

# **Product Data Sheet**

### **Anti-BTK Antibody**

Catalog #	Source	Reactivity	ŀ	Applications		
CPA5784	Rabbit	H, M, R, B, D, Mk	١	NB, IH		
Description	Ral	bbit polyclonal antibody to	ВТК			
Immunogen	KLł	A-conjugated synthetic pep	tide encompas	sing a sequence within the C-term		
	reg	region of human BTK. The exact sequence is proprietary.				
Purification	The	e antibody was purified by i	immunogen aff	inity chromatography.		
Specificity	Red	cognizes endogenous levels	of BTK protein	l.		
Clonality Polyclonal						
Conjugation						
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	d 0.01% sodium azide.				
Dilution	WE	3 (1/500 - 1/1000), IH (1/50 -	1/100)			
Gene Symbol	BTI	К				
Alternative Na	ames AG	MX1; ATK; BPK; Tyrosine-pr	rotein kinase B <sup>-</sup>	TK; Agammaglobulinemia tyrosine		
	kin	ase; ATK; B-cell progenitor	kinase; BPK; Br	uton tyrosine kinase		
Entrez Gene	695	5 (Human); 12229 (Mouse)				
SwissProt	Q0	6187 (Human); P35991 (Mo	ouse)			
Storage/Stabi	lity Shi	pped at 4°C. Upon delivery	aliquot and sto	ore at -20°C for one year. Avoid		
	fre	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

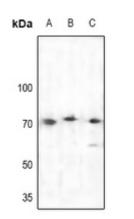
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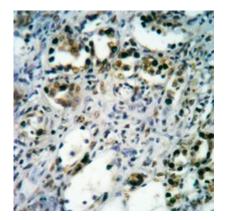


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Western blot analysis of BTK expression in mouse liver (A), rat brain (B), rat liver (C) whole cell lysates. (Predicted band size: 76 kD; Observed band size: 75 kD)



Immunohistochemical analysis of BTK 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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