

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

VEGF121 (Human) Recombinant Protein

Catalog Number: P9700

Regulation Status: For research use only (RUO)

Product Description: Human VEGF121 (P15692-9, 27 a.a. - 147 a.a.) partial recombinant protein expressed in

HEK293 cells.

Sequence:

APMAEGGGQNHHEVVKFMDVYQRSYCHPIETLVDIFQ EYPDEIEYIFKPSCVPLMRCGGCCNDEGLECVPTEES NITMQIMRIKPHQGQHIGEMSFLQHNKCECRPKKDRA RQEKCDKPRR

Host: Human

Theoretical MW (kDa): 14.07

Protocols: See our web site at

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

page for detailed protocols

Form: Liquid

Preparation Method: Mammalian cell (Expi293, high-

yield transient HEK293) expression system

Purity: > 95% as determined by Tris-Bis PAGE; > 90%

as determined by HPLC

Endotoxin Level: < 1 EU per 1 ug of protein

(determined by LAL method)

Activity: The EC_{50} was 21.7 ng/mL, messured by ELISA

at 0.5 ug/mL.

Recommend Usage: Biological Activity

ELISA SDS-PAGE

The optimal working dilution should be determined by

the end user.

Storage Buffer: In PBS pH 7.4

Storage Instruction: Store at -80°C for 12 Month. Aliquot to avoid repeated freezing and thawing.

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 promoting cell migration, and 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 RefSeq]