

Product datasheet for TL314719

OriGene Technologies, Inc.

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Apolipoprotein H (APOH) Human shRNA Plasmid Kit (Locus ID 350)

Product data:

Product Type: shRNA Plasmids

Product Name: Apolipoprotein H (APOH) Human shRNA Plasmid Kit (Locus ID 350)

Locus ID: 350

Synonyms: B2G1; B2GP1; BG

Vector: pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: APOH - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 350). 5µg

purified plasmid DNA per construct

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

RefSeq: NM 000042, NM 000042.1, NM 000042.2, BC026283, BC020703, BM005736, NM 000042.3

UniProt ID: P02749

Summary: Apolipoprotein H, also known as beta-2-glycoprotein I, is a component of circulating plasma

lipoproteins. It has been implicated in a variety of physiologic pathways including lipoprotein

metabolism, coagulation, hemostasis, and the production of antiphospholipid

autoantibodies. APOH may be a required cofactor for anionic phospholipid binding by the antiphospholipid autoantibodies found in sera of many patients with lupus and primary antiphospholipid syndrome (APS). The anti-beta (2) glycoprotein I antibodies from APS patients, mediate inhibition of activated protein C which has anticoagulant properties. Because beta-2-GPI is the main autoantigen in patients with APS, the disruption of this

pathway by autoantibodies may be an important mechanism for thrombosis in patients with

APS.[provided by RefSeq, Dec 2019]

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 custom shRNA service.







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).