

## **Product datasheet for SR308070**

## OriGene Technologies, Inc.

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## **DIP2A Human siRNA Oligo Duplex (Locus ID 23181)**

## **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 001146114, NM 001146115, NM 001146116, NM 015151, NM 206889, NM 206890,

NM 206891, NM 001353942, NM 001353943, NM 001353944

UniProt ID: Q14689

Synonyms: C21orf106; DIP2

Components: DIP2A (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 23181)

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

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

Summary: The protein encoded by this gene may be involved in axon patterning in the central nervous

system. This gene is not highly expressed. Several transcript variants encoding different

isoforms have been found for this gene. [provided by RefSeq, Mar 2009]

**Performance** OriGene guarantees that at least two of the three Dicer-Substrate duplexes in the kit will

**Guaranteed:** 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).

