

## **Product datasheet for TL513264**

## Itih4 Mouse shRNA Plasmid (Locus ID 16427)

## **Product data:**

**Product Type:** shRNA Plasmids

**Product Name:** Itih4 Mouse shRNA Plasmid (Locus ID 16427)

**Locus ID:** 16427

Synonyms: ITI-HC4; Itih; Itih-4; PK-120 Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** Itih4 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 16427).

5µg purified plasmid DNA per construct

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

RefSeq: <u>BC016500</u>, <u>BC092258</u>, <u>BC094457</u>, <u>NM 001159299</u>, <u>NM 001289632</u>, <u>NM 001289633</u>,

NM 018746, NM 018746.1, NM 018746.2, NM 018746.3, NM 018746.4, NM 001159299.1,

NM 001159299.2, NM 001289633.1, NM 001289632.1, BC021459

UniProt ID: A6X935

Summary: This gene encodes a member of the inter-alpha trypsin inhibitor (IaI) family of plasma serine

protease inhibitors with diverse functions as anti-apoptotic and matrix stabilizing molecules during development. This gene is predominantly expressed in the liver and the encoded protein was found to be a plasma kallikrein-sensitive glycoprotein. This gene is located in a cluster of related inter alpha trypsin inhibitor genes on chromosome 14. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Oct

2015]

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.



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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).