3/23/24, 10:54 AM Revision 3

Revision 5						
Phospho-Bcl-2 (Ser70) (5H2) Rabbit mAb					<b>Ell Signaling</b> C H N O L O G Y <sup>®</sup> 877-616-CELL (2355)	
ίλ Ι					orders@cellsignal.com	
				Support:	877-678-TECH (8324)	
#2827				Web:	info@cellsignal.com cellsignal.com	
3 Trask Lane   Danvers   Massachusetts   01923   USA						
For Research Use Only. Not for	_		0		Future Oraca Id	
Applications: React WB, FC-FP H		<b>MW (kDa):</b> 28	Source/Isotype: Rabbit IgG	UniProt ID: #P10415	Entrez-Gene Id: 596	
Product Usage Information						
	Application			Dilution 1:1000		
	Western Blotting	Flow Cytometry (Fixed/Permeabilized)			1:100 - 1:400	
	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than					
Storage	$0.02\%$ sodium azide. Store at $-20^{\circ}$ C. Do not aliquot the antibody.					
Specificity / Sensitivity	Phospho-Bcl-2 (Ser70) (5H2) Rabbit mAb detects endogenous levels of Bcl-2 only when phosphorylated at serine 70. The antibody does not cross-react with nonphosphorylated Bcl-2 at endogenous levels or with other Bcl-2 family members.					
Source / Purification	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding serine 70 of human Bcl-2.					
Background	Bcl-2 exerts a survival function in response to a wide range of apoptotic stimuli through inhibition of mitochondrial cytochrome c release (1). It has been implicated in modulating mitochondrial calcium homeostasis and proton flux (2). Several phosphorylation sites have been identified within Bcl-2, including Thr56, Ser70, Thr74, and Ser87 (3). It has been suggested that these phosphorylation sites may be targets of the ASK1/MKK7/JNK1 pathway and that phosphorylation of Bcl-2 may be a marker for mitotic events (4,5). Mutation of Bcl-2 at Thr56 or Ser87 inhibits its anti-apoptotic activity during glucocorticoid-induced apoptosis of T lymphocytes (6). Interleukin-3 and JNK-induced Bcl-2 phosphorylation at Ser70 may be required for its enhanced anti-apoptotic functions (7).					
Background References	<ol> <li>Murphy, K.M. et al. (2000) <i>Cell Death Differ</i> 7, 102-11.</li> <li>Zhu, L. et al. (1999) <i>J Biol Chem</i> 274, 33267-73.</li> <li>Maundrell, K. et al. (1997) <i>J Biol Chem</i> 272, 25238-42.</li> <li>Yamamoto, K. et al. (1999) <i>Mol Cell Biol</i> 19, 8469-78.</li> <li>Ling, Y.H. et al. (1998) <i>J Biol Chem</i> 273, 18984-91.</li> <li>Huang, S.T. and Cidlowski, J.A. (2002) <i>FASEB J</i> 16, 825-32.</li> <li>Deng, X. et al. (2001) <i>J Biol Chem</i> 276, 23681-8.</li> </ol>					
Species Reactivity	Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Western Blot Buffer		IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				
Applications Key	WB: Western Blotting F	WB: Western Blotting FC-FP: Flow Cytometry (Fixed/Permeabilized)				
Cross-Reactivity Key	H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected					
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