



## CLIP-115 rabbit pAb

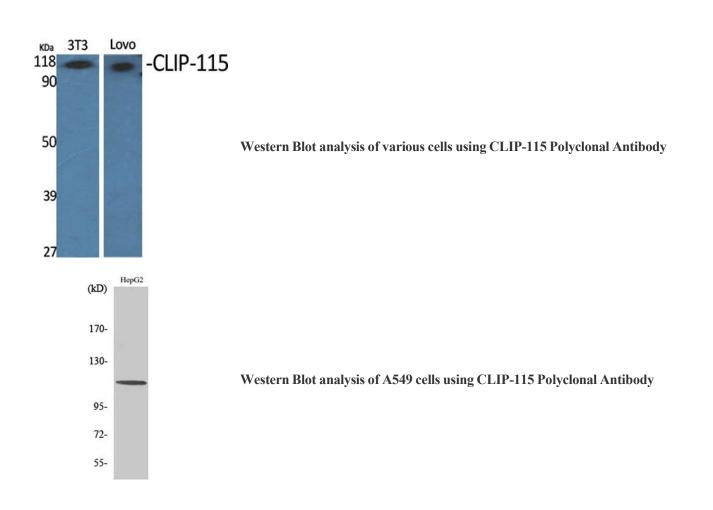
## Cat#: orb764872 (Manual)

For research use only. Not intended for diagnostic use.

| Product Name             | CLIP-115 rabbit pAb   |
|--------------------------|---|
| Host species             | Rabbit  |
| Applications             | WB;ELISA  |
| Species Cross-Reactivity | Human;Mouse;Rat   |
| Recommended dilutions    | Western Blot: 1/500 - 1/2000. ELISA: 1/5000. Not yet tested in other applications.  |
| Immunogen                | The antiserum was produced against synthesized peptide derived from human CLIP2. AA range:997-1046                        |
| Specificity              | CLIP-115 Polyclonal Antibody detects endogenous levels of CLIP-115 protein.   |
| Formulation              | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide  |
| Storage                  | Store at -20°C. Avoid repeated freeze-thaw cycles.  |
| Protein Name             | CAP-Gly domain-containing linker protein 2  |
| Gene Name                | CLIP2   |
| Cellular localization    | Cytoplasm . Cytoplasm, cytoskeleton . Localizes preferentially to the ends of tyrosinated microtubules                    |
| Purification             | The antibody was affinity-purified from rabbit antiserum by affinity-<br>chromatography using epitope-specific immunogen. |
| Clonality                | Polyclonal  |



| Concentration           | 1 mg/ml  |
|-------------------------|--|
| Observed band           | 120kD  |
| Human Gene ID           | 7461   |
| Human Swiss-Prot Number | Q9UDT6   |
| Alternative Names       | CLIP2; CYLN2; KIAA0291; WBSCR3; WBSCR4; WSCR4; CAP-Gly<br>domain-containing linker protein 2; Cytoplasmic linker protein 115; CLIP-<br>115; Cytoplasmic linker protein 2; Williams-Beuren syndrome chromosomal<br>region 3 protein; Williams-Beuren syndro   |
| Background              | 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 hemizygously 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], |





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