Product Data Sheet

Alexa Fluor® 647 anti-human CD3

Catalog # / Size: 2102110 / 25 tests

2102080 / 100 tests

Clone: UCHT1

Isotype: Mouse IgG1, κ

Reactivity: Human

Preparation: The antibody was purified by affinity

chromatography, and conjugated with

Alexa Fluor® 647 under optimal

conditions.

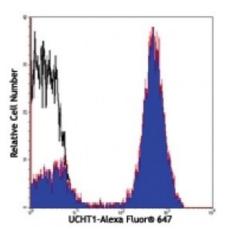
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

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

Workshop Number: III 471

Concentration: Lot-specific



Human peripheral blood lymphocytes stained with UCHT1 Alexa Fluor® 647.

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 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.

* Alexa Fluor® 647 has a maximum emission of 668 nm when it is excited at

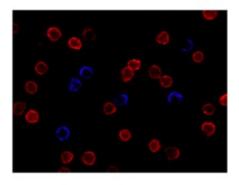
633nm / 635nm.

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 sections¹¹, immunoprecipitation1, activation of T cells^{2,3,5}, and Western blotting⁹. The LEAF™ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 300414). For highly sensitive assays, we recommend

functional assays (Cat. No. 300414). For highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 300438) with a lower endotoxin limit than standard LEAF™ purified antibodies

(Endotoxin < 0.01 EU/microg).



Human peripheral mononuclear cells were fixed with 2% paraformaldehyde (PFA), and then stained with 5 microg/ml CD19 (clone HIB19) Brilliant Violet 421™ (blue) and 5microg/ml CD3 (clone UCHT1) Alexa Fluor® 647 (red) for 30 minutes at room te

Application 1. Salmeron A, et al. 1991. J. Immunol. 147:3047. (IP)

References: 2. Graves J, et al. 1991. J. Immunol. 146:2102. (Activ)

Lafont V, et al. 2000. J. Biol. Chem. 275:19282. (Activ)
Ryschich E, et al. 2003. Tissue Antigens 62:48. (IHC)
Thompson AG, et al. 2004. J. Immunol. 173:1671. (Activ)
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.

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.