

## **Product Data Sheet**

## Anti-BIK (Phospho-T33) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3146	Rabbit	н	WB, IH		
Description	Rat	Rabbit polyclonal antibody to BIK (Phospho-T33)			
Immunogen	KLF	I-conjugated synthetic p	hosphopeptide corresponding to residues surrounding		
	Т33	3 of human BIK protein. <sup>-</sup>	The exact sequence is proprietary.		
Purification	The	e antibody was purified b	by immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous lev	els of BIK protein only when phosphorylated at T33.		
Clonality	Pol	yclonal			
Conjugation					
Form	Liqu	uid in 0.42% Potassium p	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	d 0.01% sodium azide.			
Dilution	WB	3 (1/500 - 1/1000), IH (1/5	0 - 1/100)		
Gene Symbol	BIK	,			
Alternative Na	ames NB	K; Bcl-2-interacting killer	; Apoptosis inducer NBK; BIP1; BP4		
Entrez Gene	638	8 (Human)			
SwissProt	Q13	3323 (Human)			
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	ery 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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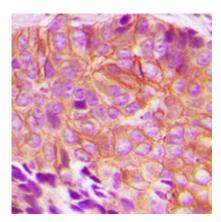
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Western blot analysis of BIK (Phospho-T33) expression in HEK293T (A), HEK293T-EGF (B), SHSY5Y (C), HepG2 (D), A549 (E) whole cell lysates. (Predicted band size: 18 kD; Observed band size: 27 kD)



Immunohistochemical analysis of BIK (Phospho-T33) 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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