

# Product datasheet for TL511795

## Asxl2 Mouse shRNA Plasmid (Locus ID 75302)

## **Product data:**

#### OriGene Technologies, Inc.

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| Product Type:                | shRNA Plasmids   |
|------------------------------|--|
| Product Name:                | Asxl2 Mouse shRNA Plasmid (Locus ID 75302)   |
| Locus ID:                    | 75302  |
| Synonyms:                    | 4930556B16Rik; mKIAA1685   |
| Vector:                      | pGFP-C-shLenti (TR30023)   |
| E. coli Selection:           | Chloramphenicol (34 ug/ml)   |
| Mammalian Cell<br>Selection: | Puromycin  |
| Format:                      | Lentiviral plasmids  |
| Components:                  | Asxl2 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 75302).<br>5µg purified plasmid DNA per construct<br>29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.   |
| RefSeq:                      | <u>NM 001270988, NM 172421, NM 172421.1, NM 172421.2, NM 172421.3, NM 172421.4, NM 172421.4, NM 172421.5, NM 001270988.1, BC141140, BC166033</u>   |
| UniProt ID:                  | <u>Q8BZ32</u>  |
| Summary:                     | This gene encodes a homolog of the Drosophila Asx gene, which interacts with genes<br>involved in axial patterning. Mice with mutations in this gene display abnormal patterning of<br>the axial skeleton, suggesting a similar function in mice as in Drosophila. This gene may also<br>be involved in bone mineral density, specifically osteoclastogenesis. [provided by RefSeq, Aug<br>2012] |
| 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 <u>techsupport@origene.com</u> .<br>If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .   |



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### **GRIGENE** Asxl2 Mouse shRNA Plasmid (Locus ID 75302) – TL511795

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. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

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