

## Product datasheet for TG311558

## MBD4 Human shRNA Plasmid Kit (Locus ID 8930)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** MBD4 Human shRNA Plasmid Kit (Locus ID 8930)

Locus ID: 8930 MED1 Synonyms:

Vector: pGFP-V-RS (TR30007)

E. coli Selection: Kanamycin Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: MBD4 - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 8930).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free.

NM 001276270, NM 001276271, NM 001276272, NM 001276273, NM 003925, NM 003925.1, RefSeq:

NM 003925.2, NM 001276273.1, NM 001276272.1, NM 001276271.1, NM 001276270.1,

BC011752, BC034463, NM 001276270.2, NM 001276272.2, NM 003925.3

UniProt ID: O95243

Summary: The protein encoded by this gene is a member of a family of nuclear proteins related by the

> presence of a methyl-CpG binding domain (MBD). These proteins are capable of binding specifically to methylated DNA, and some members can also repress transcription from methylated gene promoters. This protein contains an MBD domain at the N-terminus that functions both in binding to methylated DNA and in protein interactions and a C-terminal mismatch-specific glycosylase domain that is involved in DNA repair. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. [provided

by RefSeq, Jan 2013]

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).