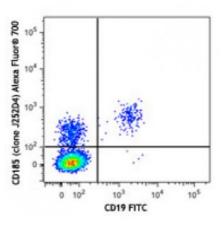
Product Data Sheet

Alexa Fluor® 700 anti-human CD185 (CXCR5)

Catalog # / Size:	2384575 / 25 tests 2384580 / 100 tests
Clone:	J252D4
Isotype:	Mouse IgG1, к
Immunogen:	Human CXCR5-transfected cells
Reactivity:	Human
Preparation:	The antibody was purified by affinity chromatography and conjugated with Alexa Fluor® 700 under optimal conditions.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).
Concentration:	Lot-specific



Human peripheral blood lymphocytes were stained with CD19 FITC and CD185 (clone [252D4) Alexa Fluor® 700 (top) or mouse IgG1, κ Alexa Fluor® 700 isotype control (bottom).

Applications:

Applications:		8 10 ⁵		
Applications:	Flow Cytometry	orte 70		
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.	Mouse IgG1, K Alexa Fluort 700	0 10 ²	10 ³ 10 ⁴ 10 ⁵ CD19 FITC
	* Alexa Fluor® 700 has a maximum emission of 719 nm when it is excited at 633 nm / 635 nm. Prior to using Alexa Fluor® 700 conjugate for flow cytometric analysis, please verify your flow cytometer's capability of exciting and detecting the fluorochrome.			
Description:	CD185, also known as CXCR5, is a 42 kD G-protein coupled receptor with seven transmembrane regions. CXCR5 is expressed by mature B cells, follicular helper T cells, Burkitt's lymphoma cells and a subset of neurons, and mediates cell migration to the B cell follicles in the secondary lymphoid organs. The ligand of CXCR5 is CXCL13 (BLC).			
Antigen References:	1. Ma CS, <i>et al.</i> 2012. <i>J. Exp. Med.</i> 209:124 2. León B, <i>et al.</i> 2012. <i>Nat. Immunol.</i> 13:68			

5.	2. LEON D, <i>EL AI</i> . 2012. <i>NAL. IIIIIIUIDI</i> . 15:001.
	3. Crotty S. 2011. Annu. Rev. Immunol. 29:621.

4. Kerfoot SM, et al. 2011. Im

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