



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](https://www.linkedin.com/company/szaboscandic) 

ENG (Human) IP-WB Antibody Pair

Catalog # : H00002022-PW1

規格 : [1 Set]

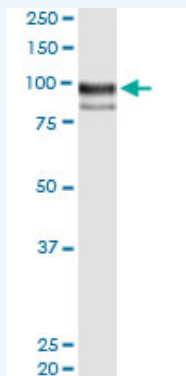
List All

Specification

Product Description: This IP-WB antibody pair set comes with one antibody for immunoprecipitation and another to detect the precipitated protein in western blot.

Reactivity: Human

Quality Control Testing: Immunoprecipitation-Western Blot (IP-WB)



Immunoprecipitation of ENG transfected lysate using rabbit polyclonal anti-ENG and Protein A Magnetic Bead ([U0007](#)), and immunoblotted with mouse polyclonal anti-ENG.

Supplied Product: Antibody pair set content:
1. Antibody pair for IP: rabbit polyclonal anti-ENG (300 ul)
2. Antibody pair for WB: mouse polyclonal anti-ENG (50 ul)

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

Immunoprecipitation-Western Blot

 [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

Immunoprecipitation-Western Blot

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](#)

... see more

[服務條款](#) | [隱私權政策](#) | [著作及商標](#) | [網站地圖](#)

©2016 亞諾法生技股份有限公司 Abnova Corporation. 版權所有.