

Product datasheet for TR316498

CISH Human shRNA Plasmid Kit (Locus ID 1154)

Product data:

Product Type: shRNA Plasmids

Product Name: CISH Human shRNA Plasmid Kit (Locus ID 1154)

Locus ID: 1154

Synonyms: BACTS2; CIS; CIS-1; G18; SOCS

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

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

1154). 5µg purified plasmid DNA per construct

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

RefSeq: NM 013324, NM 145071, NM 145071.1, NM 013324.2, NM 013324.3, NM 013324.4,

NM 013324.5, BC031590, BC031590.1, BC064354, NM 013324.6, NM 145071.4

UniProt ID: Q9NSE2

Summary: The protein encoded by this gene contains a SH2 domain and a SOCS box domain. The

protein thus belongs to the cytokine-induced STAT inhibitor (CIS), also known as suppressor of cytokine signaling (SOCS) or STAT-induced STAT inhibitor (SSI), protein family. CIS family members are known to be cytokine-inducible negative regulators of cytokine signaling. The expression of this gene can be induced by IL2, IL3, GM-CSF and EPO in hematopoietic cells. Proteasome-mediated degradation of this protein has been shown to be involved in the inactivation of the erythropoietin receptor. Multiple transcript variants encoding different

isoforms have been found for this gene. [provided by RefSeq, Aug 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>.



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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).