

## LAB (Phospho Tyr136) rabbit pAb

<b>Catalog No :</b>	YP1719
<b>Reactivity :</b>	Human;Mouse;Rat
<b>Applications :</b>	WB
<b>Target :</b>	LAB
<b>Gene Name :</b>	LAT2 LAB NTAL WBS15 WBSCR15 WBSCR5 HSPC046
<b>Protein Name :</b>	LAB (Phospho-Tyr136)
<b>Human Gene Id :</b>	7462
<b>Human Swiss Prot No :</b>	Q9GZY6
<b>Mouse Gene Id :</b>	56743
<b>Mouse Swiss Prot No :</b>	Q9JHL0
<b>Rat Gene Id :</b>	317676
<b>Rat Swiss Prot No :</b>	Q8CGL2
<b>Immunogen :</b>	Synthesized peptide derived from human LAB (Phospho-Tyr136)
<b>Specificity :</b>	This antibody detects endogenous levels of LAB (Phospho-Tyr136) at Human, Mouse,Rat
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Polyclonal, Rabbit,IgG
<b>Dilution :</b>	WB 1:500-2000
<b>Purification :</b>	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 :** 30kD

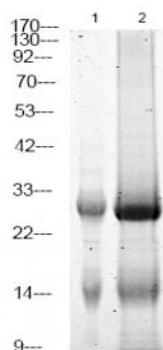
**Background :** This gene is one of the contiguous genes at 7q11.23 commonly deleted in Williams syndrome, a multisystem developmental disorder. This gene consists of at least 14 exons, and its alternative splicing generates 3 transcript variants, all encoding the same protein. [provided by RefSeq, Jul 2008],

**Function :** disease:Defects in LAT2 may be a cause of certain cardiovascular and musculo-skeletal abnormalities observed in Williams-Beuren syndrome (WBS) [MIM:194050]. WBS is a rare developmental disorder. It is a contiguous gene deletion syndrome involving genes from chromosome band 7q11.23.,function:Involved in FCER1 (high affinity immunoglobulin epsilon receptor)-mediated signaling in mast cells. May also be involved in BCR (B-cell antigen receptor)-mediated signaling in B-cells and FCGR1 (high affinity immunoglobulin gamma Fc receptor I)-mediated signaling in myeloid cells. Couples activation of these receptors and their associated kinases with distal intracellular events through the recruitment of GRB2.,PTM:May be polyubiquitinated.,PTM:Phosphorylated on tyrosines following cross-linking of BCR in B-cells, FCGR1 in myeloid cells, or FCER1 in mast cells; which induces the recruitment of GRB2.,s

**Subcellular Location :** Cell membrane ; Single-pass type III membrane protein . Present in lipid rafts.

**Expression :** Highly expressed in spleen, peripheral blood lymphocytes, and germinal centers of lymph nodes. Also expressed in placenta, lung, pancreas and small intestine. Present in B-cells, NK cells and monocytes. Absent from T-cells (at protein level).

## Products Images



Western Blot analysis of 1 HepG2 cell, 2 Serum-free treated ,using primary antibody at 1:1000 dilution. Secondary antibody(catalog#:RS23920) was diluted at 1:10000