

bsm-33313M**[Primary Antibody]**

Gamma Tubulin Mouse mAb

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— DATASHEET —

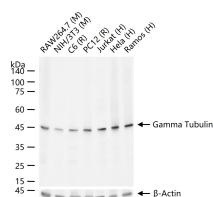
Host: Mouse**Clonality:** Monoclonal**GeneID:** 27175**Target:** Gamma Tubulin**Purification:** affinity purified by Protein G**Concentration:** 1mg/ml

Storage: Size : 50ul/100ul/200ul
0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
Size : 200ug (PBS only)
0.01M PBS
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

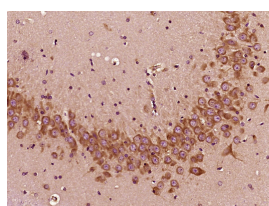
Background: Gamma tubulin, a member of the tubulin superfamily, is a ubiquitous and highly conserved protein within the microtubule organizing centre (MTOC). Gamma tubulin is not a component of microtubules, rather it functions as the microtubule nucleator at the MTOC, is responsible for binding microtubule minus ends and mediating the link between microtubules and the centrosome. By binding to the beta tubulin subunit of the tubulin molecule, it establishes the polarity of a microtubule leaving the alpha tubulin subunit exposed at the positive end.
The abundance of Gamma tubulin is less than 1% of the level of either alpha or beta tubulin. It shares approximately 28-32% identity with alpha tubulin from various organisms and 32-36% identity with beta tubulins. The detection, localization and characterization of proteins involved in microtubule function is fundamental to the understanding of mitosis, meiosis and the microtubule cytoskeleton.

Isotype: IgG**CloneNo.:** 8D11**SWISS:** P23258**Applications:** WB (1:500-5000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (1ug/Test)**Reactivity:** Human, Mouse, Rat**Predicted**
MW.: 50 kDa**Subcellular**
Location: Cytoplasm

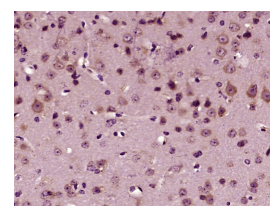
— VALIDATION IMAGES —



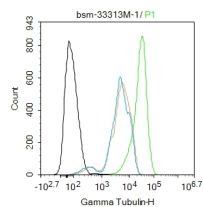
25 ug total protein per lane of various lysates (see on figure) probed with Gamma Tubulin monoclonal antibody, unconjugated (bsm-33313M) at 1:2000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Paraformaldehyde-fixed, paraffin embedded (Rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Gamma Tubulin) Polyclonal Antibody, Unconjugated (bsm-33313M) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Gamma Tubulin) Polyclonal Antibody, Unconjugated (bsm-33313M) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control (black line) :Hela. Primary
 Antibody (green line):Mouse Anti-Gamma
 Tubulin antibody (bsm-33313M)
 Dilution:1ug/Test; Secondary Antibody (white
 blue line) : Goat anti-Mouse IgG-AF488 Dilution:
 0.5ug/Test. Isotype control (orange line) :
 Normal Mouse IgG Protocol The cells were fixed
 with 4% PFA (10min at room temperature)and
 then permeabilized with 90% ice-cold methanol
 for 20 min at -20°C, The cells were then
 incubated in 5%BSA to block non-specific
 protein-protein interactions for 30 min at room
 temperature .Cells stained with Primary
 Antibody for 30 min at room temperature. The
 secondary antibody used for 40 min at room
 temperature. Acquisition of 20,000 events was
 performed.