

## **Product datasheet for SR325847**

## OriGene Technologies, Inc.

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## **CNIH3 Human siRNA Oligo Duplex (Locus ID 149111)**

**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 001322302, NM 001322303, NM 001322304, NM 001322305, NM 152495, NR 136287,

NR 136288, NR 136289, NR 136290, NR 136291, NR 136292, NR 136293, NR 136294,

NR 136295, NR 136296, NR 136297

UniProt ID: Q8TBE1
Synonyms: CNIH-3

Components: CNIH3 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 149111)

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

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

**Summary:** Regulates the trafficking and gating properties of AMPA-selective glutamate receptors

(AMPARs). Promotes their targeting to the cell membrane and synapses and modulates their gating properties by regulating their rates of activation, deactivation and desensitization.

[UniProtKB/Swiss-Prot Function]







## 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).