

Product datasheet for TR303042

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OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

NLRP7 Human shRNA Plasmid Kit (Locus ID 199713)

Product data:

Product Type: shRNA Plasmids

Product Name: NLRP7 Human shRNA Plasmid Kit (Locus ID 199713)

Locus ID:

CLR19.4; HYDM; NALP7; NOD12; PAN7; PYPAF3 Synonyms:

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell Puromycin

Selection: Format:

Retroviral plasmids

NLRP7 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

199713). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

NM 001127255, NM 139176, NM 206828, NM 206828.1, NM 206828.2, NM 206828.3, RefSeq:

NM 139176.1, NM 139176.2, NM 139176.3, NM 001127255.1, BC109125, BC109124

UniProt ID: Q8WX94

Summary: This gene encodes a member of the NACHT, leucine rich repeat, and PYD containing (NLRP)

> protein family. It has an N-terminal pyrin domain, followed by a NACHT domain, a NACHTassociated domain (NAD), and a C-terminal leucine-rich repeat (LRR) region. NLRP proteins are implicated in the activation of proinflammatory caspases through multiprotein complexes called inflammasomes. This gene may act as a feedback regulator of caspase-1-dependent interleukin 1-beta secretion. Alternative splicing results in multiple transcript variants

encoding different isoforms. [provided by RefSeg, Jul 2008]

These shRNA constructs were designed against multiple splice variants at this gene locus. To shRNA Design:

> be certain that your variant of interest is targeted, please contact techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.





Performance Guaranteed:

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, 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 shRNA control (Western Blot data preferred).