

## **Product Data Sheet**

## **Anti-MAST3 Antibody**

Catalog #	Source	Reactivity	Applications
CPA2561	Rabbit	H, M, R, B, D, P	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to	MAST3
Immunogen	KLH-	conjugated synthetic per	otide encompassing a sequence within the C-term
	regio	n of human MAST3. The	exact sequence is proprietary.
Purification	Thea	antibody was purified by	immunogen affinity chromatography.
Specificity	Reco	gnizes endogenous level	s of MAST3 protein.
Clonality	Poly	lonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB (	1/500 - 1/1000), IH (1/100	) - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	MAS	ГЗ	
Alternative Na	ames KIAA	0561; Microtubule-asso	ciated serine/threonine-protein kinase 3
Entrez Gene	2303	1 (Human); 546071 (Mo	use)
SwissProt	O603	807 (Human); Q3U214 (N	/louse)
Storage/Stabi	lity Ship	oed at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid
	freez	e/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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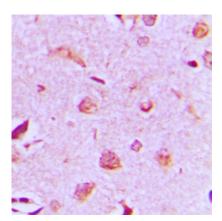
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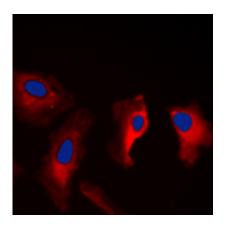
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Western blot analysis of MAST3 expression in HepG2 (A), rat kidney (B) whole cell lysates. (Predicted band size: 143 kD; Observed band size: 143 kD)



Immunohistochemical analysis of MAST3 staining in human brain 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 conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of MAST3 staining in HepG2 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 humidified 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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