Phospho-c-Cbl (Tyr774) (R3B8) rabbit mAb

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Applications	Detection	Clonality	Isotype
Flow Cytometry	Anti-Rabbit IgG	Monoclonal	Rabbit IgGk

Format: Unconjugated

Cross Reactivity: Predicted to work with mouse, rat and other homologues.

Formulation: 1X PBS, 0.02% NaN3, 50% Glycerol, 0.1% BSA

Preparation: Protein A+G

Reactivity: Human, Mouse

Recommended

Usage: 1µg/mL ? 0.001µg/mL. It is recommended that the reagent be titrated for optimal performance for

each application. See product image legends for additional information.

Immunogen: A synthetic phospho-peptide corresponding to residues surrounding Tyr774 of human phospho c-Cbl

Description: The c-Cbl (Casitas B-lineage Lymphoma) proto-oncogene is a ubiquitously expressed cytoplasmic

adaptor protein that contains multiple functional domains, including an amino-terminal tyrosine kinase-binding (TKB) domain, a RING finger motif, and a proline-rich region. The TKB recognizes phosphorylated tyrosines on activated receptor tyrosine kinases (RTKs) and on other nonreceptor tyrosine kinases, while 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 (1). The proline-rich region contains 14-3-3 protein-binding and SH3 domain-binding motifs. c-Cbl is phosphorylated at Y700, Y731, and Y774 by Syk- and Src-family kinases after the stimulation of some integrins and a wide variety of receptors for immunoglobulins, antigens, hormones, growth factors, and cytokines. Phosphorylated Y774 interacts with the SH2 domain of Crk (1,2). The c-Cbl adapter protein is expressed in the cytoplasm in all tissues, with especially high levels of expression in hematopoietic cells (3,4). Through its many functional sites, c-Cbl plays key roles in the positive and negative regulation of vital cell functions, including T Cell Receptor-mediated cellular immune responses. In human cancer tissues, c-Cbl is frequently tyrosine-phosphorylated in a tumor-specific

manner (5).

References: 1. Christine, B.F. et al. (2001) Nat. Rev. Mol. Cell Biol. 2: 294-307.

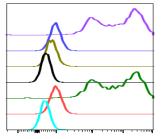
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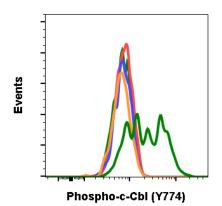
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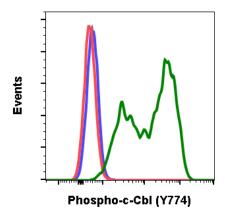


Г	lgG	Treatment	Peptide Block	Median : BL1-A
	R3B8	IFN	Non-phos.	83984
	R3B8	Ctrl	Non-phos.	958
	R3B8	IFN	Phospho	725
	R3B8	Ctrl	Phospho	420
	R3B8	IFN	-	77948
	R3B8	Ctrl	-	980
	2' only	Ctrl	-	376

Peptide blocking flow cytometric analysis of Daudi cells secondary antibody only negative control (light blue) or untreated (red) or treated with IFN? + IL-4 + pervanadate (green) or untreated and blocked with phospho-peptide (black) or treated and blocked with phospho peptide (gold) or untreated and blocked with non-phospho peptide (dark blue) or treated and blocked with non-phospho peptide (purple) using Phospho-c-CbI (Tyr774) antibody CbIY774-R3B8 at 0.1 µg/mL. Cat. #2316.



Flow cytometric analysis of 3T3 cells secondary antibody only negative control (blue) or untreated (gray) or treated with IFN? + IL-4 + pervanadate (orange) using 0.1 μ g/mL isotype control Cat. #2141 or untreated (red) or treated (green) using Phospho-c-Cbl (Tyr774) antibody CblY774-R3B8 at 0.01 μ g/mL. Cat. #2316.



Flow cytometric analysis of Daudi cells secondary antibody only negative control (blue) or untreated (red) or treated with IFN? + IL-4 + pervanadate (green) using Phospho-c-Cbl (Tyr774) antibody CblY774-R3B8 at 0.01 μ g/mL. Cat. #2316.

