

## **Product Data Sheet**

### **Anti-ZAK Antibody**

Catalog #	Source	e Reactivity	Applications		
CPA5545	Rabbit	: H	WB, IH		
Description		Rabbit polyclonal antibody t	o ZAK		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human ZAK. The e	xact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of ZAK protein.		
Clonality		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 (1/50	) - 1/100)		
Gene Symbol		MLTK			
Alternative Names		MLTK; Mitogen-activated protein kinase kinase kinase MLT; Human cervical cancer			
		suppressor gene 4 protein; I	HCCS-4; Leucine zipper- and sterile alpha		
		motif-containing kinase; ML	K-like mitogen-activated protein triple kinase; Mixed		
		lineage kinase-related kinas	e; MLK-related kinase; MRK; Sterile alpha motif- and		
		leucine zipper-containing ki	nase AZK		
Entrez Gene		51776 (Human)			
SwissProt		Q9NYL2 (Human)			
Storage/Stability		Shipped at 4°C. Upon delivery 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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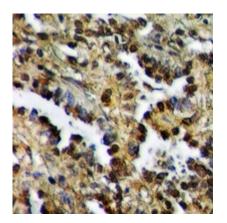
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For research purposes only, not for human use

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Western blot analysis of ZAK expression in HEK293T (A), HCT116 (B) whole cell lysates. (Predicted band size: 91 kD; Observed band size: 91 kD)



Immunohistochemical analysis of ZAK staining in human prostate 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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