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Product datasheet for TR308903

TCF4 Human shRNA Plasmid Kit (Locus ID 6925)

Product data:

Product Type:	shRNA Plasmids
Product Name:	TCF4 Human shRNA Plasmid Kit (Locus ID 6925)
Locus ID:	6925
Synonyms:	bHLHb19; CDG2T; E2-2; FECD3; ITF-2; ITF2; PTHS; SEF-2; SEF2; SEF2-1; SEF2-1A; SEF2-1B; SEF2-1D; TCF-4
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	TCF4 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 6925). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	NM_001083962, NM_001243226, NM_001243227, NM_001243228, NM_001243230, NM_001243231, NM_001243232, NM_001243233, NM_001243234, NM_001243235, NM_001243236, NM_003199, NM_001306207, NM_001306208, NM_001330604, NM_001330605, NM_001348211, NM_001348212, NM_001348213, NM_001348214, NM_001348215, NM_001348216, NM_001348217, NM_001348218, NM_001348219, NM_001348220, NM_003199.1, NM_003199.2, NM_001083962.1, NM_001243235.1, NM_001243236.1, NM_001243234.1, NM_001243233.1, NM_001243232.1, NM_001243231.1, NM_001243227.1, NM_001243230.1, NM_001243228.1, NM_001243226.1, NM_001243226.2, BC125084, BC031056, BC125085, BM011452, NM_001369567, NM_001369571, NM_001369574, NM_001369577, NM_001369579, NM_001369580, NM_001369581, NM_001369585, NM_001369586, NM_001369568, NM_001369569, NM_001369570, NM_001369572, NM_001369573, NM_001369575, NM_001369576, NM_001369578, NM_001369582, NM_001369583, NM_001369584, NM_001243227.2, NM_001243228.2, NM_001243236.2, NM_001243234.2, NM_001243231.2, NM_001243235.2, NM_001243233.2, NM_001083962.2
UniProt ID:	P15884

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Summary:

This gene encodes transcription factor 4, a basic helix-loop-helix transcription factor. The encoded protein recognizes an Ephrussi-box ('E-box') binding site ('CANNTG') - a motif first identified in immunoglobulin enhancers. This gene is broadly expressed, and may play an important role in nervous system development. Defects in this gene are a cause of Pitt-Hopkins syndrome. In addition, an intronic CTG repeat normally numbering 10-37 repeat units can expand to >50 repeat units and cause Fuchs endothelial corneal dystrophy. Multiple alternatively spliced transcript variants that encode different proteins have been described. [provided by RefSeq, Jul 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.

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).