

Product datasheet for TR303353

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MAP2 Human shRNA Plasmid Kit (Locus ID 4133)

Product data:

Product Type: shRNA Plasmids

Product Name: MAP2 Human shRNA Plasmid Kit (Locus ID 4133)

Locus ID: 4133

Synonyms: MAP-2; MAP2A; MAP2B; MAP2C

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: MAP2 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

4133). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001039538, NM 002374, NM 031845, NM 031846, NM 031847, NM 002374.1,

NM 002374.2, NM 002374.3, NM 031847.1, NM 031847.2, NM 031845.1, NM 031845.2, NM 001039538.1, BC027583, BC038857, BC066648, BC110423, BC117123, BC143245, BC172263, BC172568, NM 001363911, NM 001363910, NM 001363913, NM 001039538.2,

NM 031847.3, NM 031845.3

UniProt ID: P11137

Summary: This gene encodes a protein that belongs to the microtubule-associated protein family. The

proteins of this family are thought to be involved in microtubule assembly, which is an essential step in neurogenesis. The products of similar genes in rat and mouse are neuron-specific cytoskeletal proteins that are enriched in dentrites, implicating a role in determining and stabilizing dentritic shape during neuron development. A number of alternatively spliced variants encoding distinct isoforms have been described. [provided by RefSeq, Jan 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 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).