

Product datasheet for **TL319507V**

HMGA2 Human shRNA Lentiviral Particle (Locus ID 8091)

Product data:

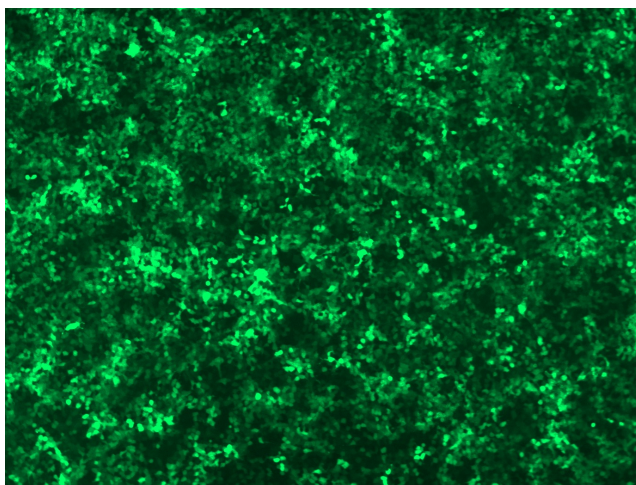
Product Type:	shRNA Lentiviral Particles
Product Name:	HMGA2 Human shRNA Lentiviral Particle (Locus ID 8091)
Locus ID:	8091
Synonyms:	BABL; BABL, LIPO, HMGIC, HMGI-C; high-mobility group (nonhistone chromosomal) protein isoform I-C; High-mobility group protein HMGI-C; high mobility group AT-hook 2; HMGI-C; HMGIC; LIPO; STQTL9
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	HMGA2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, $>10^7$ TU/ml.
RefSeq:	NM_001015886 , NM_001300918 , NM_001300919 , NM_001330190 , NM_003483 , NM_003484 , NM_003483.1 , NM_003483.2 , NM_003483.3 , NM_003483.4 , NM_003484.1 , NM_001300918.1 , NM_001300919.1 , NM_001015886.1 , BC160115 , BM782659 , NM_003483.6
UniProt ID:	P52926
Summary:	This gene encodes a protein that belongs to the non-histone chromosomal high mobility group (HMG) protein family. HMG proteins function as architectural factors and are essential components of the enhancosome. This protein contains structural DNA-binding domains and may act as a transcriptional regulating factor. Identification of the deletion, amplification, and rearrangement of this gene that are associated with myxoid liposarcoma suggests a role in adipogenesis and mesenchymal differentiation. A gene knock out study of the mouse counterpart demonstrated that this gene is involved in diet-induced obesity. Alternate transcriptional splice variants, encoding different isoforms, have been characterized. [provided by RefSeq, Jul 2008]
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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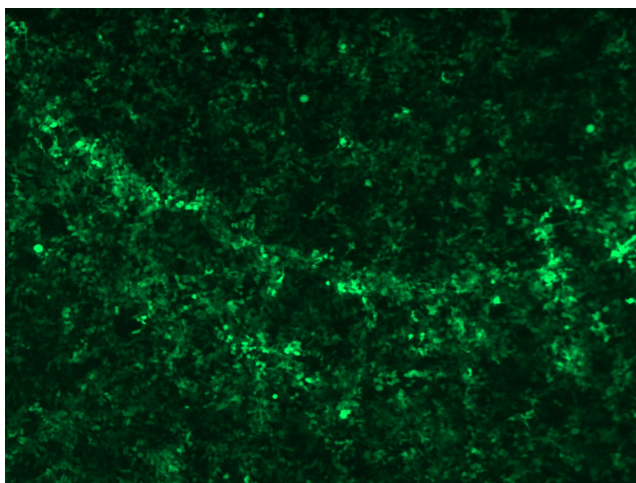
**Performance
Guaranteed:**

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.

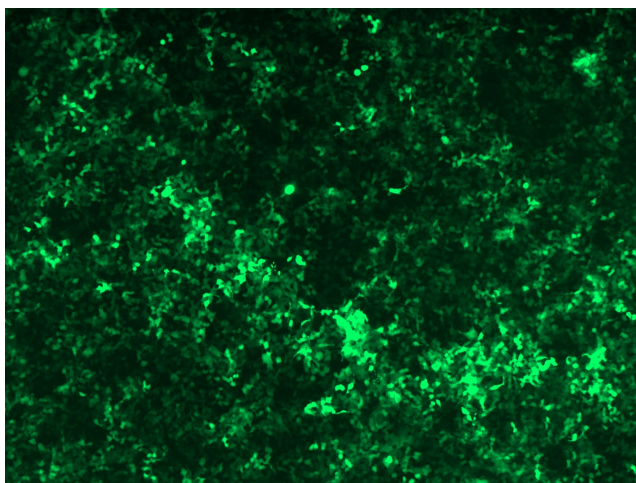
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

Product images:


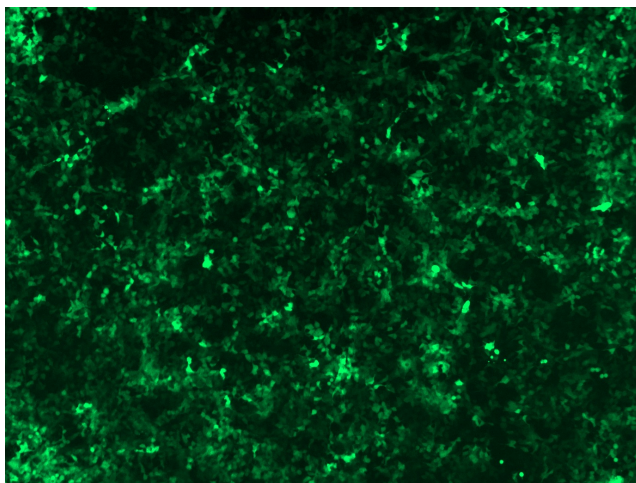
GFP signal was observed under microscope at 48 hours after transduction of TL319507A virus into HEK293 cells. TL319507A virus was prepared using lenti-shRNA TL319507A and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of TL319507B virus into HEK293 cells. TL319507B virus was prepared using lenti-shRNA TL319507B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL319507C] virus into HEK293 cells. [TL319507C] virus was prepared using lenti-shRNA [TL319507C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL319507D] virus into HEK293 cells. [TL319507D] virus was prepared using lenti-shRNA [TL319507D] and [TR30037] packaging kit.