# ANTIBODIES ONLINE

# Datasheet for ABIN7505839 anti-alpha Tubulin antibody

3 Images



## Overview

Quantity:	0.1 mg
Target:	alpha Tubulin (TUBA1)
Reactivity:	Human, Mouse, Rat, Saccharomyces cerevisiae, Mammalian, Avian
Host:	Rat
Clonality:	Monoclonal
Conjugate:	This alpha Tubulin antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Immunocytochemistry (ICC), Flow Cytometry (FACS), Immunohistochemistry (Frozen Sections) (IHC (fro))

# Product Details

Purpose:	Anti-alpha-Tubulin Purified
Immunogen:	Yeast tubulin
Clone:	YOL1-34
Isotype:	lgG2a
Specificity:	The rat monoclonal antibody YOL1/34 recognizes an epitope of alpha-tubulin localized between amino acids 414-422. It has higher affinity for fixed microtubules than for native ones.
Purification:	Purified by protein-G affinity chromatography.
Target Details	

Target:

alpha Tubulin (TUBA1)

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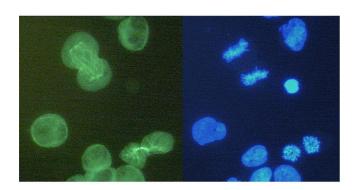
Target Details	
Alternative Name:	alpha-Tubulin (TUBA1 Products)
Background:	Tubulin alpha 1,The microtubules are intracellular dynamic polymers made up of evolutionarily
	conserved polymorphic alpha/beta-tubulin heterodimers and a large number of microtubule-
	associated proteins (MAPs). The microtubules consist of 13 protofilaments and have an outer
	diameter 25 nm. Microtubules have their intrinsic polarity, highly dynamic plus ends and less
	dynamic minus ends. Microtubules are required for vital processes in eukaryotic cells including
	mitosis, meiosis, maintenance of cell shape and intracellular transport. Microtubules are also
	necessary for movement of cells by means of flagella and cilia. In mammalian tissue culture
	cells microtubules have their minus ends anchored in microtubule organizing centers (MTOCs).
	The GTP (guanosintriphosphate) molecule is an essential for tubulin heterodimer to associate
	with other heterodimers to form microtubule. In vivo, microtubule dynamics vary considerably.
	Microtubule polymerization is reversible and a populations of microtubules in cells are on their
	minus ends either growing or shortening –, this phenomenon is called dynamic instability of
	microtubules. On a practical level, microtubules can easily be stabilized by the addition of non-
	hydrolysable analogues of GTP (eg. GMPPCP) or more commonly by anti-cancer drugs such as
	Taxol. Taxol stabilizes microtubules at room temperature for many hours. Using limited
	proteolysis by enzymes both tubulin subunits can be divided into N-terminal and C-terminal
	structural domains. The alpha-tubulin (relative molecular weight around 50 kDa) is globular
	protein that exists in cells as part of soluble alpha/beta-tubulin dimer or it is polymerized into
	microtubules. In different species it is coded by multiple tubulin genes that form tubulin classes
	(in human 6 genes). Expressed tubulin genes are named tubulin isotypes. Some of the tubulin
	isotypes are expressed ubiquitously, while some have more restricted tissue expression. Alpha-
	tubulin is also subject of numerous post-translational modifications. Tubulin isotypes and their
	posttranslational modifications are responsible for multiple tubulin charge variants - tubulin
	isoforms. Heterogeneity of alpha-tubulin is concentrated in C-terminal structural domain.,TUBA
Gene ID:	7277
UniProt:	Q71U36
Pathways:	Microtubule Dynamics
Application Details	
Application Notes:	Immunocytochemistry: Recommended dilution 4-8 µg/mL.
	Western blotting: Recommended dilution 1-2 $\mu$ g/mL, reducing conditions.
Restrictions:	For Research Use only

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## Handling

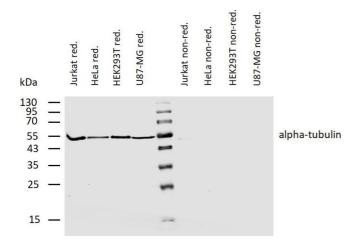
Concentration:	1 mg/mL
Buffer:	Phosphate buffered saline (PBS), pH 7.4, 15 mM sodium azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C
Storage Comment:	Store at 2-8°C. Do not freeze.

#### Images



#### Immunocytochemistry

**Image 1.** Immunocytochemistry staining of alpha-tubulin in K562 cells using purified rat monoclonal antibody YOL1/34 (concentration in sample 6  $\mu$ g/mL, DAR FITC, left picture) vs. Hoechst 34580 nuclear staining (right picture).

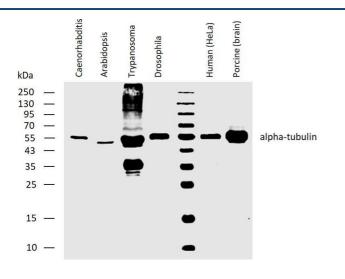


#### Western Blotting

**Image 2.** Western blotting analysis of human alpha-tubulin using rat monoclonal antibody YOL1/34 on lysates of various cell lines under reducing and non-reducing conditions. Nitrocellulose membrane was probed with 2 µ g/mL of rat anti-alpha-tubulin monoclonal antibody followed by IRDye800-conjugated anti-rat secondary antibody. A specific band was detected for alpha-tubulin at approximately 54 kDa.

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Images



#### Western Blotting

**Image 3.** Western blotting analysis of human alpha-tubulin using rat monoclonal antibody YOL1/34 on lysates of Caenorhabditis elegans, Arabidopsis thaliana, Trypanosoma brucei, Drosophila melanogaster, human HeLa cell line, and porcine brain, all under reducing conditions. Each lane contains 20  $\mu$ g of total protein. Nitrocellulose membrane was probed with 2  $\mu$ g/mL of rat anti-alpha-tubulin monoclonal antibody followed by IRDye800-conjugated antirat secondary antibody.

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