

Product datasheet for **TL518960V**

Fkbp1a Mouse shRNA Lentiviral Particle (Locus ID 14225)

Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	Fkbp1a Mouse shRNA Lentiviral Particle (Locus ID 14225)
Locus ID:	14225
Synonyms:	Fkb; Fkbp; Fkbp1; FKBP12
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	Fkbp1a - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 ⁷ TU/ml.
RefSeq:	<u>BC004671</u> , <u>NM_001302077</u> , <u>NM_001302078</u> , <u>NM_001302079</u> , <u>NM_001302080</u> , <u>NM_008019</u> , <u>NR_126058</u> , <u>NM_008019.1</u> , <u>NM_008019.2</u> , <u>NM_008019.3</u> , <u>NM_001302079.1</u> , <u>NM_001302080.1</u> , <u>NM_001302077.1</u> , <u>NM_001302078.1</u>
UniProt ID:	<u>P26883</u>
Summary:	This gene is a member of the immunophilin family. The encoded protein is a cis-trans prolyl isomerase that binds the immunosuppressants FK506 and rapamycin, and is associated with immunoregulation, protein folding, receptor signaling, protein trafficking and T-cell activation. It may modulate the calcium release activity of the ryanodine receptor Ryr1. It also interacts with the type I TGF-beta receptor. Disruption of this gene in mouse causes severe ventricular defects. Pseudogenes of this gene have been defined on chromosomes 4, 10, 14, and 16. Alternative splicing results in multiple transcript variants. [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 techsupport@origene.com . If you need a special design or shRNA sequence, please utilize our custom shRNA service .



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