

c-Cbl (Phospho Tyr731) rabbit pAb

Catalog No: YP1291

Reactivity: Human; Rat; Mouse;

Applications: WB

Target: Cbl

Fields: >>ErbB signaling pathway;>>Ubiquitin mediated

proteolysis;>>Endocytosis;>>Insulin signaling pathway;>>Bacterial invasion of epithelial cells;>>Pathways in cancer;>>Proteoglycans in cancer;>>Chronic

myeloid leukemia

P22681

P22682

Gene Name: CBL CBL2 RNF55

Protein Name: c-Cbl (Tyr731)

Human Gene Id: 867

Human Swiss Prot

No:

Mouse Gene Id: 12402

Mouse Swiss Prot

No:

Immunogen: Synthesized phosho peptide around human c-Cbl (Tyr731)

Specificity: This antibody detects endogenous levels of Human c-Cbl (phospho-Tyr731)

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:1000-2000

Purification: The antibody was affinity-purified from rabbit serum by affinity-chromatography

using specific immunogen.



Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 100kD

Cell Pathway: ErbB_HER;Ubiquitin mediated

proteolysis;Endocytosis;Jak_STAT;T_Cell_Receptor;Insulin_Receptor;Pathways

in cancer; Chronic myeloid leukemia;

Background: Cbl proto-oncogene(CBL) Homo sapiens This gene is a proto-oncogene that

encodes a RING finger E3 ubiquitin ligase. The encoded protein is one of the enzymes required for targeting substrates for degradation by the proteasome. This protein mediates the transfer of ubiquitin from ubiquitin conjugating enzymes

(E2) to specific substrates. This protein also contains an N-terminal

phosphotyrosine binding domain that allows it to interact with numerous tyrosine-phosphorylated substrates and target them for proteasome degradation. As such it functions as a negative regulator of many signal transduction pathways. This gene has been found to be mutated or translocated in many cancers including acute myeloid leukaemia, and expansion of CGG repeats in the 5' UTR has

cause of Noonan syndrome-like disorder. [provided by RefSeq, Jul 2016],

been associated with Jacobsen syndrome. Mutations in this gene are also the

Function: disease:Can be converted to an oncogenic protein by deletions or mutations that

disturb its ability to down-regulate RTKs.,domain:The N-terminus is composed of the phosphotyrosine binding (PTB) domain, a short linker region and the RING-type zinc finger. The PTB domain, which is also called TKB (tyrosine kinase binding) domain, is composed of three different subdomains: a four-helix bundle (4H), a calcium-binding EF hand and a divergent SH2 domain..domain:The RING-

type zinc finger domain mediates binding to an E2 ubiquitin-conjugating enzyme.,function:Participates in signal transduction in hematopoietic cells.

Adapter protein that functions as a negative regulator of many signaling pathways that start from receptors at the cell surface. Acts as an E3 ubiquitin-protein ligase, which accepts ubiquitin from specific E2 ubiquitin-conjugating enzymes, and then

transfers it to substrates promo

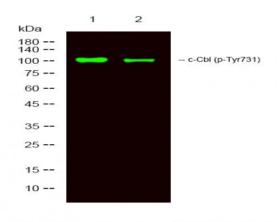
Subcellular Location:

Cytoplasm. Cell membrane. Cell projection, cilium. Golgi apparatus.

Colocalizes with FGFR2 in lipid rafts at the cell membrane.

Expression : Epithelium, T-cell,

Products Images



Western Blot analysis of 1 K562 treated with LPS, 2 K562,using primary antibody at 1:1000 dilution. Secondary antibody(catalog#:RS23920) was diluted at 1:10000