

## Product datasheet for **TR519309**

### Cisd2 Mouse shRNA Plasmid (Locus ID 67006)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	Cisd2 Mouse shRNA Plasmid (Locus ID 67006)
Locus ID:	67006
Synonyms:	1500009M05Rik; 1500026J14Rik; 1500031D15Rik; AI848398; B630006A20Rik; Miner1; Noxp70
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	Cisd2 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 67006). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<a href="#">BC058279</a> , <a href="#">NM_025902</a> , <a href="#">NM_025902.1</a> , <a href="#">NM_025902.2</a> , <a href="#">NM_025902.3</a> , <a href="#">BC025614</a> , <a href="#">BC037182</a> , <a href="#">BC052543</a>
UniProt ID:	<a href="#">Q9CQB5</a>
Summary:	Regulator of autophagy that contributes to antagonize BECN1-mediated cellular autophagy at the endoplasmic reticulum. Participates in the interaction of BCL2 with BECN1 and is required for BCL2-mediated depression of endoplasmic reticulum Ca(2+) stores during autophagy. Contributes to BIK-initiated autophagy, while it is not involved in BIK-dependent activation of caspases. Involved in life span control, probably via its function as regulator of autophagy (By similarity).[UniProtKB/Swiss-Prot Function]
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).