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

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic

ENG (Human) Matched Antibody Pair

Catalog # : H00002022-AP11 **規格 :** [1 Set]

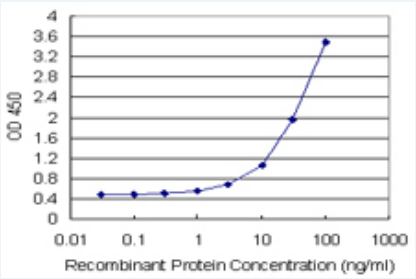
List All

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human ENG.

Reactivity: Human

Quality Control Testing: Standard curve using recombinant protein (H00002022-P02) as an analyte.



Sandwich ELISA detection sensitivity ranging from 1 ng/ml to 100 ng/ml.

Supplied Product: Antibody pair set content:
1. Capture antibody: rabbit MaxPab® affinity purified polyclonal anti-ENG (100 ug)
2. Detection antibody: mouse monoclonal anti-ENG, IgG1 Kappa (20 ug)
*Reagents are sufficient for at least 1-2 x 96 well plates using recommended protocols.

Storage Instruction: Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.

MSDS: Download

Applications

ELISA Pair (Recombinant protein)

Protocol Download

Gene Information

Entrez GeneID: 2022

Gene Name: ENG

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

Gene Description: endoglin

Omim ID: 131195, 187300

Gene Ontology: Hyperlink

Application Image

ELISA Pair (Recombinant protein)

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

Other Designations: CD105 antigen,OTTHUMP00000022221,Osler-Rendu-Weber syndrome 1

Related Disease

Aneurysm Aneurysm, Ruptured Arteriovenous Malformations Cardiovascular Diseases Central Nervous System Vascular Malformations Chromosome Aberrations Diabetes Mellitus, Type 2 Edema Epistaxis Genetic Predisposition to Disease Heart Defects, Congenital Hypertension, Pulmonary Intracranial Aneurysm Intracranial Arteriovenous Malformations Kidney Failure, Chronic Liver Cirrhosis Liver Diseases Lung Neoplasms Pre-Eclampsia

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