

Product datasheet for SR313758

OriGene Technologies, Inc.

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HSH2D Human siRNA Oligo Duplex (Locus ID 84941)

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 001291274, NM 032855, NR 111903, NR 111904, NM 001352265, NM 001352266,

NM 001369808, NR 163150, NR 163152, NR 163153, NR 163155, NM 001369809,

NR 163151, NR 163154, NR 163156

UniProt ID: Q96|Z2

Synonyms: ALX; HSH2

Components: HSH2D (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 84941)

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

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

Summary: T-cell activation requires 2 signals: recognition of antigen by the T-cell receptor (see TCR; MIM

186880) and a costimulatory signal provided primarily by CD28 (MIM 186760) in naive T cells.

HSH2 is a target of both of these signaling pathways (Greene et al., 2003 [PubMed

12960172]).[supplied by OMIM, Mar 2008]





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).