

## **Product datasheet for TL304282**

## OriGene Technologies, Inc.

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## Glycoprotein VI (GP6) Human shRNA Plasmid Kit (Locus ID 51206)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Glycoprotein VI (GP6) Human shRNA Plasmid Kit (Locus ID 51206)

**Locus ID:** 51206

Synonyms: BDPLT11; GPIV; GPVI

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

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** GP6 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 51206).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001083899, NM 001256017, NM 016363, NM 001083899.1, NM 001083899.2,

NM 016363.1, NM 016363.2, NM 016363.4, NM 016363.5, NM 001256017.1,

NM 001256017.2, BC069485, BC104832, BC111963

UniProt ID: Q9HCN6

**Summary:** This gene encodes a platelet membrane glycoprotein of the immunoglobulin superfamily.

The encoded protein is a receptor for collagen and plays a critical role in collagen-induced platelet aggregation and thrombus formation. The encoded protein forms a complex with the Fc receptor gamma-chain that initiates the platelet activation signaling cascade upon collagen binding. Mutations in this gene are a cause of platelet-type bleeding disorder-11 (BDPLT11). Alternatively spliced transcript variants encoding multiple isoforms have been observed for

this gene. [provided by RefSeq, Dec 2011]

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).