

Product Data Sheet

Anti-BCL2 Antibody

Catalog #	Source	Reactivity	Applications
CPA9112	Mouse	H, M, R	WB, IH, IF/IC
Description	Mouse monoclonal antibody to BCL2		
Immunogen	KLH-conjugated synthetic peptide encompassing a sequence of human BCL2. The exact sequence is proprietary.		
Purification			
Specificity	Recognizes endogenous levels of BCL2 protein.		
Clonality	Monoclonal		
Conjugation			
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
Dilution	WB (1/1000 - 1/2000), IH (1/200 - 1/500), IF/IC (1/100 - 1/200)		
Gene Symbol	BCL2		
Alternative Names	Apoptosis regulator Bcl-2		
Entrez Gene	596 (Human); 12043 (Mouse); 24224 (Rat)		
SwissProt	P10415 (Human); P10417 (Mouse); P49950 (Rat)		
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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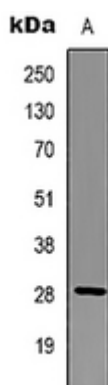
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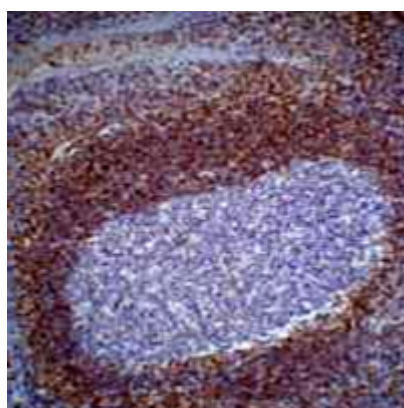
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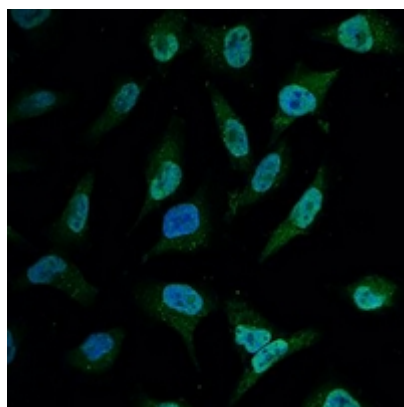
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Western blot analysis of BCL2 expression in HeLa (A) whole cell lysates. (Predicted band size: 26 kD; Observed band size: 28 kD)



Immunohistochemical analysis of BCL2 staining in human tonsil 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 staining in HeLa 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 humidified chamber. Cells were washed with PBST and incubated with a FITC-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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