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Schillerstr. 5

AP26397PU-N Polyclonal Antibody to AGGF1 / VG5Q - Purified

Alternate names: Angiogenic factor VG5Q, Angiogenic factor with G patch and FHA domains 1, G patch

domain-containing protein 7, GPATC7, GPATCH7, Vasculogenesis gene on 5q protein

 Quantity:
 0.1 mg

 Concentration:
 0.1 mg/ml

 Uniprot ID:
 Q8N302

 NCBI:
 NP 060516.2

GenelD: <u>55109</u>

Host / Isotype: Rabbit / IgG

Immunogen: VG5Q functions as an angiogenic factor in promoting angiogenesis and suppression

of VG5Q expression inhibits vessel formation. Angiogenic factors are critical to the initiation of angiogenesis and maintenance of the vascular network. Angiogenesis has an essential role in pathological conditions such as cancer and various ischaemic

and inflammatory diseases.

VG5Q can bind to endothelial cells and promote cell proliferation, suggesting that the protein may act in an autocrine fashion. VG5Q interacts with TWEAK (also known as

TNFSF12), another secreted angiogenic factor.

VG5Q shows strong expression in blood vessels and is secreted when vessel

formation is initiated. VG5Q protein was detected mostly in the cytoplasm and around the nuclei of human microvascular endothelial cells (HMVECs). Furthermore VG5Q is detected in human umbilical vein endothelial cells (HUVECs), human heart fibroblast (HHF) and ovarian cancer cells (OV-3), but low expression was detected in kidney

cancer cells (RP-45), HeLa Cells and bladder cancer cells.

Format: State: Liquid 0.2 µm filtered Ig fraction

Purification: Protein A Buffer System: PBS

Preservatives: 0.02% sodium azide **Stabilizers:** 0.1% bovine serum albumin

Applications: Immunohistochemistry on frozen sections.

Immunohistochemistry on paraffin sections.

Immunoassays.
Immunoprecipitation.

Western blot.

Other applications not tested. Optimal dilutions are dependent on conditions and

should be determined by the user.

Specificity: The antibody reacts with human VG5Q, a 84 kDa protein.

Species Reactivity: Tested: Human
Storage: Store at 2 - 8 °C.

Shelf life: one year from despatch.



General Readings:

1. Tian XL, Kadaba R, You SA, Liu M, Timur AA, Yang L, et al. Identification of an angiogenic factor that when mutated causes susceptibility to Klippel-Trenaunay syndrome. Nature. 2004 Feb 12;427(6975):640-5. PubMed PMID: 14961121.