

## **Product datasheet for SR306195**

#### OriGene Technologies, Inc.

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### SLC22A6 Human siRNA Oligo Duplex (Locus ID 9356)

#### **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 004790, NM 153276, NM 153277, NM 153278, NM 153279

UniProt ID: Q4U2R8

**Synonyms:** HOAT1; OAT1; PAHT; ROAT1

Components: SLC22A6 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 9356)

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 is involved in the sodium-dependent transport and

excretion of organic anions, some of which are potentially toxic. The encoded protein is an

integral membrane protein and may be localized to the basolateral membrane. Four

transcript variants encoding four different isoforms have been found for this gene. [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).