

## **Product datasheet for TR320681**

## OriGene Technologies, Inc.

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## STK36 Human shRNA Plasmid Kit (Locus ID 27148)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** STK36 Human shRNA Plasmid Kit (Locus ID 27148)

**Locus ID:** 27148

Synonyms: FU

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: STK36 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

27148). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001243313, NM 015690, NM 015690.1, NM 015690.2, NM 015690.3, NM 015690.4,

NM 001243313.1, BC026158, BC026158.2, NM 001369423, NM 015690.5, NM 001243313.2

UniProt ID: Q9NRP7

Summary: This gene encodes a member of the serine/threonine kinase family of enzymes. This family

member is similar to a Drosophila protein that plays a key role in the Hedgehog signaling pathway. This human protein is a positive regulator of the GLI zinc-finger transcription factors. Knockout studies of the homologous mouse gene suggest that defects in this human gene may lead to congenital hydrocephalus, possibly due to a functional defect in motile cilia. Because Hedgehog signaling is frequently activated in certain kinds of gastrointestinal cancers, it has been suggested that this gene is a target for the treatment of these cancers.

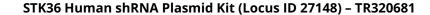
Alternative splicing of this gene results in multiple transcript variants. [provided by RefSeq,

Aug 2011]

**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.







## 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).