

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

VEGFA (Human) Recombinant Protein

Catalog Number: P9304

Regulation Status: For research use only (RUO)

Product Description: Human VEGFA recombinant

protein expressed in? Pichia Pastoris.

Host: Yeast

Theoretical MW (kDa): 42

Protocols: See our web site at

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

page for detailed protocols

Form: Liquid

Preparation Method: yeast expression system

Purification: chromatographic

Purity: > 90% as determined by SDS-PAGE.

 $\mbox{\bf Activity:} \ \mbox{ED}_{50} \ \mbox{is 2-6 ng/mL}, \ \mbox{determined by the ability to} \\ \mbox{stimulate 3H-Thymidine incorporation in human umbilical}$

vein endothelial cells.

Storage Buffer: Solution containing 20 mM PBS, pH

7.4.

Storage Instruction: Store at 15°C for two weeks 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 RefSeq1