

Product datasheet for **TR302716**

P2Y6 (P2RY6) Human shRNA Plasmid Kit (Locus ID 5031)

Product data:

Product Type:	shRNA Plasmids
Product Name:	P2Y6 (P2RY6) Human shRNA Plasmid Kit (Locus ID 5031)
Locus ID:	5031
Synonyms:	P2Y6
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	P2RY6 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 5031). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	NM_001277204 , NM_001277205 , NM_001277206 , NM_001277207 , NM_001277208 , NM_004154 , NM_176796 , NM_176797 , NM_176798 , NM_176797.1 , NM_176797.2 , NM_004154.1 , NM_004154.2 , NM_004154.3 , NM_176796.1 , NM_176796.2 , NM_176798.1 , NM_176798.2 , NM_001277204.1 , NM_001277205.1 , NM_001277206.1 , NM_001277207.1 , NM_001277208.1 , BC000571 , BC000571.2 , BC009391 , NM_001277204.2
UniProt ID:	Q15077
Summary:	The product of this gene belongs to the family of P2 receptors, which is activated by extracellular nucleotides and subdivided into P2X ligand-gated ion channels and P2Y G-protein coupled receptors. This family has several receptor subtypes with different pharmacological selectivity, which overlaps in some cases, for various adenosine and uridine nucleotides. This receptor, which is a G-protein coupled receptor, is responsive to UDP, partially responsive to UTP and ADP, and not responsive to ATP. It is proposed that this receptor mediates inflammatory responses. Alternative splicing results in multiple transcript variants that encode different protein isoforms. [provided by RefSeq, Mar 2013]
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).