

Product Data Sheet

Anti-MATK Antibody

Catalog #	Source	e Reactivity	Applications	
CPA6616	Rabbit	: H	WB, IH	
Description		Rabbit polyclonal antibody	to MATK	
Immunogen		KLH-conjugated synthetic p	eptide encompassing a sequence within the center	
		region of human MATK. The	e exact sequence is proprietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous lev	els of MATK protein.	
Clonality Po		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/5) - 1/200)	
Gene Symbol		MATK		
Alternative Na	ames	CTK; HYL; Megakaryocyte-a	ssociated tyrosine-protein kinase; CSK homologous	
		kinase; CHK; Hematopoietio	consensus tyrosine-lacking kinase; Protein kinase HYL;	
		Tyrosine-protein kinase CT		
Entrez Gene		4145 (Human)		
SwissProt		P42679 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
		freeze/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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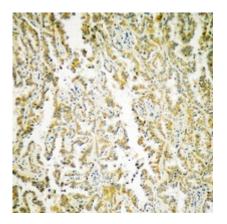
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For research purposes only, not for human use

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Western blot analysis of MATK expression in K562 (A), MCF7 (B) whole cell lysates. (Predicted band size: 56; 51 kD; Observed band size: 56; 51 kD)



Immunohistochemical analysis of MATK staining in human lung 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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