

Product Data Sheet

Anti-TUFM Antibody

Catalog #	Source	Reactivity	Applications
CPA5007	Rabbit	H, M, R, B, D, Mk, P	WB, IH
Description	Ra	abbit polyclonal antibody to	TUFM
Immunogen	KL	H-conjugated synthetic pep.	tide encompassing a sequence within the center
	re	gion of human TUFM. The e	xact sequence is proprietary.
Purification	Th	ne antibody was purified by i	mmunogen affinity chromatography.
Specificity	Re	ecognizes endogenous levels	of TUFM protein.
Clonality	Pc	olyclonal	
Conjugation			
Form	Lic	quid in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	nd 0.01% sodium azide.	
Dilution	W	/B (1/500 - 1/1000), IH (1/100	- 1/200)
Gene Symbol	TU	JFM	
Alternative Na	ames Ele	ongation factor Tu mitochor	drial; EF-Tu; P43
Entrez Gene	72	284 (Human); 233870 (Mous	e); 293481 (Rat)
SwissProt	P4	19411 (Human); Q8BFR5 (M	ouse); P85834 (Rat)
Storage/Stabi	lity Sh	nipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	fre	eeze/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

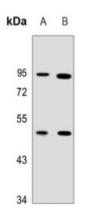
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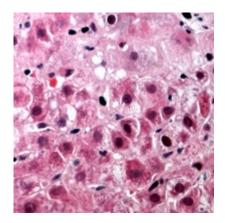
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For research purposes only, not for human use

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Western blot analysis of TUFM expression in Jurkat (A), Hela (B) whole cell lysates. (Predicted band size: 49 kD; Observed band size: 50 kD)



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

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