

## Product datasheet for **TR312411**

### HLAC (HLA-C) Human shRNA Plasmid Kit (Locus ID 3107)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	HLAC (HLA-C) Human shRNA Plasmid Kit (Locus ID 3107)
Locus ID:	3107
Synonyms:	D6S204; HLA-JY3; HLAC; HLC-C; MHC; PSORS1
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	HLA-C - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 3107). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<a href="#">BC002463</a> , <a href="#">BC008457</a> , <a href="#">NM_001243042</a> , <a href="#">NM_002117</a> , <a href="#">NM_002117.1</a> , <a href="#">NM_002117.2</a> , <a href="#">NM_002117.3</a> , <a href="#">NM_002117.4</a> , <a href="#">NM_002117.5</a> , <a href="#">NM_001243042.1</a> , <a href="#">BC002463.1</a> , <a href="#">BC008457.1</a> , <a href="#">BC004489</a> , <a href="#">BC007814</a> , <a href="#">BC010542</a> , <a href="#">BC033293</a> , <a href="#">BC041078</a> , <a href="#">NM_002117.6</a>
UniProt ID:	<a href="#">P10321</a>



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**Summary:**

HLA-C belongs to the HLA class I heavy chain paralogues. This class I molecule is a heterodimer consisting of a heavy chain and a light chain (beta-2 microglobulin). The heavy chain is anchored in the membrane. Class I molecules play a central role in the immune system by presenting peptides derived from endoplasmic reticulum lumen. They are expressed in nearly all cells. The heavy chain is approximately 45 kDa and its gene contains 8 exons. Exon one encodes the leader peptide, exons 2 and 3 encode the alpha1 and alpha2 domain, which both bind the peptide, exon 4 encodes the alpha3 domain, exon 5 encodes the transmembrane region, and exons 6 and 7 encode the cytoplasmic tail. Polymorphisms within exon 2 and exon 3 are responsible for the peptide binding specificity of each class one molecule. Typing for these polymorphisms is routinely done for bone marrow and kidney transplantation. About 6000 HLA-C alleles have been described. The HLA system plays an important role in the occurrence and outcome of infectious diseases, including those caused by the malaria parasite, the human immunodeficiency virus (HIV), and the severe acute respiratory syndrome coronavirus (SARS-CoV). The structural spike and the nucleocapsid proteins of the novel coronavirus SARS-CoV-2, which causes coronavirus disease 2019 (COVID-19), are reported to contain multiple Class I epitopes with predicted HLA restrictions. Individual HLA genetic variation may help explain different immune responses to a virus across a population.[provided by RefSeq, Aug 2020]

**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](mailto: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](mailto: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).