

Product datasheet for SR323737

OriGene Technologies, Inc.

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RNF167 Human siRNA Oligo Duplex (Locus ID 26001)

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 001320356, NM 001320357, NM 001320358, NM 001320359, NM 001320360,

NM 001320361, NM 001320362, NM 001320363, NM 001320364, NM 001320365, NM 015528, NM 001370306, NM 001370307, NM 001370311, NM 001370303,

NM 001370304, NM 001370305, NM 001370308, NM 001370313

UniProt ID: Q9H6Y7

Synonyms: 5730408C10Rik; LP2254; RING105

Components: RNF167 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 26001)

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

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

Summary: RNF167 is an E3 ubiquitin ligase that interacts with TSSC5 (SLC22A18; MIM 602631) and,

together with UBCH6 (UBE2E1; MIM 602916), facilitates TSSC5 polyubiquitylation (Yamada

and Gorbsky, 2006 [PubMed 16314844]).[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).