

## **Product datasheet for TL320414**

MATK Human shRNA Plasmid Kit (Locus ID 4145)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** MATK Human shRNA Plasmid Kit (Locus ID 4145)

Locus ID: 4145

Synonyms: CHK; CTK; HHYLTK; HYL; HYLTK; Lsk

**Vector:** pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: MATK - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 4145).

5µg purified plasmid DNA per construct

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

RefSeq: NM 002378, NM 139354, NM 139355, NM 139354.1, NM 139354.2, NM 139355.1,

NM 139355.2, NM 002378.1, NM 002378.2, NM 002378.3, BC000114, BC000114.1, BC003109,

BM925586, NM 002378.4, NM 139355.3

UniProt ID: P42679

Summary: The protein encoded by this gene has amino acid sequence similarity to Csk tyrosine kinase

and has the structural features of the CSK subfamily: SRC homology SH2 and SH3 domains, a

catalytic domain, a unique N terminus, lack of myristylation signals, lack of a negative regulatory phosphorylation site, and lack of an autophosphorylation site. This protein is thought to play a significant role in the signal transduction of hematopoietic cells. It is able to phosphorylate and inactivate Src family kinases, and may play an inhibitory role in the control of T-cell proliferation. This protein might be involved in signaling in some cases of breast cancer. Three alternatively spliced transcript variants that encode different isoforms have

been described for this gene. [provided by RefSeq, Jul 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).