



SZABO SCANDIC

Part of Europa Biosite

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

ENG MaxPab rabbit polyclonal antibody (D01)

Catalog Number: H00002022-D01

Regulatory Status: For research use only (RUO)

Product Description: Rabbit polyclonal antibody raised against a full-length human ENG protein.

Immunogen: ENG (AAH14271.1, 1 a.a. ~ 658 a.a) full-length human protein.

Sequence:

MDRGTPLPLAVALLLASC SLSP TSAETVHCDLQPVGPE
RDEVYTTTSQVSKGCVAQAPNAILEVHVLFLEFPTGPS
QLELTQASKQNGTWPREVLLVLSVNSSVFLHLQALGI
PLHLAYNSSLVTFQEPGPVNTTELPSFKTQILEWAAE
RGPITSAEELNDPQSILLRLGQAQGSLSFCMLEASQD
MGRTLEWRPRTPALVRGCHLEGVAGHKEAHILRVLPG
HSAGPRTVTVKVELSCAPGDLDAVLILQGPPYVSWLID
ANHNMQIWTGGEYSFKIFPEKNIRGFKLPDTPQGLLGE
ARMLNASIVASFVELPLASIVSLHASSCGGRLQTS PAPI
QTTTPPKDTCSPELLMSLIQTKCADDAMTLVLKELVAH
LKCTITGLTFWDPSCEAEDRGDKFVLR SAYSSCGMQV
SASMISNEAVVNILSSSSPQRKKVHCLNMDLSLQGLGL
YLSPHFLQASNTIEPGQQSFVQVRVSPSVSEFLQLDS
CHLDLGPEGGTVELIQGRAAKGNCVLLSPSPEGDPR
FSLLHFYTVPIPKTGTLSCTVALRPKTGSQDQEVHRT
VFMRLNIISPDLGCTSKGLVLPVAVLGITFGAFLIGALLT
AALWYIYSHTRSPSKREPVVAVAAPASSESSSTNHSIG
STQSTPCSTSSMA

Host: Rabbit

Reactivity: Human

Applications: IP, WB-Tr

(See our web site product page for detailed applications information)

Protocols: See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Storage Buffer: No additive

Storage Instruction: Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 2022

Gene Symbol: ENG

Gene Alias: CD105, END, FLJ41744, HHT1, ORW, ORW1

Gene Summary: This gene encodes a homodimeric transmembrane protein which is a major glycoprotein of the vascular endothelium. This protein is a component of the transforming growth factor beta receptor complex and it binds TGFB1 and TGFB3 with high affinity. Mutations in this gene cause hereditary hemorrhagic telangiectasia, also known as Osler-Rendu-Weber syndrome 1, an autosomal dominant multisystemic vascular dysplasia. Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq]