

## Anti-βIII-Tubulin Antibody

Catalog # AN2005

## **Specification**

#### Anti-βIII-Tubulin Antibody - Product Information

Application WB
Primary Accession Q13509
Reactivity Bovine
Host Rabbit

Clonality Rabbit Polyclonal

Isotype IgG
Calculated MW 50433

## Anti-βIII-Tubulin Antibody - Additional Information

Gene ID **10381** 

**Other Names** 

TUBB3

**Dilution** 

WB~~1:1000

#### 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- $\beta$ III-Tubulin Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

#### **Shipping**

Blue Ice

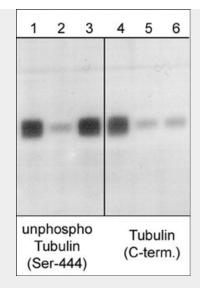
# Anti-βIII-Tubulin Antibody - Protocols

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

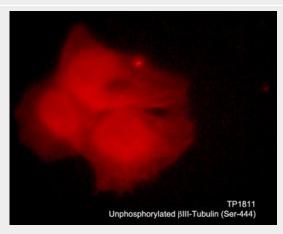
- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

# Anti-βIII-Tubulin Antibody - Images





Western blot analysis of mouse brain. The blot was probed with anti-unphosphorylated  $\beta$ III-Tubulin (Ser-444) (lanes 1-3) and anti- $\beta$ III-Tubulin (C-terminus) (lanes 4-6) polyclonal antibodies. Both antibodies were used in the presence of unphosphorylated  $\beta$ III-Tublin (Ser-444) peptide (lanes 2 & 5; TX1815) and phospho- $\beta$ III-Tublin (Ser-444) peptide (lanes 3 & 6; TX1695).



Immunocytochemical labeling of  $\beta$ -tubulin in aldehyde fixed and NP-40 permeabilized human NCI-H1299 lung carcinoma cells. The cells were labeled with rabbit polyclonal anti-unphosphorylated  $\beta$ -Tubulin (TP1811). The antibody was detected using goat anti-rabbit DyLight® 594.

#### Anti-BIII-Tubulin Antibody - Background

Microtubules (MTs) are cytoskeletal elements that play an essential role in cell division and cytoplasmic organization. MTs are dynamic polymers of  $\alpha/\beta$ -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  $\beta$ -Tubulin isoforms. Multiple kinases can phosphorylate Ser-444 at the C-terminus of  $\beta$ III-Tubulin in vitro. Unphosphorylated Ser-444 in  $\beta$ III-Tubulin is an early marker for cells of neuronal lineage, while phosphorylation of Ser-444 is upregulated after neuronal maturation and may preferentially occur in assembled MTs. By contrast, Cdk1 phosphorylation of Ser-172 in  $\beta$ -Tubulin occurs in mitotic cells and may impair tubulin incorporation into microtubules.