



Immobilized human VEGFR2-His-Avi dimer protein (CSP-25127-03) at 2 μ g/mL (100 μ L/well) can bind anti-human VEGFR2 monoclonal antibody with half maximal effective concentration (EC50) range of 13-51.8 ng/mL (QC tested).

Bioactivity – Ligand Binding





Immobilized human VEGF-A 2 μ g/mL (100 μ L/well) can bind human VEGFR2 dimer protein, His-Avi tag (Cat. No. CSP-25127-03), with half maximal effective concentration (EC50) range of 38.8-155.2 μ g/mL (QC tested).



MW: Molecular Weight marker reduced condition NR: VEGFR2 dimer under non-reduced condition

The migration range of the dimer protein with glycosylation under non-reducing condition is >190 kDa on SDS PAGE.



Bioactive, Human VEGFR2 Dimer, His-Avi Tag Product Code: CSP-25127-03 For Research Use Only (RUO)

Expression Host HEK293T

Purity

Greater than 90% dimer form as determined by SDS-PAGE under non-reducing condition

Protein Construct

VEGFR2 dimer protein contains a VEGFR2 extracellular domain (UniProt# P35968) fused with a proprietary dimer motif followed by a tandem His-Avi tag at the Cterminus. Expressed in HEK293T cell line.

SDS-Page Molecular Weight

187 kDa. The migration range of the dimer protein with glycosylation under non-reducing condition is >190 kDa on SDS PAGE.

Shipping Conditions

Frozen Dry Ice

Protein Name VEGFR2

Alternate Name(s)

Kinase insert domain receptor, KDR, cluster of differentiation 309, CD309, Fetal Liver Kinase 1, Flk1, VEGFR, VEGFR-2

Amino Acid Range

A20-E764

Formulation

0.22µm filtered PBS, pH 7.4

Stability & Storage -80°C

Background

Human vascular endothelial growth factor receptor 2 (VEGFR2) belongs to the Type IV receptor tyrosine kinase (RTK) family. VEGFR2 is a key receptor in the VEGF (Vascular Endothelial Growth Factor) signaling pathway, involved in angiogenesis, the formation of new blood vessels. VEGFR2 is also known as kinase insert domain receptor (KDR), cluster of differentiation 309 (CD309), and fetal liver kinase 1 (Flk1). VEGFR2, a Type I transmembrane protein, contains an extracellular domain with 7 immunoglobulin-like (Ig-like) domains, a single transmembrane domain, and an intracellular domain. VEGFR2 is mainly expressed on vascular endothelial cells and can bind VEGF-A and VEGF-D. VEGF binding by VEGFR2 causes it to homodimerize which is essential for it to stimulate cellular responses such as vasculogenesis and angiogenesis. In pathological conditions, ligand-independent dimerization of VEGFR2 can contribute to abnormal angiogenesis. Abnormal angiogenesis is associated with a variety of diseases such as tumor neovascularization, diabetic retinopathy, rheumatoid arthritis, and age-related macular degeneration. Abnormal angiogenesis is a major contributing factor in the growth and spread of a variety of cancers and inhibition of VEGFR2 activity offers a potential and promising approach to cancer therapy.