

1F-250-C100

Monoclonal Antibody to alpha-tubulin Fluorescein (FITC) conjugated (0.1 mg)

Clone:	TU-01
Isotype:	Mouse IgG1
Specificity:	The antibody TU-01 recognizes the defined epitope (aa 65-97) on N-terminal structural domain of alpha-tubulin.
Regulatory Status:	RUO
Immunogen:	Fraction of tubulin purified from porcine brain by two cycles of polymerization - depolymerization.
Species Reactivity:	Broad species reactivity
Preparation:	The purified antibody is conjugated with Fluorescein isothiocyanate (FITC) under optimum conditions. The reagent is free of unconjugated FITC.
Concentration:	1 mg/ml
Storage Buffer:	Phosphate buffered saline (PBS) with 15 mM sodium azide, approx. pH 7.4
Storage / Stability:	Store in the dark at 2-8°C. Do not freeze. Avoid prolonged exposure to light. Do not use after expiration date stamped on vial label.
Usage:	The reagent is designed for direct immunofluorescence analysis on fixed and permeabilized cells. Suggested working dilution is 1:50. It is recommended that the user titrates the reagent for use in the particular testing system.
Expiration:	See vial label
Lot Number:	See vial label

For laboratory research only, not for drug, diagnostic or other use.



Background:

The microtubules are intracellular dynamic polymers made up of evolutionarily conserved polymorphic alpha/beta-tubulin heterodimers and a large number of microtubule-associated proteins (MAPs). The microtubules consist of 13 protofilaments and have an outer diameter 25 nm. Microtubules have their intrinsic polarity; highly dynamic plus ends and less dynamic minus ends. Microtubules are required for vital processes in eukaryotic cells including mitosis, meiosis, maintenance of cell shape and intracellular transport. Microtubules are also necessary for movement of cells by means of flagella and cilia. In mammalian tissue culture cells microtubules have their minus ends anchored in microtubule organizing centers (MTOCs). The GTP (guanosintriphosphate) molecule is an essential for tubulin heterodimer to associate with other heterodimers to form microtubule. In vivo, microtubule dynamics vary considerably. Microtubule polymerization is reversible and a populations of microtubules in cells are on their minus ends either growing or shortening – this phenomenon is called dynamic instability of microtubules. On a practical level, microtubules can easily be stabilized by the addition of non-hydrolysable analogues of GTP (eg. GMPPCP) or more commonly by anti-cancer drugs such as Taxol. Taxol stabilizes microtubules at room temperature for many hours. Using limited proteolysis by enzymes both tubulin subunits can be divided into N-terminal and C-terminal structural domains.

The alpha-tubulin (relative molecular weight around 50 kDa) is globular protein that exists in cells as part of soluble alpha/beta-tubulin dimer or it is polymerized into microtubules. In different species it is coded by multiple tubulin genes that form tubulin classes (in human 6 genes). Expressed tubulin genes are named tubulin isotypes. Some of the tubulin isotypes are expressed ubiquitously, while some have more restricted tissue expression.

Alpha-tubulin is also subject of numerous post-translational modifications. Tubulin isotypes and their posttranslational modifications are responsible for multiple tubulin charge variants - tubulin isoforms. Heterogeneity of alpha-tubulin is concentrated in C-terminal structural domain.

For laboratory research only, not for drug, diagnostic or other use.





Antibodies

References:

*Viklicky V, Draber P, Hasek J, Bartek J: Production and characterization of a monoclonal antitubulin antibody. Cell Biol Int Rep. 1982 Aug;6(8):725-31.

*Draber P, Draberova E, Zicconi D, Sellitto C, Viklicky V, Cappuccinelli P: Heterogeneity of microtubules recognized by monoclonal antibodies to alpha-tubulin. Eur J Cell Biol. 1986 Jun;41(1):82-8.

*Grimm M, Breitling F, Little M: Location of the epitope for the alpha-tubulin monoclonal antibody TU-O1. Biochim Biophys Acta. 1987 Jul 24;914(1):83-8.

*Draber P, Draberova E, Linhartova I, Viklicky V: Differences in the exposure of Cand N-terminal tubulin domains in cytoplasmic microtubules detected with domain-specific monoclonal antibodies. J Cell Sci. 1989 Mar;92 (Pt 3):519-28.

*Draber P, Draberova E, Viklicky V: Immunostaining of human spermatozoa with tubulin domain-specific monoclonal antibodies. Recognition of a unique beta tubulin epitope in the sperm head. Histochemistry. 1991;95(5):519-24. *Linhartova I, Draber P, Draberova E, Viklicky V: Immunological discrimination of

*Linhartova I, Draber P, Draberova E, Viklicky V: Immunological discrimination of beta-tubulin isoforms in developing mouse brain. Post-translational modification of non-class-III beta-tubulins. Biochem J. 1992 Dec 15;288 (Pt 3):919-24.

*Nováková M, Dráberová E, Schürmann W, Czihak G, Víklický V, Dráber P: gamma-Tubulin redistribution in taxol-treated mitotic cells probed by monoclonal antibodies. Cell Motil Cytoskeleton. 1996;33(1):38-51.

*Smertenko A, Blume Y, Viklicky V, Opatrny Z, Draber P: Post-translational modifications and multiple tubulin isoforms in Nicotiana tabacum L. cells. Planta. 1997;201(3):349-58.

*Kukharskyy V, Sulimenko V, Macůrek L, Sulimenko T, Dráberová E, Dráber P: Complexes of gamma-tubulin with nonreceptor protein tyrosine kinases Src and Fyn in differentiating P19 embryonal carcinoma cells. Exp Cell Res. 2004 Aug 1;298(1):218-28.

*Lukas J, Mazna P, Valenta T, Doubravska L, Pospichalova V, Vojtechova M, Fafilek B, Ivanek R, Plachy J, Novak J, Korinek V: Dazap2 modulates transcription driven by the Wnt effector TCF-4. Nucleic Acids Res. 2009 Mar 20. [Epub ahead of print]

*Smertenko A, Blume Y, Viklický V, Dráber P: Exposure of tubulin structural domains in Nicotiana tabacum microtubules probed by monoclonal antibodies. Eur J Cell Biol. 1997 Feb;72(2):104-12.

Unless indicated otherwise, all products are For Research Use Only and not for diagnostic or therapeutic use. Not for resale or transfer either as a stand-alone product or as a component of another product without written consent of EXBIO. EXBIO will not be held responsible for patent infringement or other violations that may occur with the use of our products. All orders are accepted subject to EXBIO's term and conditions which are available at www.exbio.cz.

For laboratory research only, not for drug, diagnostic or other use.