

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

VEGFA (Human) Recombinant Protein

Catalog Number: P9307

Regulation Status: For research use only (RUO)

Product Description: Human VEGFA recombinant

protein expressed in Insect Cells.

Sequence:

APMAEGGGQNHHEVVKFMDVYQRSYCHPIETLVDIFQ EYPDEIEYIFKPSCVPLMRCGGCCNDEGLECVPTEES NITMQIMRIKPHQGQHIGEMSFLQHNKCECRPKKDRA RQEKCDKPRR

Host: insect

Theoretical MW (kDa): 36

Protocols: See our web site at

http://www.abnova.com/support/protocols.asp or product

page for detailed protocols

Form: Lyophilized

Preparation Method: Insect cells expression system

Purification: chromatographic

Purity: > 95% as determined by SDS-PAGE.

Activity: ED_{50} is 2-10 ng/mL, measured in a cell proliferation assay using primary HUVEC cells.

Storage Buffer: Lyophilized from a solution containing 50 mM acetic acid. Reconstitute the lyophilized powder

in 50 mM acetic acid to 50 ug/mL.

Storage Instruction: Lyophilized protein at room temperature for 3 weeks, should be stored at -20°C. Protein aliquots at 4°C for 2-7 days and should be stored at -20°C to -80°C. For long term storage it is recommended to add a carrier protein (0.1% HSA or BSA).

Avoid repeated freeze/thaw cycles.

Entrez GenelD: 7422

Gene Symbol: VEGFA

Gene Alias: MGC70609, VEGF, VEGF-A, VPF

Gene Summary: This gene is a member of the PDGF/VEGF growth factor family and encodes a protein that is often found as a disulfide linked homodimer. This protein is a glycosylated mitogen that specifically acts on endothelial cells and has various effects, including mediating increased vascular permeability, inducing angiogenesis, vasculogenesis and endothelial cell growth, promoting cell migration, and inhibiting apoptosis. Elevated levels of this protein is linked to POEMS syndrome, also known as Crow-Fukase syndrome. Mutations in this gene have been associated proliferative and nonproliferative retinopathy. Alternatively spliced transcript variants, encoding either freely secreted or cell-associated isoforms, have been characterized. There is also

evidence for the use of non-AUG (CUG) translation

initiation sites upstream of, and in-frame with the first AUG, leading to additional isoforms. [provided by