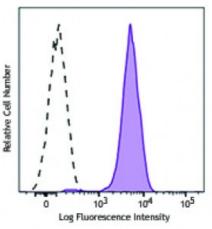
Product Data Sheet

Brilliant Violet 421[™] anti-ERK1/2 Phospho (Thr202/Tyr204)

Catalog # / Size:	2447545 / 25 tests 2447550 / 100 tests	
Clone:	6B8B69	
Isotype:	Mouse IgG2a, κ	
Immunogen:	Synthetic peptide (TGFLT*EY*VATRC) conjugated to KLH.	
Reactivity:	Human,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.	ŀ
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).	
Concentration:	Lot-specific	(t



Human peripheral blood lymphocytes were stimulated with (filled histogram) or without (open histogram) Cell Activation Cocktail (without Brefeldin A) for 15 minutes, then fixed with Fixation Buffer, permeabilized with True-Phos[™] Perm Buffer, and int

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by intracellular flow cytometry . 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.
Application Notes:	Clone 6B8B69 was found to strongly cross-react with mouse ERK1/2 Phospho (Thr202/Tyr204) when tested in-house on C57BL/6 mouse splenocytes.
Description:	ERK1/2 are members of mitogen-activated kinases (MAPKs) of serine/threonine protein kinases. ERK1/2 can be activated by a range of extracellular stimuli, such as mitogen, growth factors, neurotransmitters, chemokines, and cytokines, through receptor tyrosine kinases (RTK), G protein-coupled receptors (GPCRs), or protein kinase C (PKC). Upon stimulation, ERK1/2 are phosphorylated by the upstream kinase MEK on residues Thr202 and Tyr204 and in turn phosphorylate many other downstream molecules that are involved in a range of cellular processes such as cell proliferation, differentiation, motility and cell death.
Antigen References:	 Futran AS, <i>et al.</i> 2013. <i>Curr. Biol.</i> 23:R972. Mendoza MC, <i>et al.</i> 2011. <i>Trends Biochem. Sci.</i> 36:320. Chambard JC, <i>et al.</i> 2007. <i>Biochim. Biophys. Acta.</i> 1773:1299. Roux PP,

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