

## **Product datasheet for SR306547**

## OriGene Technologies, Inc.

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## **KIAA0586 Human siRNA Oligo Duplex (Locus ID 9786)**

**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 001244189, NM 001244190, NM 001244191, NM 001244192, NM 001244193,

NM 001329943, NM 001329944, NM 001329945, NM 001329946, NM 001329947,

NM 014749, NM 001364700, NM 001364701

UniProt ID: Q9BVV6

Synonyms: JBTS23; SRTD14; Talpid3

Components: KIAA0586 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 9786)

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

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

**Summary:** This gene encodes a conserved centrosomal protein that functions in ciliogenesis and

responds to hedgehog signaling. Mutations in this gene causes Joubert syndrome 23.

Alternative splicing results in multiple transcript variants and protein isoforms. [provided by

RefSeq, Aug 2016]







## 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).