

## Product datasheet for **TL305216V**

### CRLF2 Human shRNA Lentiviral Particle (Locus ID 64109)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	CRLF2 Human shRNA Lentiviral Particle (Locus ID 64109)
Locus ID:	64109
Synonyms:	CRL2; CRLF2Y; TSLPR
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	CRLF2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">NM_001012288</a> , <a href="#">NM_022148</a> , <a href="#">NR_110830</a> , <a href="#">NM_022148.1</a> , <a href="#">NM_022148.2</a> , <a href="#">NM_022148.3</a> , <a href="#">NM_001012288.1</a> , <a href="#">NM_001012288.2</a> , <a href="#">BC160055</a> , <a href="#">NM_022148.4</a>
UniProt ID:	<a href="#">Q9HC73</a>
Summary:	This gene encodes a member of the type I cytokine receptor family. The encoded protein is a receptor for thymic stromal lymphopoietin (TSLP). Together with the interleukin 7 receptor (IL7R), the encoded protein and TSLP activate STAT3, STAT5, and JAK2 pathways, which control processes such as cell proliferation and development of the hematopoietic system. Rearrangement of this gene with immunoglobulin heavy chain gene (IGH) on chromosome 14, or with P2Y purinoceptor 8 gene (P2RY8) on the same X or Y chromosomes is associated with B-progenitor acute lymphoblastic leukemia (ALL) and Down syndrome ALL. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Sep 2014]
shRNA Design:	These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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**Performance  
Guaranteed:**

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at [techsupport@origene.com](mailto:techsupport@origene.com). Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).