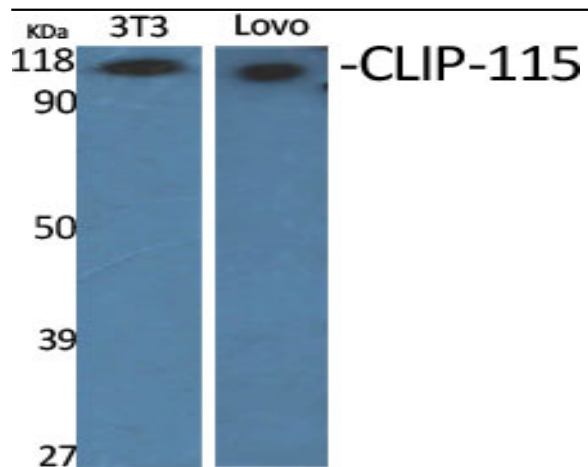


## CLIP-115 Polyclonal Antibody

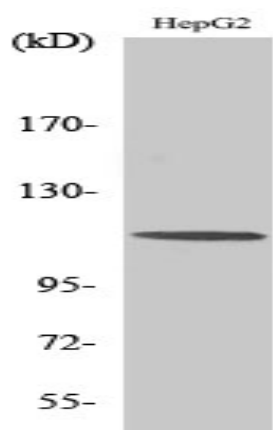
<b>Catalog No :</b>	YT0967
<b>Reactivity :</b>	Human;Mouse;Rat
<b>Applications :</b>	WB;ELISA
<b>Target :</b>	CLIP-115
<b>Gene Name :</b>	CLIP2
<b>Protein Name :</b>	CAP-Gly domain-containing linker protein 2
<b>Human Gene Id :</b>	7461
<b>Human Swiss Prot No :</b>	Q9UDT6
<b>Mouse Gene Id :</b>	269713
<b>Mouse Swiss Prot No :</b>	Q9Z0H8
<b>Rat Gene Id :</b>	29264
<b>Rat Swiss Prot No :</b>	O55156
<b>Immunogen :</b>	The antiserum was produced against synthesized peptide derived from human CLIP2. AA range:997-1046
<b>Specificity :</b>	CLIP-115 Polyclonal Antibody detects endogenous levels of CLIP-115 protein.
<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 - 1:2000. ELISA: 1:5000. Not yet tested in other applications.
<b>Purification :</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.

<b>Concentration :</b>	1 mg/ml
<b>Storage Stability :</b>	-15°C to -25°C/1 year(Do not lower than -25°C)
<b>Observed Band :</b>	120kD
<b>Cell Pathway :</b>	Regulation of Microtubule Dynamics
<b>Background :</b>	<p>The protein encoded by this gene belongs to the family of cytoplasmic linker proteins, which have been proposed to mediate the interaction between specific membranous organelles and microtubules. This protein was found to associate with both microtubules and an organelle called the dendritic lamellar body. This gene is hemizygotously deleted in Williams syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at 7q11.23. Alternative splicing of this gene generates 2 transcript variants. [provided by RefSeq, Jul 2008],</p>
<b>Function :</b>	<p>disease:Haploinsufficiency of CLIP2 may be the 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:Seems to link microtubules to dendritic lamellar body (DLB), a membranous organelle predominantly present in bulbous dendritic appendages of neurons linked by dendrodendritic gap junctions. May operate in the control of brain-specific organelle translocations.,similarity:Contains 2 CAP-Gly domains.,subcellular location:Associated with the cytoskeleton.,subunit:Interacts with CLASP1 and CLASP2.,</p>
<b>Subcellular Location :</b>	Cytoplasm . Cytoplasm, cytoskeleton . Localizes preferentially to the ends of tyrosinated microtubules. .
<b>Expression :</b>	Brain,Clones donated by Kazusa DNA Research Inst.,Epitheliu

## Products Images



Western Blot analysis of various cells using CLIP-115 Polyclonal Antibody



Western Blot analysis of A549 cells using CLIP-115 Polyclonal Antibody