

beta II Tubulin Antibody

Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP22106a

Specification

beta II Tubulin Antibody - Product Information

Application Primary Accession Other Accession

Reactivity Predicted Host Clonality Isotype WB, FC, IF,E <u>O7TMM9</u> <u>P09203</u>, <u>O9NFZ7</u>, <u>O13885</u>, <u>O4R5B3</u>, <u>P85108</u>, <u>O6B856</u>, <u>O9BVA1</u>, <u>O9CWF2</u>, <u>O3KRE8</u>, <u>P32882</u>, <u>P13602</u>, <u>O9NFZ5</u>, <u>P30156</u>, <u>P20802</u>, <u>O59837</u>, <u>P02554</u> Human, Mouse, Rat Chicken, Monkey, Bovine, Xenopus, Pig Rabbit polyclonal Rabbit IgG

beta II Tubulin Antibody - Additional Information

Gene ID 22151

Other Names Tubulin beta-2A chain, Tubb2a, Tubb2

Target/Specificity

This antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 14-46 amino acids from human.

Dilution WB~~1:8000 FC~~1:25 IF~~1:25 E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

beta II Tubulin Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

beta II Tubulin Antibody - Protein Information



Name Tubb2a

Synonyms Tubb2

Function Tubulin is the major constituent of microtubules, a cylinder consisting of laterally associated linear protofilaments composed of alpha- and beta-tubulin heterodimers. Microtubules grow by the addition of GTP-tubulin dimers to the microtubule end, where a stabilizing cap forms. Below the cap, tubulin dimers are in GDP-bound state, owing to GTPase activity of alpha-tubulin.

Cellular Location Cytoplasm, cytoskeleton.

beta II Tubulin Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- <u>Western Blot</u>
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- <u>Cell Culture</u>

beta II Tubulin Antibody - Images



Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized C2C12 (mouse myoblast cell line) cells labeling beta II Tubulin with AP22106a at 1/25 dilution, followed by Dylight® 488-conjugated goat anti-rabbit IgG (NK179883) secondary antibody at 1/200 dilution (green). Immunofluorescence image showing cytoplasm staining on C2C12 cell line. The nuclear counter stain is DAPI (blue).



All lanes : Anti-beta II Tubulin at 1:8000 dilution Lane 1: mouse brain lysate Lane 2: Hela whole cell lysate Lane 3: Jurkat whole cell lysate Lane 4: PC-12 whole cell lysate Lane 5: NIH/3T3 whole cell lysate Lysates/proteins at 20 μ g per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 50 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



Overlay histogram showing NIH/3T3 cells stained with AP22106a (green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (AP22106a, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Rabbit lgG, **DyLight**® 488 Conjugated Highly Cross-Adsorbed(OH191631) at 1/200 dilution for 40 min at 37°C. Isotype control antibody (blue line) was rabbit IgG $(1\mu g/1 \times 10^{6} \text{ cells})$ used under the same conditions. Acquisition of >10, 000 events was performed.

beta II Tubulin Antibody - Background

Tubulin is the major constituent of microtubules. It binds two moles of GTP, one at an exchangeable site on the beta chain and one at a non-exchangeable site on the alpha chain (By similarity).

beta II Tubulin Antibody - References

Carninci P., et al. Science 309:1559-1563(2005). Lubec G., et al. Submitted (JAN-2009) to UniProtKB.



Janke C., et al. Science 308:1758-1762(2005). Rogowski K., et al. Cell 137:1076-1087(2009). Yoshida K., et al. Biochem. Biophys. Res. Commun. 389:506-511(2009).