

## **Product datasheet for SR303089**

### OriGene Technologies, Inc.

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## Nck (NCK1) Human siRNA Oligo Duplex (Locus ID 4690)

#### **Product data:**

**Product Type:** siRNA Oligo Duplexes

Purity: HPLC purified

**Quality Control:** Tested by ESI-MS

Sequences: Available with shipment

**Stability:** One year from date of shipment when stored at -20°C.

# of transfections: Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final

conc. 10 nM).

 Note:
 Single siRNA duplex (10nmol) can be ordered.

 RefSeq:
 NM 001190796, NM 001291999, NM 006153

UniProt ID: P16333

Synonyms: NCK; nck-1; NCKalpha

Components: NCK1 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 4690)

Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol

Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 ml

**Summary:** The protein encoded by this gene is one of the signaling and transforming proteins

containing Src homology 2 and 3 (SH2 and SH3) domains. It is located in the cytoplasm and is

an adaptor protein involved in transducing signals from receptor tyrosine kinases to

downstream signal recipients such as RAS. Alternatively spliced transcript variants encoding

different isoforms have been found. [provided by RefSeq, Jun 2010]



# Performance Guaranteed:

OriGene guarantees that at least two of the three Dicer-Substrate duplexes in the kit will provide at least 70% or more knockdown of the target mRNA when used at 10 nM concentration by quantitative RT-PCR when the TYE-563 fluorescent transfection control duplex (cat# SR30002) indicates that >90% of the cells have been transfected and the HPRT positive control (cat# SR30003) provides 90% knockdown efficiency.

For non-conforming siRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the siRNA kit. To arrange for a free replacement with newly designed duplexes, 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 siRNA control (quantitative RT-PCR data required).