

## **Product datasheet for SR310625**

## OriGene Technologies, Inc.

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## **DNAI7 Human siRNA Oligo Duplex (Locus ID 55259)**

**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 001082972, NM 001082973, NM 001204101, NM 001204102, NM 001319977,

NM 001319978, NM 018272, NM 001352061, NM 001352062, NM 001352063, NM 001352064, NM 001352065, NM 001352066, NM 001352067, NM 001352068,

NR 147909, NR 147910, NR 147911

UniProt ID: Q6TDU7

Synonyms: CASC1; LAS1; PPP1R54

Components: CASC1 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 55259)

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

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

**Summary:** Via its association with the multisubunit axonemal dynein complex, is potentially involved in

the regulation of cilia function. May also act as a cell cycle regulator. [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).