

Brilliant Violet 711™ anti-human CD3

Catalog # / Size: 2186640 / 100 tests
2186635 / 25 tests

Clone: OKT3

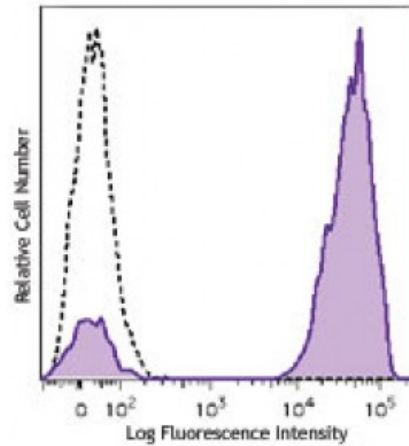
Isotype: Mouse IgG2a, κ

Reactivity: Human

Preparation: The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 711™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 711™ and unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).

Concentration: Lot-specific



Human peripheral lymphocytes were stained with CD3 (clone OKT3) Brilliant Violet 711™.

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.

Brilliant Violet 711™ excites at 405 nm and emits at 711 nm. The bandpass filter 710/50 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. **Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel.** Refer to your instrument manual or manufacturer for support. Brilliant Violet 711™ is a trademark of Sirigen Group Ltd.

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Application Notes: The OKT3 monoclonal antibody reacts with an epitope on the epsilon-subunit within the human CD3 complex.

Clone OKT3 can block the binding of clones SK7 and UCHT1.4 The OKT3 antibody is able to induce T cell activation. Additional reported applications (for the relevant formats) include: immunohistochemical staining of acetone-fixed frozen sections and activation of T cells. The LEAF™ purified antibody (Endotoxin <0.1 EU/μg, Azide-Free, 0.2 μm filtered) is recommended for functional assays (Cat. No. 317304). For highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 317326) with a lower endotoxin limit than standard LEAF™ purified antibodies (Endotoxin <0.01 EU/microg).

Application 1. Schlossman S, *et al.* Eds. 1995. Leucocyte Typing V. Oxford University Press.

- References:** New York.
2. Knapp W. 1989. Leucocyte Typing IV. Oxford University Press New York.
 3. Barclay N, *et al.* 1997. The Leucocyte Antigen Facts Book. Academic Press Inc. San Diego.
 4. Li B, *et al.* 2005. *Immunology* 116:487.
 5. Jeong HY, *et al.* 2008. *J. Leuckocyte Biol.* 83:755. [PubMed](#)
 6. Alter G, *et al.* 2008. *J. Virol.* 82:9668. [PubMed](#)
 7. Manevich-Mendelson E, *et al.* 2009. *Blood* 114:2344. [PubMed](#)
 8. Pinto JP, *et al.* 2010. *Immunology.* 130:217. [PubMed](#)
 9. Biggs MJ, *et al.* 2011. *J. R. Soc. Interface.* 8:1462. [PubMed](#)
 10. Willmann KL, *et al.* 2014. *Nat Commun.* 5:5360. [PubMed](#)
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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 lymphocytes, NK T 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.