For Research Use Only

## Acetyl-Tubulin (Lys40) Monoclonal antibody, PBS Only

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**Purification Method:** 

Protein G purification

CloneNo.:

7E5H8

Catalog Number:66200-1-PBS

**Basic Information** 

Catalog Number: 66200-1-PBS

Concentration:

1000 μg/ml Source: Mouse

Isotype: lgG1

GenBank Accession Number: NM\_006009

GeneID (NCBI):

7846

**UNIPROT ID:** Q71U36 Full Name:

tubulin, alpha 1a Calculated MW:

52 kDa

Observed MW: 50-55 kDa

**Applications** 

**Tested Applications:** 

WB, IHC, IF/ICC, IF-P, ELISA, Indirect ELISA

Species Specificity:

human, mouse, rat, pig, canine

## **Background Information**

Tubulin, composed of heterodimers of alpha and beta tubulin, is the mainly component of microtubules which play important roles in cell motility, mitosis, and intracellular vesicle transport. Both alpha and beta tubulin undergo several posttranslational modifications such as polyglutamylation and acetylation/deacetylation. Tubulin acetylation occurs on lysine-40 at the N-terminal of alpha tubulin and is conserved across species. The histone deacetylase HDAC6 and SIRT2 has been identified as tubulin deacetylases. Reversible acetylation of alpha tubulin may be implicated in regulating microtubule stability, cell motility, and axon regeneration. The level of acetylated tubulin has been linked to epithelial malignancies and sensitivity to chemotherapy. In addition, acetylated tubulin has been widely used as a marker for primary cilia. This antibody is specific to the acetylated tubulin; it does not recognize non-acetylated tubulin. (24268707, 23881549)

Storage

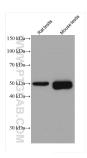
Storage:

Store at -80°C.

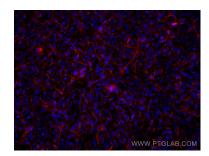
The product is shipped with ice packs. Upon receipt, store it immediately at -80°C

PBS only, pH7.3

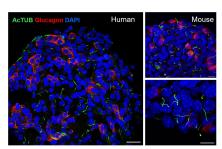
## Selected Validation Data



Various lysates were subjected to SDS PAGE followed by western blot with 66200-1-1g (Acetyl-Tubulin (Lys40) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. This data was developed using the same antibody clone with 66200-1-PBS in a different storage buffer formulation

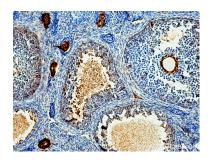


Immunofluorescent analysis of (4% PFA) fixed hTERT-RPE1 cells using Acetyl-Tubulin (Lys40) antibody (66200-1-Ig, Clone: 7E5H8) at dilution of 1:1000 and Multi-rAb Coralite® Plus 594-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO. RGAM004). This data was developed using the same antibody clone with 66200-1-PBS in a different storage buffer formulation.

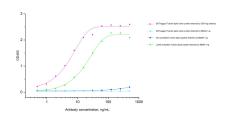


(left) Confocal image of a single-plane cross-section of a healthy human islet, showing primary cilia on both alpha cells (glucagon, red) and non-alpha cells (non-red). Acetylated alpha tubulin, cilia (green, Cat. No 66200-1-lg, 1:400), nuclei (blue), scale 20 μm. (right) Primary cilia in wildtype 86 mouse islets: acetylated alpha tubulin (green, Cat. No 66200-1-lg, 1:400), glucagon (red), nuclei (blue), single-plane images, scales 10 μm. (Fig from Dr. Jing Hughes) This data was

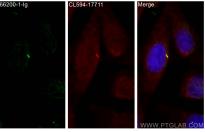




Immunohistochemical analysis of paraffinembedded mouse ovary tissue slide using 66200-1-Ig (Acetyl-Tubulin (Lys40) antibody) at dilution of 1:2000 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0). This data was developed using the same antibody clone with 66200-1-PBS in a different storage buffer formulation.



Specificity validation by indirect ELISA.
Indirect ELISA was carried out by coating
E.coli expressed GST-tagged Tubulin alpha
(70 ng/well), non-acetylated Tubulin alpha,
and Lys40 acetylated Tubulin alpha (with
excessive amount) followed by blocking and
adding serial diluted GST tag primary
antibody (Cat.NO. 66001-2-lg) and Acetylated
Tubuliln antibody 66200-1-lg respectively.
HRP-Goat anti-mouse secondary antibody was
used for detection. Signal was developed with



Immunofluorescent analysis of (4% PFA) fixed MDCK cells using Acetyl-Tubulin (Lys40) antibody (66200-1-lg, Clone: 7E5H8) at dilution of 1:400 and CoraLite@488-Conjugated Goat Anti-Mouse IgG(H+L) (SA00013-1), CoraLite®594 ARL13B antibody (CL594-17711, red). This data was developed using the same antibody clone with 66200-1-PBS in a different storage buffer formulation.