facilitating the uptake of iron via ferrotransferrin binding and the recycling of apotransferrin to the cell surface.	
Antigen References:	1. Hentze M, <i>et al.</i> 1996. <i>P. Natl. Acad. Sci. USA</i> 93:8175. 2. Trowbridge I, <i>et al.</i> 1993. <i>Annu. Rev. Cell Biol.</i> 9:129.	

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