OriGene Technologies, Inc.
9620 Medical Center Drive, Ste 200
Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com
techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

Product datasheet for TF320691

MAP4K6 (MINK1) Human shRNA Plasmid Kit (Locus ID 50488)

Product data:

Product Type:	shRNA Plasmids
Product Name:	MAP4K6 (MINK1) Human shRNA Plasmid Kit (Locus ID 50488)
Locus ID:	50488
Synonyms:	B55; MAP4K6; MINK; YSK2; ZC3
Vector:	pRFP-C-RS (TR30014)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	MINK1 - Human, 4 unique 29mer shRNA constructs in retroviral RFP vector(Gene ID = 50488). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRFP-C-RS Vector, TR30015, included for free.
RefSeq:	NM_001024937, NM_001321236, NM_015716, NM_153827, NM_170663, NM_153827.1, NM_153827.2, NM_153827.3, NM_153827.4, NM_015716.1, NM_015716.2, NM_015716.3, NM_015716.4, NM_001024937.1, NM_001024937.2, NM_001024937.3, NM_170663.1, NM_170663.2, NM_170663.3, NM_170663.4, BC034673, BC034673.1, BC028888, BC094686, BM674875, NM_015716.5, NM_170663.5, NM_001024937.4
UniProt ID:	Q8N4C8
Summary:	This gene encodes a serine/threonine kinase belonging to the germinal center kinase (GCK) family. The protein is structurally similar to the kinases that are related to NIK and may belong to a distinct subfamily of NIK-related kinases within the GCK family. Studies of the mouse homolog indicate an up-regulation of expression in the course of postnatal mouse cerebral development and activation of the cJun N-terminal kinase (JNK) and the p38 pathways. [provided by RefSeq, Mar 2016]
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).