Spark Violet™ 538 anti-human CD3

Catalog # / 2102420 / 100 tests

Size: 2102415 / 25 tests

Clone: UCHT1

Isotype: Mouse IgG1, κ

Immunogen: Human T cells from a T-ALL patient.

Reactivity: Human, Other

Preparation: The antibody was purified by affinity

chromatography and conjugated with Spark Violet™ 538 under optimal

conditions.

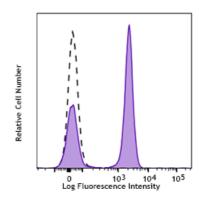
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

0.2% (w/v) BSA (origin USA)

Workshop Number: III 471

Concentration: Lot-specific



Human peripheral blood lymphocytes were stained with CD3 (clone UCHT1) Spark Violet™ 538 (filled histogram) or mouse lgG1, k Spark Violet™ 538 isotype control (open histogram).

Applications:

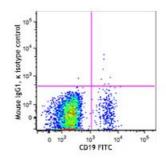
Applications: Flow Cytometry

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 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.

* Spark Violet™ 538 has a maximum excitation of 396 nm and a maximum emission of 538 nm.



Human peripheral blood lymphocytes were stained with anti-human CD4 FITC and antihuman CD25 (clone M-A251) Spark YG™ 581 (left) or antihuman CD4 FITC only (right).

Application Notes:

Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen sections^{4,6,7} and formalin-fixed paraffin-embedded sections 11, immunoprecipitation 1, activation of T cells^{2,3,5}, and Western blotting 9 . The LEAF $^{\mathrm{m}}$ purified antibody (Endotoxin < 0.1 EU/μg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 300413, 300414, and 300432). For highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 300437, 300438, 300465, 300466, 300473, 300474) with a lower endotoxin limit than standard LEAF[™] purified antibodies (Endotoxin $< 0.01 EU/\mu g$).

Application References:

- 1. Salmeron A, et al. 1991. J. Immunol. 147:3047. (IP)
- 2. Graves J, et al. 1991. J. Immunol. 146:2102. (Activ)
- 3. Lafont V, et al. 2000. J. Biol. Chem. 275:19282. (Activ)
- 4. Ryschich E, et al. 2003. Tissue Antigens 62:48. (IHC)
- 5. Thompson AG, et al. 2004. J. Immunol. 173:1671. (Activ)
- 6. Sakkas LI, et al. 1998. Clin. Diagn. Lab. Immun. 5:430. (IHC)
- 7. Mack CL, et al. 2004. Pediatr. Res. 56:79. (IHC)
- 8. Thakral D, et al. 2008. J. Immunol. 180:7431. (FC) PubMed
- 9. Van Dongen JJM, et al. 1988. Blood 71:603. (WB)
- 10. Yoshino N, et al. 2000. Exp. Anim. (Tokyo) 49:97. (FC)
- 11. Pollard, K. et al. 1987. J. Histochem. Cytochem. 35:1329. (IHC)
- 12. Luckashenak N, et al. 2013. J. Immunol. 190:27. PubMed
- 13. Laurent AJ, et al. 2014. PLoS One. 9:103683. PubMed
- 14. Li J, et al. 2015. Cancer Res. 75:508. PubMed
- 15. Stoeckius M, et al. 2017. Nat. Methods. 14:865-868. (PG)

Description:

CD3 ϵ is a 20 kD chain of the CD3/T-cell receptor (TCR) complex which is composed of two CD3 ϵ , one CD3 γ , one CD3 δ , one CD3 ζ (CD247), and a T-cell receptor (α/β or γ/δ) heterodimer. It is found on all mature T cells, NKT cells, and some thymocytes. CD3, also known as T3, is a member of the immunoglobulin superfamily that plays a role in antigen recognition, signal transduction, and T cell activation.

Antigen References:

- 1. Barclay N, et al. 1993. The Leucocyte FactsBook. Academic Press. San Diego.
- 2. Beverly P, et al. 1981. Eur. J. Immunol. 11:329.
- 3. Lanier L, et al. 1986. J. Immunol. 137:2501-2507.