

Product datasheet for SR416910

OriGene Technologies, Inc.

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Nrf1 Mouse siRNA Oligo Duplex (Locus ID 18181)

Product data:

Product Type: siRNA Oligo Duplexes

HPLC purified **Purity:**

Quality Control: Tested by ESI-MS

Available with shipment Sequences:

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.

NM 001164226, NM 001164227, NM 001164228, NM 001164229, NM 001164230, RefSeq:

NM 010938, NM 001361692, NM 001361693, NM 001361694, NM 001361695

Q9WU00 **UniProt ID:**

Synonyms: C87038; D6Ertd415e

Components: Nrf1 (Mouse) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 18181)

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

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

Transcription factor that activates the expression of the EIF2S1 (EIF2-alpha) gene. Links the **Summary:**

> transcriptional modulation of key metabolic genes to cellular growth and development. Implicated in the control of nuclear genes required for respiration, heme biosynthesis, and

mitochondrial DNA transcription and replication (By similarity).[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).