γ Tubulin (D-10): sc-17788



The Power to Question

BACKGROUND

Tubulin is a major cytoskeleton component that has five distinct forms, designated $\alpha,\,\beta,\,\gamma,\,\delta$ and ϵ Tubulin. α and β Tubulins form heterodimers which multimerize to form a microtubule filament. Multiple β Tubulin isoforms ($\beta1,\,\beta2,\,\beta3,\,\beta4,\,\beta5,\,\beta6$ and $\beta8$) have been characterized and are expressed in mammalian tissues. $\beta1$ and $\beta4$ are present throughout the cytosol, $\beta2$ is present in the nuclei and nucleoplasm, and $\beta3$ is a neuron-specific cytoskeletal protein. γ Tubulin forms the gammasome, which is required for nucleating microtubule filaments at the centrosome. Both δ Tubulin and ϵ Tubulin are associated with the centrosome. δ Tubulin is a homolog of the $\it{Chlamydomonas}\,\delta$ Tubulin Uni3 and is found in association with the centrioles, whereas ϵ Tubulin localizes to the pericentriolar material. ϵ Tubulin exhibits a cell-cycle-specific pattern of localization, first associating with only the older of the centrosomes in a newly duplicated pair and later associating with both centrosomes.

REFERENCES

- Weisenberg, R. 1981. Invited review: the role of nucleotide triphosphate in Actin and Tubulin assembly and function. Cell Motil. 1: 485-497.
- 2. Burns, R.G. 1991. α -, β -, and γ -Tubulins: sequence comparisons and structural constraints. Cell Motil. Cytoskeleton 20: 181-189.

CHROMOSOMAL LOCATION

Genetic locus: TUBG1/TUBG2 (human) mapping to 17q21.2; Tubg1/Tubg2 (mouse) mapping to 11 D.

SOURCE

 γ Tubulin (D-10) is a mouse monoclonal antibody raised against amino acids 269-451 mapping at the C-terminus of γ Tubulin of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_{2b}$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

 γ Tubulin (D-10) is available conjugated to agarose (sc-17788 AC), 500 $\mu g/0.25$ ml agarose in 1 ml, for IP; to HRP (sc-17788 HRP), 200 $\mu g/ml$, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-17788 PE), fluorescein (sc-17788 FITC), Alexa Fluor® 488 (sc-17788 AF488), Alexa Fluor® 546 (sc-17788 AF546), Alexa Fluor® 594 (sc-17788 AF594) or Alexa Fluor® 647 (sc-17788 AF647), 200 $\mu g/ml$, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-17788 AF680) or Alexa Fluor® 790 (sc-17788 AF790), 200 $\mu g/ml$, for Near-Infrared (NIR) WB, IF and FCM.

In addition, γ Tubulin (D-10) is available conjugated to Alexa Fluor® 405 (sc-17788 AF405, 200 μ g/ml), for IF, IHC(P) and FCM.

Alexa Fluor® is a trademark of Molecular Probes, Inc., Oregon, USA

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

APPLICATIONS

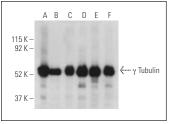
 γ Tubulin (D-10) is recommended for detection of γ Tubulin of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1,000), immunoprecipitation [1-2 μg per 100-500 μg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500), flow cytometry (1 μg per 1 x 10 6 cells) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for γ Tubulin siRNA (h): sc-29322, γ Tubulin siRNA (m): sc-29323, γ Tubulin shRNA Plasmid (h): sc-29322-SH, γ Tubulin shRNA Plasmid (m): sc-29323-SH, γ Tubulin shRNA (h) Lentiviral Particles: sc-29322-V and γ Tubulin shRNA (m) Lentiviral Particles: sc-29323-V.

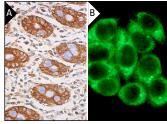
Molecular Weight of γ Tubulin: 50 kDa.

Positive Controls: K-562 whole cell lysate: sc-2203, A-431 whole cell lysate: sc-2201 or Hep G2 cell lysate: sc-2227.

DATA







γ Tubulin (D-10): sc-17788. Immunoperoxidase staining of formalin fixed, paraffin-embedded human rectum tissue showing cytoplasmic and membrane staining of glandular cells (**A**). γ Tubulin (D-10) AF488: sc-17788 AF488. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization (**B**).

SELECT PRODUCT CITATIONS

- Jeong Seo, E., et al. 2004. The role of HPV oncoproteins and cellular factors in maintenance of hTERT expression in cervical carcinoma cells. Gynecol. Oncol. 94: 40-47.
- 2. Barbieri, E., et al. 2018. Targeted enhancer activation by a subunit of the integrator complex. Mol. Cell 71: 103-116.e7.
- 3. Ghaleb, A., et al. 2019. Irradiation induces p53 loss of heterozygosity in breast cancer expressing mutant p53. Commun. Biol. 2: 436.
- 4. Park, Y., et al. 2020. Nonsense-mediated mRNA decay factor UPF1 promotes aggresome formation. Nat. Commun. 11: 3106.
- Senatore, E., et al. 2021. The TBC1D31/praja2 complex controls primary ciliogenesis through PKA-directed OFD1 ubiquitylation. EMBO J. 40: e106503.

RESEARCH USE

For research use only, not for use in diagnostic procedures.