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Product datasheet for TA389228

TUBA1A Mouse Antibody [Clone ID: DM1A]

Product data:

Product Type:	Primary Antibodies
Clone Name:	DM1A
Applications:	ICC, IHC, IP, WB
Recommended Dilution:	WB : 1:500 ICC : 1:200
Reactivity:	Human, Rat, Mouse
Host:	Mouse
lsotype:	lgG1
Immunogen:	Clone DM1A was generated from purified chick brain tubulin. The antibody recognizes an epitope within the C-terminal end of α -tubulin isoforms. The epitope is highly conserved in α -tubulin isoforms from most vertebrate species.
Specificity:	The antibody detects a 55 kDa* protein corresponding to the molecular mass of α -Tubulin on SDS-PAGE immunoblots of human, rat, and mouse cells and tissues.
Formulation:	PBS + 1 mg/ml BSA, 0.05% NaN3 and 50% glycerol
Concentration:	lot specific
Purification:	Protein A Purified
Conjugation:	Unconjugated
Storage:	Storage at -20°C is recommended, as aliquots may be taken without freeze/thawing due to presence of 50% glycerol. Stable for at least 1 year at -20°C.
Stability:	After date of receipt, stable for at least 1 year at -20°C.
Predicted Protein Size:	55
Database Link:	<u>Q71U36</u>



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GRIGENE TUBA1A Mouse Antibody [Clone ID: DM1A] – TA389228

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 βIII-
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.

Note: Protein G purified tissue culture supernatant.

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