

Product datasheet for **TR316428**

P2RX7 Human shRNA Plasmid Kit (Locus ID 5027)

Product data:

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| Product Type: | shRNA Plasmids |
| Product Name: | P2RX7 Human shRNA Plasmid Kit (Locus ID 5027) |
| Locus ID: | 5027 |
| Synonyms: | P2X7 |
| Vector: | pRS (TR20003) |
| E. coli Selection: | Ampicillin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | P2RX7 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 5027). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free. |
| RefSeq: | NM_002562 , NM_177427 , NR_033948 , NR_033949 , NR_033950 , NR_033951 , NR_033952 , NR_033953 , NR_033954 , NR_033955 , NR_033956 , NM_002562.1 , NM_002562.2 , NM_002562.3 , NM_002562.4 , NM_002562.5 , BC007679 , BC011913 , BC121157 , BC121158 , NM_002562.6 |
| UniProt ID: | Q99572 |
| Summary: | The product of this gene belongs to the family of purinoceptors for ATP. This receptor functions as a ligand-gated ion channel and is responsible for ATP-dependent lysis of macrophages through the formation of membrane pores permeable to large molecules. Activation of this nuclear receptor by ATP in the cytoplasm may be a mechanism by which cellular activity can be coupled to changes in gene expression. Multiple alternatively spliced variants have been identified, most of which fit nonsense-mediated decay (NMD) criteria. [provided by RefSeq, Jul 2010] |
| shRNA Design: | These shRNA constructs were designed against multiple splice variants at this gene locus. To 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 . |



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**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).