

Product datasheet for **SR318298**

C16orf88 (KNOP1) Human siRNA Oligo Duplex (Locus ID 400506)

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:	<u>NM_001012991</u> , <u>NM_001348527</u> , <u>NM_001348528</u> , <u>NM_001348529</u> , <u>NM_001348530</u> , <u>NM_001348531</u> , <u>NM_001348532</u> , <u>NM_001348533</u> , <u>NM_001348534</u> , <u>NM_001348535</u> , <u>NM_001348536</u> , <u>NM_001348537</u>
UniProt ID:	<u>Q1ED39</u>
Synonyms:	101F10.1; C16orf88; FAM191A; TSG118
Components:	KNOP1 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 400506) 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 a nucleolar protein that interacts with zinc finger 106 protein. The encoded protein has several of the same characteristics as nucleostemin and may be involved in testis development. [provided by RefSeq, Feb 2017]



[View online »](#)

**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).