

Alexa Fluor® 488 anti-Tubulin β 3 (TUBB3)

Catalog # / Size: 3887020 / 100 µg
3887015 / 25 µg

Clone: AA10

Isotype: Mouse IgG2a, κ

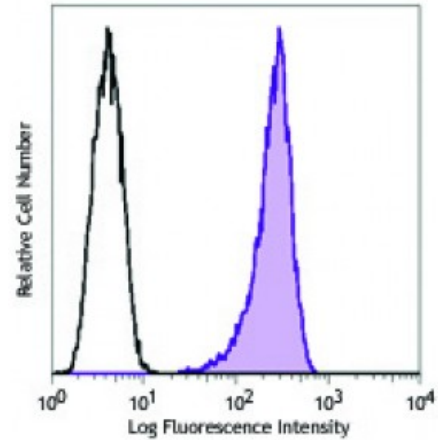
Immunogen: Fusion protein

Reactivity: Human, Mouse, Rat

Preparation: The antibody was purified by affinity chromatography and conjugated with Alexa Fluor® 488 under optimal conditions.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

Concentration: 0.5



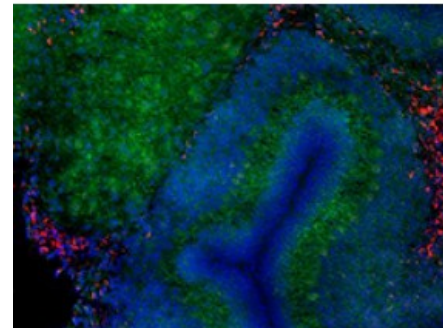
Human lung adenocarcinoma cell line A549 was treated with BioLegend's Fixation Buffer (Cat. No. 420801) and Permeabilization Wash Buffer (Cat. No. 421002), and then stained with TUBB3 (clone AA10) Alexa Fluor® 488 (filled histogram) or mouse I

Applications:

Applications: 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 ≤0.125 microg per million cells in 100 microL volume. For immunohistochemical staining on frozen tissue sections, the suggested use is 1.25-5 microg/mL. It is recommended that the reagent be titrated for optimal performance for each application.

* Alexa Fluor® 488 has a maximum emission of 519 nm when it is excited at 488 nm.



C57BL/6 mouse frozen brain tissue was fixed with 4% paraformaldehyde (PFA) for ten minutes, permeabilized with 0.5 % Triton X-100 for ten minutes, and blocked with 5% FBS for 1 hour. Then the tissue was stained with 5 microg/ml of Alexa Fluor® 488 a

Description: Tubulin is the main component of microtubules. In adults, tubulin β 3 (TUBB3) is primarily expressed in neurons and is commonly used as a neuronal marker. It plays an important role in neuronal cell proliferation and differentiation. Mutations in this gene cause congenital fibrosis of the type 3 extraocular muscles. Tubulin β 3 (TUBB3) is also found in a wide range of tumors. Studies indicate that it is a predictive and prognostic marker in various tumors.

Antigen 1. Katsetos CD, *et al.* 2003. *J. Child Neurol.* 18:851.

- References:**
2. Mobarakeh ZT, *et al.* 2012. *Cell Biol. Int. Rep. (2010)* 19:e00015.
 3. Locher H, *et al.* 2013. *Differentiation*. 85:173.
 4. Kar