

## Product datasheet for SR308941

## OriGene Technologies, Inc.

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## TZFP (ZBTB32) Human siRNA Oligo Duplex (Locus ID 27033)

## **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 001316902, NM 001316903, NM 014383

UniProt ID: Q9Y2Y4

**Synonyms:** FAXF; FAZF; Rog; TZFP; ZNF538

Components: ZBTB32 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 27033)

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

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

**Summary:** DNA-binding protein that binds to the to a 5'-TGTACAGTGT-3' core sequence. May function as

a transcriptional transactivator and transcriptional repressor. Probably exerts its repressor

effect by preventing GATA3 from binding to DNA. May play a role in regulating the

differentiation and activation of helper T-cells (By similarity).[UniProtKB/Swiss-Prot Function]

**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).

