

## Product datasheet for RC224491L4V

## OriGene Technologies, Inc.

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## CYLN2 (CLIP2) (NM\_003388) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: CYLN2 (CLIP2) (NM\_003388) Human Tagged ORF Clone Lentiviral Particle

Symbol: CYLN2

Synonyms: CLIP; CLIP-115; CYLN2; WBSCR3; WBSCR4; WSCR3; WSCR4

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM\_003388

ORF Size: 3138 bp

**ORF Nucleotide** 

OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC224491).

Sequence:

**Domains:** 

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeq:** <u>NM 003388.3, NP 003379.3</u>

CAP\_GLY

RefSeq Size: 5563 bp
RefSeq ORF: 3141 bp
Locus ID: 7461
UniProt ID: Q9UDT6
Cytogenetics: 7q11.23

**MW:** 115.6 kDa







## **Gene Summary:**

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]