

OriGene Technologies, Inc.

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Product datasheet for SR309021

IL36 alpha (IL36A) Human siRNA Oligo Duplex (Locus ID 27179)

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 014440</u>
UniProt ID:	Q9UHA7
Synonyms:	FIL1; FIL1(EPSILON); FIL1E; IL-1F6; IL1(EPSILON); IL1F6
Components:	IL36A (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 27179) 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 cytokine that can activate NF-kappa-B and MAPK signaling pathways to generate an inflammatory response. The encoded protein functions primarily in skin and demonstrates increased expression in psoriasis. In addition, decreased expression of this gene has been linked to a poor prognosis in both hepatocellular carcinoma and colorectal cancer patients. [provided by RefSeq, Nov 2015]



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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).

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