

Product datasheet for TL510718

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OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

Dag1 Mouse shRNA Plasmid (Locus ID 13138)

Product data:

Product Type: shRNA Plasmids

Product Name: Dag1 Mouse shRNA Plasmid (Locus ID 13138)

Locus ID:

D9Wsu13; D9Wsu13e; DG; Dp71; Dp427 Synonyms:

pGFP-C-shLenti (TR30023) Vector:

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: Dag1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 13138).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

BC007150, NM 001276481, NM 001276482, NM 001276485, NM 001276486, NM 001276492, RefSeq:

> NM 001276493, NM 001276494, NM 010017, NM 010017.1, NM 010017.2, NM 010017.3, NM 010017.4, NM 001276481.1, NM 001276482.1, NM 001276485.1, NM 001276486.1,

NM 001276492.1, NM 001276493.1, NM 001276494.1

UniProt ID: Q62165

Summary: This gene encodes dystroglycan, a central component of dystrophin-glycoprotein complex

> that links the extracellular matrix and the cytoskeleton in the skeletal muscle. The encoded preproprotein undergoes O- and N-glycosylation, and proteolytic processing to generate alpha and beta subunits. A complete lack of the encoded protein in mice results in embryonic lethality due to the disorganization of Reichert's membrane. Chimeric mice deficient in the

encoded protein overcome embryonic lethality but develop a progressive muscular

dystrophy. Alternative splicing results in multiple transcript variants, all encoding the same

protein. [provided by RefSeq, Nov 2015]

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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.





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).