

# CD105 (PN0640) Nb-FC recombinant antibody

CatalogNo: YA0047 Recombinant R

## Key Features

Reactivity

Human

Applications
• ELISA

#### **Recommended Dilution Ratios**

ELISA 1:5000-100000

#### **Storage**

**Storage\*** -15°C to -25°C/1 year(Avoid freeze / thaw cycles)

Formulation Phosphate-buffered solution

#### **Basic Information**

| Source              | Camel, chimeric fusion of Nanobody (VHH) and mouse IgG1 Fc domain , recombinantly produced from 293F cell |
|---------------------|---|
| Purification        | Camel, chimeric fusion of Nanobody (VHH) and mouse IgG1 Fc domain , recombinantly produced from 293F cell |
| <b>Clone Number</b> | PN0640  |

# Immunogen Information

| Immunogen   | Purified recombinant Human CD105   |
|-------------|--|
| Specificity | This recombinant monoclonal antibody can detects endogenous levels of CD105 protein. |

## **Target Information**

| Gene name                | ENG END   |              |                |
|--------------------------|---|--------------|----------------|
| Protein Name             | Endoglin (CD antigen CD105)<br><b>Organism</b>  | Gene ID      | UniProt ID     |
|                          | Human   | <u>3684;</u> | <u>P17813;</u> |
| Cellular<br>Localization | Cell membrane ; Single-pass type I membrane protein .   |              |                |
| Tissue specificity       | Predominantly expressed in monocytes and granulocytes (PubMed:1346576). Expressed in neutrophils (at protein level) (PubMed:2119347).   |              |                |
| Function                 | Disease:Defects in ENG are the cause of hereditary hemorrhagic telangiectasia type 1<br>(HHT1) [MIM:187300, 108010]; also known as Osler-Rendu-Weber syndrome 1 (ORW1).<br>HHT1 is an autosomal dominant multisystemic vascular dysplasia, characterized by<br>recurrent epistaxis, muco-cutaneous telangiectases, gastro-intestinal hemorrhage, and<br>pulmonary (PAVM), cerebral (CAVM) and hepatic arteriovenous malformations; all<br>secondary manifestations of the underlying vascular dysplasia. Although the first symptom<br>of HHT1 in children is generally nose bleed, there is an important clinical<br>heterogeneity.,Major glycoprotein of vascular endothelium. May play a critical role in the<br>binding of endothelial cells to integrins and/or other RGD receptors.,subunit:Homodimer<br>that forms an heteromeric complex with the signaling receptors for transforming growth<br>factor-beta: TGF-beta receptors I and/or II. It is able to bind TGF-beta 1, and 3 efficiently and<br>TGF-beta 2 less efficiently. Interacts with TCTEX1D4.,tissue specificity:Endoglin is restricted<br>to endothelial cells in all tissues except bone marrow., |              |                |

# Validation Data

# **Contact information**

| Orders:    | order@immunoway.com                      |
|------------|--|
| Support:   | tech@immunoway.com                       |
| Telephone: | 877-594-3616 (Toll Free), 408-747-0185   |
| Website:   | http://www.immunoway.com                 |
| Address:   | 2200 Ringwood Ave San Jose, CA 95131 USA |
|            |  |



Please scan the QR code to access additional product information: CD105 (PN0640) Nb-FC recombinant antibody

For Research Use Only. Not for Use in Diagnostic Procedures.

Antibody | ELISA Kits | Protein | Reagents