

Product datasheet for DA3513

VEGF-A (Isoform 121) Human Protein

Product data:

Product Type: Recombinant Proteins

Description: VEGF-A (Isoform 121) human protein, 5 μg

Species: Human
Expression Host: E. coli
Predicted MW: 28.4 kDa

Purity: >98% pure by SDS-PAGE and Silver staining.

Buffer: Presentation State: Purified

State: Lyophilized without buffer and stabilizer.

Buffer System: 50 mM Acetic Acid.

Biological: The ED50 for stimulation of 3H-thymidine incorporation and cell proliferation by

human umbilical vein endothelial cells for VEGF121 has been determined to be in the range

of 1-4 ng/ml.

Endotoxin: $< 0.1 \text{ ng per } \mu\text{g of VEGF121}.$

Reconstitution Method: Restore in PBS or medium containing at least 0.1% Human or Bovine Serum Albumin to a

concentration not lower than 50 µg/ml.

Preparation: Lyophilized without buffer and stabilizer.

Protein Description: Human VEGF is a 28.4 kDa protein consisting of two 121 amino acid polypeptide chains.

Note: Centrifuge vials before opening! Storage: Store lyophilized protein at -20°C.

Reconstituted VEGF121 should be stored in working aliquots at -20°C.

Avoid repeated freeze-thaw cycles!

Stability: Shelf life: One year from despatch.

RefSeq: NP 001020537

 Locus ID:
 7422

 UniProt ID:
 P15692

 Cytogenetics:
 6p21.1

Synonyms: MVCD1; VEGF; VPF



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Summary:

This gene is a member of the PDGF/VEGF growth factor family. It encodes a heparin-binding protein, which exists as a disulfide-linked homodimer. This growth factor induces proliferation and migration of vascular endothelial cells, and is essential for both physiological and pathological angiogenesis. Disruption of this gene in mice resulted in abnormal embryonic blood vessel formation. This gene is upregulated in many known tumors and its expression is correlated with tumor stage and progression. Elevated levels of this protein are found in patients with POEMS syndrome, also known as Crow-Fukase syndrome. Allelic variants of this gene have been associated with microvascular complications of diabetes 1 (MVCD1) and atherosclerosis. Alternatively spliced transcript variants encoding different isoforms have been described. There is also evidence for alternative translation initiation from upstream non-AUG (CUG) codons resulting in additional isoforms. A recent study showed that a C-terminally extended isoform is produced by use of an alternative inframe translation termination codon via a stop codon readthrough mechanism, and that this isoform is antiangiogenic. Expression of some isoforms derived from the AUG start codon is regulated by a small upstream open reading frame, which is located within an internal ribosome entry site. The levels of VEGF are increased during infection with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), thus promoting inflammation by facilitating recruitment of inflammatory cells, and by increasing the level of angiopoietin II (Ang II), one of two products of the SARS-CoV-2 binding target, angiotensin-converting enzyme 2 (ACE2). In turn, Ang II facilitates the elevation of VEGF, thus forming a vicious cycle in the release of inflammatory cytokines. [provided by RefSeq, Jun 2020]

Protein Families:

Druggable Genome, Secreted Protein

Protein Pathways:

Bladder cancer, Cytokine-cytokine receptor interaction, Focal adhesion, mTOR signaling pathway, Pancreatic cancer, Pathways in cancer, Renal cell carcinoma, VEGF signaling pathway

Product images:

