

Product Data Sheet

Anti-MATK Antibody

Catalog #	Source	Reactivity	Applications		
CQA3204	Rabbit	H, M, R	WB, IH		
Description	R	abbit polyclonal antibody t	o MATK		
Immunogen	R	ecombinant full length pro	tein of human MATK		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	els of MATK protein.		
Clonality		Polyclonal			
Conjugation					
Form	Li	iquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	а	nd 0.01% sodium azide.			
Dilution	W	VB (1/500 - 1/2000), IH (1/50	- 1/200)		
Gene Symbol	N	ЛАТК			
Alternative Na	ames C	TK; HYL; Megakaryocyte-as	sociated tyrosine-protein kinase; CSK homologous		
	ki	inase; CHK; Hematopoietic	consensus tyrosine-lacking kinase; Protein kinase HYL;		
	Ţ	yrosine-protein kinase CTK			
Entrez Gene	4	145 (Human); 17179 (Mou	se); 60450 (Rat)		
SwissProt	P	42679 (Human); P41242 (N	1ouse); P41243 (Rat)		
Storage/Stabi	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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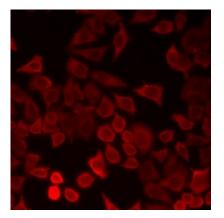
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Western blot analysis of MATK expression in MCF7 (A), mouse brain (B) whole cell lysates. (Predicted band size: 51; 56 kD; Observed band size: 56 kD)



Immunohistochemical analysis of MATK 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 conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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