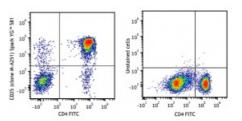
Spark YG[™] 581 anti-human CD25

Catalog # / Size:	2380765 / 25 tests 2380770 / 100 tests	
Clone:	M-A251	
lsotype:	Mouse IgG1, к	
Immunogen:	Human PHA-induced lymphocyte cells	
Reactivity:	Human, Non-human primate, Other	
Preparation:	The antibody was purified by affinity chromatography and conjugated with Spark YG™	
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA)	
Workshop Number:	581 under optimal conditions.	
Concentration:	Lot-specific	

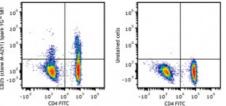


PHA-stimulated (3 days) human peripheral blood lymphocytes were stained with anti-human CD4 FITC and anti-human CD25 (clone M-A251) Spark YG[™] 581 (left) or CD4 FITC only (right).

Applications:

References:

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 in 100 μ L staining volume 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.	Human peripheral blo
	* Spark YG™ 581 has a maximum excitation of 562 nm and a maximum emission of 581 nm.	lymphocytes were sta anti-human CD4 FITC human CD25 (clone N Spark YG™ 581 (left)
Application Notes:	Additional reported applications (for the relevant formats) include: immunohistochemical staining of paraformaldehyde fixed frozen sections. ¹	human CD4 FITC only
	The CD25 molecule reveals three epitope regions: A, B, and C. M-A251 antibody recognizes epitope region B. Unlike other CD25 antibody clones, M-A251 can detect CD25 after fixation with paraformaldehyde.	
Application	1. Li H and Pauza CD. 2015. Eur. J. Im	munol. 45:298. (IHC)



iman peripheral blood mphocytes were stained with ti-human CD4 FITC and anti-Iman CD25 (clone M-A251) ark YG™ 581 (left) or anti-Iman CD4 FITC only (right).

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Description: CD25 is a 55 kD type I transmembrane glycoprotein also known as low affinity IL-2 receptor α chain or Tac. It is expressed on progenitor lymphocytes, activated T and B cells, and activated monocytes/macrophages. CD25 is also expressed on a subset of non-stimulated CD4⁺ T cells termed T regulatory cells. Soluble CD25/IL-2Rα is produced as a consequence of lymphocyte stimulation and is found in biological fluids following inflammatory responses. CD25 associates with IL-2 receptor β (CD122) and common γ (CD132) chains to form a high affinity IL-2R complex.
Antigen 1. Knapp W, *et al.* 1989. Leucocyte Typing IV: White Cell Differentiation

References:Antigens. Oxford University Press.2. Schlossman S, et al. 1995. Leucocyte Typing V: White Cell Differentiation

Antigens. Oxford University Press.Barclay N, *et al.* 1997. The Leukocyte Antigen FactsBook. Academic Press Inc.

4. Taniguchi T and Minami Y. et al. 1993. Cell 73:5.

5. Waldmann T. 1991. J. Biol. Chem. 266:2681.