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Product datasheet for TR313381

DPP6 Human shRNA Plasmid Kit (Locus ID 1804)

Product data:

Product Type:	shRNA Plasmids
Product Name:	DPP6 Human shRNA Plasmid Kit (Locus ID 1804)
Locus ID:	1804
Synonyms:	DPL1; DPPX; MRD33; VF2
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	DPP6 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 1804). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	NM_001039350, NM_001290252, NM_001290253, NM_001936, NM_130797, NM_130797.1, NM_130797.2, NM_130797.3, NM_001936.1, NM_001936.2, NM_001936.3, NM_001936.4, NM_001039350.1, NM_001039350.2, NM_001290253.1, NM_001290252.1, BC035912, BC150304, BM931444, NM_001364497, NM_001364499, NM_001364500, NM_001364502, NM_001364498, NM_001364501, NR_157195, NR_157196, NM_001290253.2, NM_001290252.2, NM_001039350.3, NM_001936.5
UniProt ID:	P42658
Summary:	This gene encodes a single-pass type II membrane protein that is a member of the peptidase S9B family of serine proteases. This protein has no detectable protease activity, most likely due to the absence of the conserved serine residue normally present in the catalytic domain of serine proteases. However, it does bind specific voltage-gated potassium channels and alters their expression and biophysical properties. Variations in this gene may be associated with susceptibility to amyotrophic lateral sclerosis and with idiopathic ventricular fibrillation. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 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).