

Product Data Sheet

Anti-Beta-tubulin Antibody

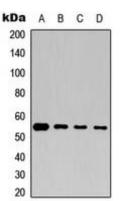
Catalog #	Source	Reactivity	Applications	
CPA4302	Rabbit	H, M, R	WB, IH, IF/IC	
Description	R	abbit polyclonal antibody	to Beta-tubulin	
Immunogen	К	LH-conjugated synthetic p	eptide encompassing a sequence within the C-term	
	re	egion of human Beta-tubu	in. The exact sequence is proprietary.	
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.	
Specificity	R	ecognizes endogenous lev	els of Beta-tubulin protein.	
Clonality	Р	olyclonal		
Form	Li	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	а	nd 0.01% sodium azide.		
Dilution	V	VB (1/500 - 1/1000), IH (1/10	00 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	Т	UBB3		
Alternative N	ames T	UBB4; Tubulin beta-3 chai	n; Tubulin beta-4 chain; Tubulin beta-III	
Entrez Gene	1	.0381 (Human); 22152 (Mc	ouse); 246118 (Rat)	
SwissProt	Q	213509 (Human); Q9ERD7	(Mouse); Q4QRB4 (Rat)	
Storage/Stabi	i lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	fr	reeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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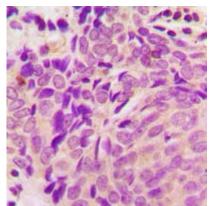




For research purposes only, not for human use

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Western blot analysis of Beta-tubulin expression in Jurkat (A), K562 (B), mouse brain (C), rat brain (D) whole cell lysates.



Immunohistochemical analysis of Beta-tubulin staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Immunofluorescent analysis of Beta-tubulin staining in Jurkat cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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