

## a-tubulin Monoclonal Antibody

Catalog No: YM3115

**Reactivity:** Zebrafish

**Applications:** WB

Target: Tubulin a

**Fields:** >>Phagosome;>>Gap junction;>>Alzheimer disease;>>Parkinson

disease;>>Amyotrophic lateral sclerosis;>>Huntington disease;>>Prion disease;>>Pathways of neurodegeneration - multiple diseases;>>Pathogenic

Escherichia coli infection;>>Salmonella infection

Gene Name: TUBA1A

Protein Name: Tubulin alpha-1A chain

**Human Gene Id:** 7846/10376

**Human Swiss Prot** 

.

No:

Mouse Gene ld: 22142/22143

**Rat Gene Id:** 64158/500929

Rat Swiss Prot No: P68370/Q6P9V9

Immunogen: Recombinant Protein of Tubulin alpha-1A chain

Q71U36/P68363

**Specificity:** The antibody detects Zebrafish endogenous α-tubulin protein.

**Formulation:** PBS, pH 7.4, containing 0.5%BSA, 0.02% sodium azide as Preservative and

50% Glycerol.

Source: Monoclonal, Mouse

**Dilution:** WB 1:500-10000

1/3



**Purification:** The antibody was affinity-purified from mouse ascites by affinity-

chromatography using specific immunogen.

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 52kD

**Background :** Microtubules of the eukaryotic cytoskeleton perform essential and diverse

functions and are composed of a heterodimer of alpha and beta tubulins. The genes encoding these microtubule constituents belong to the tubulin superfamily, which is composed of six distinct families. Genes from the alpha, beta and gamma tubulin families are found in all eukaryotes. The alpha and beta tubulins represent the major components of microtubules, while gamma tubulin plays a critical role in the nucleation of microtubule assembly. There are multiple alpha and beta tubulin genes, which are highly conserved among species. This gene encodes alpha tubulin and is highly similar to the mouse and rat Tuba1 genes. Northern blotting studies have shown that the gene expression is predominantly found in morphologically differentiated neurologic cells. This gene is one of three

alpha-tubulin genes in a cluster on chromosome 12q.

**Function:** disease:Defects in TUBA1A are the cause of lissencephaly type 3 (LIS3)

[MIM:611603]. LIS is characterized by a smooth brain surface due to the absence (agyria) or reduction (pachygyria) of surface convolutions. It is often associated with psychomotor retardation and seizures. LIS3 features include agyria or pachygyria or laminar heterotopia, severe mental retardation, motor delay, variable presence of seizures, and abnormalities of corpus callosum,

hippocampus, cerebellar vermis and brainstem.,function: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 alphachain.,PTM:Undergoes a tyrosination/detyrosination cycle, the cyclic removal and

re-addition of a C-terminal tyrosine residue by the enzymes tubulin tyrosine

carboxypeptidase (TTCP) and tubulin tyrosine ligase (TTL), resp

Subcellular Location:

Cytoplasm, cytoskeleton.

**Expression:** Expressed at a high level in fetal brain.

Tag: orthogonal

**Sort**: 1272

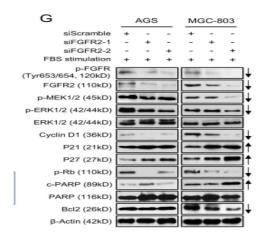
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Host: Mouse

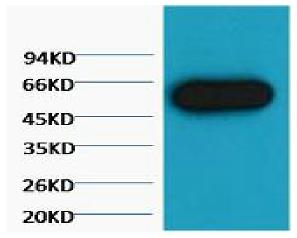
Modifications: Unmodified



## **Products Images**



Zhang, J., Wong, C.C., Leung, K.T. et al. FGF18–FGFR2 signaling triggers the activation of c-Jun–YAP1 axis to promote carcinogenesis in a subgroup of gastric cancer patients and indicates translational potential. Oncogene 39, 6647–6663 (2020).



Western blot analysis of Zebrafish skeletal muscle, (Zebrafish Specific) diluted at 1:5000.