

## **Product datasheet for TL320647**

## OriGene Technologies, Inc.

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## **FASTK Human shRNA Plasmid Kit (Locus ID 10922)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** FASTK Human shRNA Plasmid Kit (Locus ID 10922)

Locus ID: 10922 Synonyms: FAST

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** FASTK - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 10922).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001258461, NM 006712, NM 025096, NM 033015, NM 006712.1, NM 006712.2,

NM 006712.3, NM 006712.4, NM 033015.1, NM 033015.2, NM 033015.3, NM 001258461.1,

BC011770, BC011770.2, BC039026, BC039026.2, BC000377, BC006386, BM799420,

NM 006712.5

UniProt ID: 014296

**Summary:** The protein encoded by this gene is a member of the serine/threonine protein kinase family.

This kinase was shown to be activated rapidly during Fas-mediated apoptosis in Jurkat cells. In response to Fas receptor ligation, it phosphorylates TIA1, an apoptosis-promoting nuclear RNA-binding protein. The encoded protein is a strong inducer of lymphocyte apoptosis. Two transcript variants encoding different isoforms have been found for this gene. Other variants exist, but their full-length natures have not yet been determined. [provided by RefSeq, Jul

20081

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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a>.
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).