

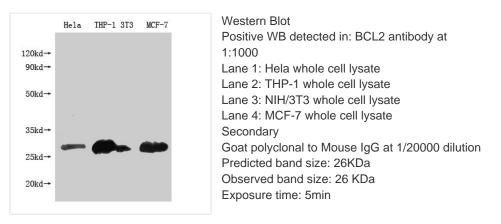
**CUSABIO TECHNOLOGY LLC** 

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## **BCL2 Monoclonal Antibody**

Product Code	CSB-MA002611A0m
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P10415
Immunogen	Recombinant Human poptosis regulator Bcl-2 protein (2-211AA)
Raised In	Mouse
Species Reactivity	Human, Mouse
Tested Applications	ELISA, WB, FC; Recommended dilution: WB: 1:1000-1:5000, FC: 1:50-1:200
Relevance	Suppresses apoptosis in a variety of cell systems including factor-dependent lymphohematopoietic and neural cells. Regulates cell death by controlling the mitochondrial membrane permeability. Appears to function in a feedback loop system with caspases. Inhibits caspase activity either by preventing the release of cytochrome c from the mitochondria and/or by binding to the apoptosis- activating factor (APAF-1). May attenuate inflammation by impairing NLRP1- inflammasome activation, hence CASP1 activation and IL1B release (PubMed:17418785).
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	
	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Purification Method	
	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
Purification Method	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified
Purification Method	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG1
Purification Method Isotype Product Type	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4 >95%, Protein G purified IgG1 Monoclonal Antibody
Purification Method Isotype Product Type Immunogen Species	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4>95%, Protein G purifiedIgG1Monoclonal AntibodyHomo sapiens (Human)
Purification Method Isotype Product Type Immunogen Species Research Area	Constituents: 50% Glycerol, 0.01M PBS, PH 7.4>95%, Protein G purifiedIgG1Monoclonal AntibodyHomo sapiens (Human)Cancer; Cell biology; Metabolism; Signal transduction

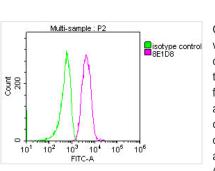
Image



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Overlay Peak curve showing Jurkat cells stained with CSB-MA002611A0m (red line) at 1:100. The cells were incubated in 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody  $(1\mu g/1*106cells)$  for 1h at 4°C. The secondary antibody used was FITC-conjugated Goat Anti-Mouse IgG(H+L) at 1/100 dilution for 30min at 4°C. Isotype control antibody (green line) was mouse IgG1  $(1\mu g/1*106cells)$  used under the same conditions. Acquisition of >10,000 events was performed.