

Product datasheet for TL312855

GADD45A Human shRNA Plasmid Kit (Locus ID 1647)

Product data:

OriGene Technologies, Inc.

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Product Type:	shRNA Plasmids
Product Name:	GADD45A Human shRNA Plasmid Kit (Locus ID 1647)
Locus ID:	1647
Synonyms:	DDIT1; GADD45
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	GADD45A - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 1647). 5μg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<u>NM 001199741</u> , <u>NM 001199742</u> , <u>NM 001924</u> , <u>NM 001924.1</u> , <u>NM 001924.2</u> , <u>NM 001924.3</u> , <u>NM 001199742.1</u> , <u>NM 001199741.1</u> , <u>BC011757</u> , <u>BC011757.2</u> , <u>NM 001199742.2</u> , <u>NM 001199741.2</u> , <u>NM 001924.4</u>
UniProt ID:	<u>P24522</u>
Summary:	This gene is a member of a group of genes whose transcript levels are increased following stressful growth arrest conditions and treatment with DNA-damaging agents. The protein encoded by this gene responds to environmental stresses by mediating activation of the p38/JNK pathway via MTK1/MEKK4 kinase. The DNA damage-induced transcription of this gene is mediated by both p53-dependent and -independent mechanisms. Alternatively spliced transcript variants encoding distinct isoforms have been found for this gene.[provided by RefSeq, Dec 2010]
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> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .



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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).

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