Product Data Sheet

Spark NIR™ 685 anti-human FOXP3

Catalog # / 2200650 / 100 tests

Size: 2200645 / 25 tests

Clone: 206D

Isotype: Mouse IgG1, κ

Immunogen: Full-length FOXP3 protein

Reactivity: Human, Non-human primate

Preparation: The antibody was purified by affinity

chromatography and conjugated with

Spark NIR™ 685 under optimal

conditions.

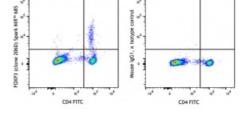
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

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

Workshop Number: 750 under optimal conditions.

Concentration: Lot-specific



Human peripheral blood lymphocytes were surface stained with CD4 FITC and then treated with True-Nuclear™ Transcription Factor Buffer set. Cells were then stained with anti-human FOXP3 (clone 206D) Spark NIR™ 685 (left) or mouse IgG1, κ Spark NIR™ 685 isotype control (right).

Applications:

Applications: Intracellular Staining for Flow

Cytometry

Recommended

Usage:

Each lot of this antibody is quality control tested by intracellular

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

each application.

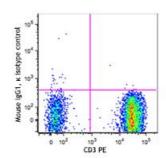
* Spark NIR™ 685 has a maximum excitation of 665 nm and a maximum

titrated for optimal performance for

emission of 685 nm.

Application Notes:

Additional reported applications (for relevant formats) include: the immunohistochemical staining acetone-fixed frozen sections 1 and paraffin-embedded formalin-fixed sections^{1,8,19-20} and Western blotting¹. The binding of 206D to FOXP3 can be partially blocked by 259D, but 206D does not show significant blocking effect on 259D binding.



Application References:

- 1. Roncador G, et al. 2005. Eur. J. Immunol. 35:1681.(IHC)
 - 2. Yang ZZ, et al. 2006. Blood 107:3639.
 - 3. Liu W, et al. 2006. J. Exp. Med. 203:1701. PubMed
 - 4. Bollyky PL, et al. 2007. J. Immunol. 179:744.
 - 5. Bell MP, et al. 2007. J. Immunol. 179:1893.
- 6. Tran DQ, et al. 2007. Blood doi:10.1182/blood-2007-06-094656. PubMed
- 7. Gao Q,et al.2007. J Clin Oncol. 25: 2586. (IHC) PubMed
- 8. Pillai V, et al. 2008. Blood 111:463. PubMed
- 9. Zheng Y, et al. 2008. J. Immunol. 181:1683. PubMed
- 10. Zonios DI, et al. 2008. Blood 112:287. PubMed
- 11. Kavanagh B, et al. 2008. Blood. PubMed
- 12. Nevala WK, et al. 2009. Clin Cancer Res. 15:1931. PubMed
- 13. Grant J, et al. 2009. Cytometry B Clin Cytom. 76:69. PubMed
- 14. Nigam P, et al. 2010. J. Immunol. 184:1690. PubMed
- 15. Kmieciak M, et al. 2009. J. Transl. Med. 7:89. (ICFC) PubMed
- 16. Hartigan-O'Connor DJ,et al.2007.J Exp Med.204:2679. PubMed
- 17. Raghaven S, et al. 2009. Ann Rheum Dis. 68:1908. PubMed
- 18. Hodi FS, et al. 2014. Cancer Immunol Res. 2:632.(IHC) PubMed
- 19. Sziros E, et al. 2015. Clin Cancer Res. 21:2840.(IHC) PubMed

Description:

FOXP3 is a 50-55 kD transcription factor, also known as Forkhead box protein P3, Scurfin, JM2, or IPEX. It is proposed to be a master regulatory gene and more specific marker of T regulatory cells than most cell surface markers (such as CD4 and CD25). Transduced expression of FOXP3 in CD4⁺/CD25⁻ cells has been shown to induce GITR, CD103, and CTLA4 and impart a T regulatory cell phenotype. FOXP3 is mutated in X-linked autoimmunity-allergic dysregulation syndrome (XLAAD or IPEX) in humans and in "scurfy" mice. Overexpression of FOXP3 has been shown to lead to a hypoactive immune state suggesting that this transcriptional factor is a central regulator of T cell activity. In human, unlike in mouse, two isoforms of FOXP3 have been reported: one (FOXP3) corresponding to the canonical full-length sequence; the other (FOXP3 δ2) lacking exon 2. The 206D antibody recognizes human FOXP3 epitope in the region of amino acids 105-235.

Antigen References:

- 1. Hori S, et al. 2003. Science 299:1057.
- References: 2. Gandhi R, et al. 2010. Nat. Immunol. 11:846.