## Phospho-c-Cbl (Tyr731) Antibody



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## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 120	Source/Isotype: Rabbit	UniProt ID: #P22681	Entrez-Gene Id: 867
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Phospho-c-Cbl (Tyr731) Antibody detects endogenous levels of c-Cbl only when phosphorylated at tyrosine 731. The antibody does not cross-react with related tyrosine-phosphorylated proteins.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr731 of human c-Cbl. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		The c-Cbl proto-oncogene is a ubiquitously expressed cytoplasmic adaptor protein that is especially predominant in hematopoietic cells (1,2). c-Cbl is rapidly tyrosine-phosphorylated in response to stimulation of a variety of cell-surface receptors and becomes associated with a number of intracellular signaling molecules such as protein tyrosine kinases, phosphatidylinositol-3 kinase, Crk, and 14-3-3 proteins (3,4). c-Cbl possesses a highly conserved amino-terminal phosphotyrosine binding domain (TKB) and a C3HC4 RING finger motif. The TKB recognizes phosphorylated tyrosines on activated receptor tyrosine kinases (RTKs) as well as other nonreceptor tyrosine kinases. The RING finger motif recruits ubiquitin-conjugating enzymes. These two domains are primarily responsible for the ubiquitin ligase activity of c-Cbl and downregulation of RTKs (3). Research studies have indicated that in human cancer tissues, c-Cbl is frequently tyrosine-phosphorylated in a tumor-specific manner (5). Phosphorylation of Tyr731 of c-Cbl provides a docking site for downstream signaling components such as p85 and Fyn (6).				
Background References		<ol> <li>Blake, T.J. et al. (1991) Oncogene 6, 653-657.</li> <li>Thien, C.B. and Langdon, W.Y. (1998) Immunol. Cell Biol. 76, 473-482.</li> <li>Christine, B.F. et al. (2001) Nat. Rev. Mol. Cell Biol. 2, 294-307.</li> <li>Feshchenko, E.A. et al. (1998) J. Biol. Chem. 273, 8323-8331.</li> <li>Kamei, T. et al. (2000) Int. J. Oncol. 17, 335-339.</li> <li>Hunter, C. et al. (1999) J. Biol. Chem. 274, 2097-2106.</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** 

W: Western Blotting

**Cross-Reactivity Key** 

H: Human

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