

## **Product datasheet for TL313928**

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## **CHRM1 Human shRNA Plasmid Kit (Locus ID 1128)**

**Product data:** 

**Product Type:** shRNA Plasmids

Product Name: CHRM1 Human shRNA Plasmid Kit (Locus ID 1128)

**Locus ID:** 1128

Synonyms: HM1; M1; M1R

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

Puromycin

Selection:

Format: Lentiviral plasmids

Components: CHRM1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 1128).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: <u>BC007740</u>, <u>NM 000738</u>, <u>NM 000738.1</u>, <u>NM 000738.2</u>, <u>BC007740.2</u>, <u>BC022984</u>, <u>BC022984</u>.1,

NM 000738.3

UniProt ID: P11229

**Summary:** The muscarinic cholinergic receptors belong to a larger family of G protein-coupled receptors.

The functional diversity of these receptors is defined by the binding of acetylcholine and

includes cellular responses such as adenylate cyclase inhibition, phosphoinositide degeneration, and potassium channel mediation. Muscarinic receptors influence many effects of acetylcholine in the central and peripheral nervous system. The muscarinic

cholinergic receptor 1 is involved in mediation of vagally-induced bronchoconstriction and in the acid secretion of the gastrointestinal tract. The gene encoding this receptor is localized to

11q13. [provided by RefSeq, Jul 2008]

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>.







## 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).