

Product datasheet for SR305595

OriGene Technologies, Inc.

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DGKD Human siRNA Oligo Duplex (Locus ID 8527)

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.

RefSeq: NM 003648, NM 152879

UniProt ID: Q16760

Synonyms: DGK-delta; dgkd-2; DGKdelta

Components: DGKD (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 8527)

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

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

This gene encodes a cytoplasmic enzyme that phosphorylates diacylglycerol to produce **Summary:**

phosphatidic acid. Diacylglycerol and phosphatidic acid are two lipids that act as second

messengers in signaling cascades. Their cellular concentrations are regulated by the encoded

protein, and so it is thought to play an important role in cellular signal transduction.

Alternative splicing results in two transcript variants encoding different isoforms. [provided

by RefSeq, Jul 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).