

## **Product Data Sheet**

### **Anti-LTK Antibody**

Catalog #	Source	Reactivity	Applications
CPA4970	Rabbit	Н, М	WB, IH, IF/IC
Description	Rat	obit polyclonal antibody	y to LTK
Immunogen	KLH	1-conjugated synthetic	peptide encompassing a sequence within the center
	reg	ion of human LTK. The	exact sequence is proprietary.
Purification	The	e antibody was purified	by immunogen affinity chromatography.
Specificity	Rec	cognizes endogenous le	evels of LTK protein.
Clonality	Pol	yclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WB	s (1/500 - 1/1000), IH (1/	50 - 1/100), IF/IC (1/50 - 1/200)
Gene Symbol	LTK		
Alternative Na	ames TYk	<1; Leukocyte tyrosine l	kinase receptor; Protein tyrosine kinase 1
Entrez Gene	405	58 (Human); 17005 (Mc	ouse)
SwissProt	P29	9376 (Human); P08923	(Mouse)
Storage/Stabi	<b>lity</b> Shi	pped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid
	free	eze/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

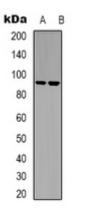
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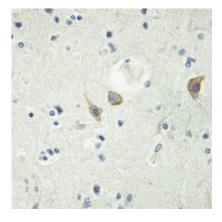
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Western blot analysis of LTK expression in Raji (A), Jurkat (B) whole cell lysates. (Predicted band size: 91 kD; Observed band size: 92 kD)



Immunohistochemical analysis of LTK 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 LTK staining in NIH-3T3 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 hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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