

## Product datasheet for AR26015PU-L

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## **VEGF-A (VEGF164) Mouse Protein**

**Product data:** 

**Product Type: Recombinant Proteins** 

**Description:** VEGF-A (VEGF164) mouse recombinant protein, 20 μg

Species: Mouse **Expression Host:** E. coli

**Expression cDNA Clone** 

APTTEGEQKS HEVIKFMDVY QRSYCRPIET LVDIFQEYPD EIEYIFKPSC VPLMRCAGCC NDEALECVPT or AA Sequence:

SESNITMQIM RIKPHQSQHI GEMSFLQHSR CECRPKKDRT KPENHCEPCS ERRKHLFQDP

QTCKCSCKNT DSRCKARQLE LNERTCRCDK PRR

**Predicted MW:** 38.4 kDa

**Purity:** >95% by SDS-PAGE and silver stain

**Buffer:** Presentation State: Purified

State: Lyophilized protein

Buffer System: 50 mM Acetic Acid

Stabilizer: None

**Bioactivity:** Biological: The ED50 for stimulation of cell proliferation by human umbilical vein endothelial

cells for VEGF164 has been determined to be in the range of 1-5 ng/ml.

**Endotoxin:** < 0.1 ng/µg of VEGF164

**Reconstitution Method:** The lyophilized VEGF164 is soluble in water and most aqueous buffers. The lyophilized

VEGF164 should be reconstituted in PBS or medium containing at least 0.1% human or

bovine serum albumin to a concentration not lower than 50µg/ml.

Preparation: Lyophilized protein

**Protein Description:** Recombinant Mouse Vascular Endothelial Growth Factor164

Note: Protein RefSeq: NP 001020421

mRNA RefSeq: NM 001025250

Storage: Store (following reconstitution in aliquots) at -20°C or below.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

RefSeq: NP 001020421

Locus ID: 22339





UniProt ID: Q00731

Cytogenetics: 17 22.79 cM

Synonyms: VEGFA, VEGF, VPF, Vascular endothelial growth factor A, Vascular permeability factor

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. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. 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 in-frame 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.[provided by RefSeq, Nov 2015]

## **Product images:**

