

Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody

Catalog # AN2000

Specification

Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody - Product Information

Primary Accession <u>O9BOE3</u>

Reactivity Bovine, Chicken, Drosophila, C.Elegans

Host Rabbi

Clonality Rabbit Polyclonal

Isotype IgG
Calculated MW 49895

Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody - Additional Information

Gene ID 84790

Other Names

alpha 6 Tubulin, Tubulin A6

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

Anti- α 6-Tubulin (Ser-165), Phosphospecific Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Shipping

Blue Ice

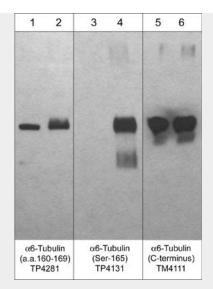
Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody - Protocols

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

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody - Images





Western blot analysis of 250 ng/lane of a1-Tubulin unphosphorylated (lanes 1, 3, & 5) or phosphorylated at Ser-165 with PKC α (lanes 2, 4, & 6). The blots were probed with anti- α 6-Tubulin (a.a. 160-169) (TP4281; lanes 1 & 2), anti- α 6-Tubulin (Ser-165) (TP4131; lanes 3 & 4), and anti- α -Tubulin (C-terminus) (TM4111; lanes 5 & 6).

Anti-α6-Tubulin (Ser-165), Phosphospecific Antibody - Background

Microtubules (MTs) are cytoskeletal elements that play an essential role in cell division and cytoplasmic organization. MTs are dynamic polymers of a/ β -Tubulin heterodimers. At least two populations of MTs, called dynamic and stable according to their rates of turnover, are readily distinguishable in cells. The proteins associated with MTs (MAPs) are among the best-known factors that regulate MT dynamics and stability. In addition, a variety of different post-translational modifications may also regulate MT dynamics and stability. Phosphorylation is one of these modifications and it can occur on serine, threonine, and tyrosine residues in α - and β -Tubulin isoforms. Multiple kinases can phosphorylate Ser-444 at the C-terminus of β III-Tubulin in vitro, and unphosphorylated Ser-444 may be an early marker for cells of neuronal lineage. Cdk1 can phosphorylate Ser-172 in β -Tubulin during mitosis and this may impair tubulin incorporation into microtubules. In α -tubulin, PKC can phosphorylate Ser-165 leading to increased cell motility in human breast cells.