

# VTN (Human) Native Protein

Catalog # P4954

Size 100 ug

## Specification

<b>Product Description</b>	Human VTN (multimeric) native protein from human plasma.
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<b>Host</b>	Human
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<b>Theoretical MW (kDa)</b>	75 monomer
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<b>Form</b>	Liquid
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<b>Preparation Method</b>	Native protein purified from human plasma
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<b>Purification</b>	Non-denaturing chromatography
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<b>Concentration</b>	3.4 mg/mL
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<b>Storage Buffer</b>	In 0.05 M sodium phosphate, 0.1 M NaCl, pH 7.4
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<b>Storage Instruction</b>	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
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## Applications

- SDS-PAGE

## Gene Info — VTN

<b>Entrez GeneID</b>	<a href="#">7448</a>
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<b>Gene Name</b>	VTN
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<b>Gene Alias</b>	V75, VN, VNT
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<b>Gene Description</b>	vitronectin
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Omim ID [193190](#)

Gene Ontology [Hyperlink](#)

**Gene Summary**

The protein encoded by this gene is a member of the pexin family. It is found in serum and tissues and promotes cell adhesion and spreading, inhibits the membrane-damaging effect of the terminal cytolytic complement pathway, and binds to several serpin serine protease inhibitors. It is a secreted protein and exists in either a single chain form or a clipped, two chain form held together by a disulfide bond. [provided by RefSeq]

**Other Designations**

complement S-protein|epibolin|serum spreading factor|somatomedin B|vitronectin (serum spreading factor, somatomedin B, complement S-protein)

## Pathway

- [ECM-receptor interaction](#)
- [Focal adhesion](#)

## Disease

- [Cardiovascular Diseases](#)
- [Diabetes Mellitus](#)
- [Edema](#)
- [Genetic Predisposition to Disease](#)
- [Macular Degeneration](#)
- [Thyroid Neoplasms](#)