

F9 Antibody (Center) Blocking Peptide
Synthetic peptide
Catalog # BP16976c**Specification**

F9 Antibody (Center) Blocking Peptide - Product InformationPrimary Accession [P00740](#)**F9 Antibody (Center) Blocking Peptide - Additional Information****Gene ID** 2158**Other Names**

Coagulation factor IX, Christmas factor, Plasma thromboplastin component, PTC, Coagulation factor IXa light chain, Coagulation factor IXa heavy chain, F9

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

F9 Antibody (Center) Blocking Peptide - Protein Information**Name** F9**Function**

Factor IX is a vitamin K-dependent plasma protein that participates in the intrinsic pathway of blood coagulation by converting factor X to its active form in the presence of Ca(2+) ions, phospholipids, and factor VIIIa.

Cellular Location

Secreted

Tissue Location

Detected in blood plasma (at protein level) (PubMed:3857619, PubMed:8295821, PubMed:2592373, PubMed:9169594, PubMed:19846852). Synthesized primarily in the liver and secreted in plasma.

F9 Antibody (Center) Blocking Peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

F9 Antibody (Center) Blocking Peptide - Images

F9 Antibody (Center) Blocking Peptide - Background

This gene encodes vitamin K-dependent coagulation factor IX that circulates in the blood as an inactive zymogen. This factor is converted to an active form by factor XIa, which excises the activation peptide and thus generates a heavy chain and a light chain held together by one or more disulfide bonds. The role of this activated factor IX in the blood coagulation cascade is to activate factor X to its active form through interactions with Ca^{+2} ions, membrane phospholipids, and factor VIII. Alterations of this gene, including point mutations, insertions and deletions, cause factor IX deficiency, which is a recessive X-linked disorder, also called hemophilia B or Christmas disease.

F9 Antibody (Center) Blocking Peptide - References

Bailey, S.D., et al. Diabetes Care 33(10):2250-2253(2010) Yang, L., et al. J. Biol. Chem. 285(37):28488-28495(2010) Kao, C.Y., et al. Thromb. Haemost. 104(2):355-365(2010) Roberts, K.E., et al. Gastroenterology 139(1):130-139(2010) Arellano, A.R., et al. J. Thromb. Haemost. 8(5):1132-1134(2010)