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Product datasheet for TL313930

Chimaerin 2 (CHN2) Human shRNA Plasmid Kit (Locus ID 1124)

Product data:

shRNA Plasmids
Chimaerin 2 (CHN2) Human shRNA Plasmid Kit (Locus ID 1124)
1124
ARHGAP3; BCH; CHN2-3; RHOGAP3
pGFP-C-shLenti (TR30023)
Chloramphenicol (34 ug/ml)
Puromycin
Lentiviral plasmids
CHN2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 1124). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
NM 001039936, NM 001293069, NM 001293070, NM 001293071, NM 001293072, NM 001293072, NM 001293073, NM 001293075, NM 001293076, NM 001293077, NM 001293078, NM 001293079, NM 001293080, NM 001293081, NM 004067, NR 120524, NR 120525, NM 004067.1, NM 004067.2, NM 001293075.1, NM 001039936.1, NM 001039936.2, NM 001293079.1, NM 001293078.1, NM 001293075.1, NM 001293077.1, NM 001293081.1, NM 001293077.1, NM 001293072.1, NM 001293
<u>P52757</u>
This gene encodes a guanosine triphosphate (GTP)-metabolizing protein that contains a phorbol-ester/diacylglycerol (DAG)-type zinc finger, a Rho-GAP domain, and an SH2 domain. The encoded protein translocates from the cytosol to the Golgi apparatus membrane upon binding by diacylglycerol (DAG). Activity of this protein is important in cell proliferation and migration, and expression changes in this gene have been detected in cancers. A mutation in this gene has also been associated with schizophrenia in men. Alternative transcript splicing and the use of alternative promoters results in multiple transcript variants. [provided by RefSeq, May 2014]



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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).

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