

## Product datasheet for TR307431

## OriGene Technologies, Inc.

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## **TIFA Human shRNA Plasmid Kit (Locus ID 92610)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** TIFA Human shRNA Plasmid Kit (Locus ID 92610)

Locus ID:

T2BP; T6BP; TIFAA Synonyms:

pRS (TR20003) Vector:

E. coli Selection: Ampicillin Mammalian Cell

Selection:

Puromycin

Format: Retroviral plasmids

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

92610). 5µg purified plasmid DNA per construct

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

NM 052864, NM 052864.1, NM 052864.2, BC014259, BC014259.2, BC008294, BM680110, RefSeq:

BM718895, NM 052864.3

UniProt ID: 096CG3

**Summary:** This gene encodes an adapter protein involved in adaptive and innate immunity. This protein

> includes a forkhead-associated (FHA) domain that specifically binds to phosphorylated serine and threonine residues. In response to bacterial infection, the encoded host cell protein undergoes an intermolecular interaction between the FHA domain and a phosphorylated threonine that leads to protein oligomerization and stimulation of the NF-kappa B and other downstream signaling pathways. This protein exhibits reduced expression in hepatocellular carcinoma and may suppress hepatocellular carcinoma progression. This protein may also

play a role in the DNA damage response. [provided by RefSeq, Jun 2018]

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).