

## Product datasheet for **TL320634V**

### ACK1 (TNK2) Human shRNA Lentiviral Particle (Locus ID 10188)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	ACK1 (TNK2) Human shRNA Lentiviral Particle (Locus ID 10188)
Locus ID:	10188
Synonyms:	ACK; ACK-1; ACK1; p21cdc42Hs
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	TNK2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">BC008884</a> , <a href="#">NM_001010938</a> , <a href="#">NM_001308046</a> , <a href="#">NM_005781</a> , <a href="#">NM_005781.1</a> , <a href="#">NM_005781.2</a> , <a href="#">NM_005781.3</a> , <a href="#">NM_005781.4</a> , <a href="#">NM_001010938.1</a> , <a href="#">BC020389</a> , <a href="#">BC028164</a> , <a href="#">BC156798</a> , <a href="#">NM_005781.5</a>
UniProt ID:	<a href="#">Q07912</a>
Summary:	This gene encodes a tyrosine kinase that binds Cdc42Hs in its GTP-bound form and inhibits both the intrinsic and GTPase-activating protein (GAP)-stimulated GTPase activity of Cdc42Hs. This binding is mediated by a unique sequence of 47 amino acids C-terminal to an SH3 domain. The protein may be involved in a regulatory mechanism that sustains the GTP-bound active form of Cdc42Hs and which is directly linked to a tyrosine phosphorylation signal transduction pathway. Several alternatively spliced transcript variants have been identified from this gene, but the full-length nature of only two transcript variants has been determined. [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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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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](mailto: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).