

## **Product datasheet for TL312532**

## OriGene Technologies, Inc.

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## **HABP2 Human shRNA Plasmid Kit (Locus ID 3026)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** HABP2 Human shRNA Plasmid Kit (Locus ID 3026)

Locus ID: 3026

**Synonyms:** FSAP; HABP; HGFAL; NMTC5; PHBP

**Vector:** pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: HABP2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 3026).

5µg purified plasmid DNA per construct

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

**RefSeq:** NM 001177660, NM 004132, NM 004132.1, NM 004132.2, NM 004132.3, NM 004132.4,

NM 001177660.1, NM 001177660.2, BC031412, BC031412.1, NM 004132.5

UniProt ID: Q14520

**Summary:** This gene encodes a member of the peptidase S1 family of serine proteases. The encoded

preproprotein is secreted by hepatocytes and proteolytically processed to generate heavy and light chains that form the mature heterodimer. Further autoproteolysis leads to smaller, inactive peptides. This extracellular protease binds hyaluronic acid and may play a role in the

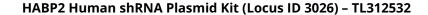
coagulation and fibrinolysis systems. Mutations in this gene are associated with

nonmedullary thyroid cancer and susceptibility to venous thromboembolism. Alternative splicing results in multiple transcript variants, at least one of which encodes a preproprotein

that is proteolytically processed. [provided by RefSeq, Jan 2016]

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).