

## **Product datasheet for TL312311**

## OriGene Technologies, Inc.

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## **HSP90AB1 Human shRNA Plasmid Kit (Locus ID 3326)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** HSP90AB1 Human shRNA Plasmid Kit (Locus ID 3326)

Locus ID: 3326

Synonyms: D6S182; HSP84; HSP90B; HSPC2; HSPCB

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: HSP90AB1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

3326). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 001271969, NM 001271970, NM 001271971, NM 001271972, NM 007355, NM 007355.1,

NM 007355.2, NM 007355.3, NM 001271971.1, NM 001271972.1, NM 001271969.1, NM 001271970.1, BC012807, BC012807.2, BC004928, BC007327, BC009206, BC014485,

BC016753, BC068474, NR 073528, NM 007355.4

UniProt ID: P08238

Summary: This gene encodes a member of the heat shock protein 90 family; these proteins are involved

in signal transduction, protein folding and degradation and morphological evolution. This gene encodes the constitutive form of the cytosolic 90 kDa heat-shock protein and is thought to play a role in gastric apoptosis and inflammation. Alternative splicing results in multiple transcript variants. Pseudogenes have been identified on multiple chromosomes. [provided

by RefSeq, Dec 2012]

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 <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.





## 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).