

Product datasheet for SR421876

OriGene Technologies, Inc.

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Herc4 Mouse siRNA Oligo Duplex (Locus ID 67345)

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 026101, NM 030114, NM 001359484

UniProt ID: Q6PAV2

Synonyms: 1700056O17Rik; 4921531D01Rik; 9530080M15Rik; mKIAA1593

Components: Herc4 (Mouse) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 67345)

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

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

Summary: Probable E3 ubiquitin-protein ligase involved in either protein trafficking or in the distribution

of cellular structures. Required for spermatozoon maturation and fertility, and for the

removal of the cytoplasmic droplet of the spermatozoon. E3 ubiquitin-protein ligases accept

ubiquitin from an E2 ubiquitin-conjugating enzyme in the form of a thioester and then

directly transfer it to targeted substrates.[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).