

## **Product datasheet for SR314605**

### OriGene Technologies, Inc.

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## AGAP1 Human siRNA Oligo Duplex (Locus ID 116987)

### **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 001037131, NM 001244888, NM 014914

UniProt ID: Q9UPQ3

**Synonyms:** AGAP-1; CENTG2; cnt-g2; GGAP1

Components: AGAP1 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 116987)

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 member of an ADP-ribosylation factor GTPase-activating protein family

involved in membrane trafficking and cytoskeleton dynamics. This gene functions as a direct

regulator of the adaptor-related protein complex 3 on endosomes. Multiple transcript

variants encoding different isoforms have been found for this gene. [provided by RefSeq, Oct

2011]







# 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).