

Product datasheet for TL516114

Pomc Mouse shRNA Plasmid (Locus ID 18976)

Product data:

Product Type: shRNA Plasmids **Product Name:** Pomc Mouse shRNA Plasmid (Locus ID 18976) Locus ID: 18976 ACT; ACTH; alp; alph; alpha-MSH; alphaMSH; BE; Beta-LPH; beta-M; beta-MSH; Clip; gamma-; Synonyms: Gamma-LPH; gamma-MSH; Npp; PO; Pomc-1; Pomc1 Vector: pGFP-C-shLenti (TR30023) E. coli Selection: Chloramphenicol (34 ug/ml) Mammalian Cell Puromycin Selection: Format: Lentiviral plasmids Pomc - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 18976). **Components:** 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free. **RefSeq:** BC061215, NM 001278581, NM 001278582, NM 001278583, NM 001278584, NM 008895, NM 008895.1, NM 008895.2, NM 008895.3, NM 008895.4, NM 001278581.1, NM 001278582.1, NM 001278583.1, NM 001278584.1 **UniProt ID:** P01193 Summary: This gene encodes a polypeptide hormone precursor that undergoes extensive, tissuespecific, post-translational processing. Processing yields several biologically active peptides, which are involved in diverse cellular functions, such as energy homeostasis, steroidogenesis, and increased melanin production in melanocytes. In mouse deficiency of this gene is associated with obesity, defects in adrenal development, and altered pigmentation. A pseudogene of this gene is located on chromosome 19. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jun 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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CRIGENE Pomc Mouse shRNA Plasmid (Locus ID 18976) – TL516114

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).

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