

## **Product Data Sheet**

## Anti-BCL2 (Phospho-S70) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3144	Rabbit	Н	WB, IH, IF/IC, IP, FC		
Description	I	Rabbit polyclonal antibody to BCL2 (Phospho-S70)			
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
	9	S70 of human BCL2 proteir	n. The exact sequence is proprietary.		
Purification	-	The antibody was purified	by immunogen affinity chromatography.		
Specificity	<b>pecificity</b> Recognizes endogenous levels of BCL2 protein only when phosphorylated at S70.				
Clonality	I	Polyclonal			
Conjugation					
Form	I	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	v	WB (1/500 - 1/1000), IH (1/5	60 - 1/100), IF/IC (1/50 - 1/200), IP (1/10 - 1/100), FC (1/100 -		
	<u>.</u>	1/200)			
Gene Symbol	I	BCL2			
Alternative Na	ames /	Apoptosis regulator Bcl-2			
Entrez Gene	ļ	596 (Human)			
SwissProt	I	P10415 (Human)			
Storage/Stabi	lity :	Shipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	f	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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kDa

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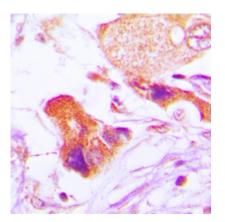
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A B C

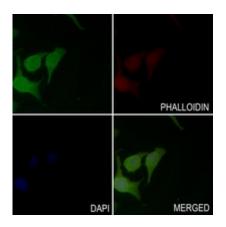
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Western blot analysis of BCL2 (Phospho-S70) expression in Jurkat (A), THP1 (B), A2780 (C) whole cell lysates. (Predicted band size: 26 kD; Observed band size: 28 kD)



Immunohistochemical analysis of BCL2 (Phospho-S70) 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.



Immunofluorescent analysis of BCL2 (Phospho-S70) staining in A549 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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