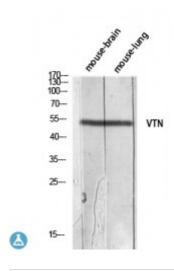


Anti-Vitronectin antibody



Description Rabbit polyclonal to Vitronectin.

Model STJ97659

Host Rabbit

Reactivity Human

Applications ELISA, WB

Immunogen Synthesized peptide derived from human Vitronectin.

Immunogen Region Internal

Gene ID <u>7448</u>

Gene Symbol VTN

Dilution range WB 1:500-1:2000ELISA 1:10000

Specificity Vitronectin Polyclonal Antibody detects endogenous levels of Vitronectin

protein.

Tissue Specificity Plasma.

Purification The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Note For Research Use Only (RUO).

Protein Name Vitronectin VN S-protein Serum-spreading factor V75 Vitronectin V65

subunit Vitronectin V10 subunit Somatomedin-B

Clonality Polyclonal

Conjugation Unconjugated

Isotype IgG

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Concentration 1 mg/ml

Store at -20°C, and avoid repeat freeze-thaw cycles. **Storage Instruction**

Database Links HGNC:12724OMIM:193190

Alternative Names Vitronectin VN S-protein Serum-spreading factor V75 Vitronectin V65

subunit Vitronectin V10 subunit Somatomedin-B

Function Vitronectin is a cell adhesion and spreading factor found in serum and tissues.

Vitronectin interact with glycosaminoglycans and proteoglycans. Is

recognized by certain members of the integrin family and serves as a cell-tosubstrate adhesion molecule. Inhibitor of the membrane-damaging effect of the terminal cytolytic complement pathway.; Somatomedin-B is a growth

hormone-dependent serum factor with protease-inhibiting activity.

Sequence and Domain Family The SMB domain mediates interaction with SERPINE1/PAI1. The heparin-

binding domain mediates interaction with insulin.

Cellular Localization Secreted, extracellular space.

Sulfated on 2 tyrosine residues. N- and O-glycosylated. Phosphorylation on Post-translational **Modifications**

Thr-69 and Thr-76 favors cell adhesion and spreading. It has been suggested that the active SMB domain may be permitted considerable disulfide bond heterogeneity or variability, thus two alternate disulfide patterns based on 3D

structures are described with 1 disulfide bond conserved in both.; Phosphorylation sites are present in the extracellular medium.

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