

Product datasheet for TR311540

OriGene Technologies, Inc.

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GANP (MCM3AP) Human shRNA Plasmid Kit (Locus ID 8888)

Product data:

Product Type: shRNA Plasmids

Product Name: GANP (MCM3AP) Human shRNA Plasmid Kit (Locus ID 8888)

Locus ID:

GANP; MAP80; PNRIID; SAC3 Synonyms:

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

MCM3AP - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

8888). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

NM 003906, NM 003906.1, NM 003906.2, NM 003906.3, NM 003906.4, BC104958, BC004497, RefSeq:

BC013285, BC104960, NM 003906.5

UniProt ID: 060318

Summary: The minichromosome maintenance protein 3 (MCM3) is one of the MCM proteins essential

> for the initiation of DNA replication. The protein encoded by this gene is a MCM3 binding protein. It was reported to have phosphorylation-dependent DNA-primase activity, which was

up-regulated in antigen immunization induced germinal center. This protein was

demonstrated to be an acetyltransferase that acetylates MCM3 and plays a role in DNA replication. The mutagenesis of a nuclear localization signal of MCM3 affects the binding of this protein with MCM3, suggesting that this protein may also facilitate MCM3 nuclear localization. This gene is expressed in the brain or in neuronal tissue. An allelic variant

encoding amino acid Lys at 915, instead of conserved Glu, has been identified in patients with

mild intellectual disability. [provided by RefSeq, Jan 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.







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).