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Recombinant Human Coagulation Factor IX/F9 Protein (His Tag)

Catalog No. PKSH031109

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Synonyms Coagulation factor 9;F9;Coagulation factor IX;Christmas factor;Plasma

thromboplastin component; Coagulation factor IXa light chain; Coagulation factor

IXa heavy chain;FIX;HEMB;P19;PTC;THPH8

Species Human

Expression Host

Sequence

Met 1-Thr 461

Accession

Calculated Molecular Weight

Observed molecular weight

Tag

HEK293 Cells

Met 1-Thr 461

AAB59620.1

50.0 kDa

60-80 kDa

C-His

Bioactivity Measured by its ability to cleave the peptide substrate, Z-D-Arg-Gly-Arg-pNA. The

specific activity is > 20pmols/min/ug.

Properties

Purity > 95 % as determined by reducing SDS-PAGE.

Endotoxin < 1.0 EU per µg of the protein as determined by the LAL method.

Storage Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to

-80°C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots

of reconstituted samples are stable at $< -20^{\circ}$ C for 3 months.

Shipping This product is provided as lyophilized powder which is shipped with ice packs.

Formulation Lyophilized from sterile PBS, pH 7.4

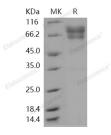
Normally 5 % - 8 % trehalose, mannitol and 0.01% Tween80 are added as

protectants before lyophilization.

Please refer to the specific buffer information in the printed manual.

Reconstitution Please refer to the printed manual for detailed information.

Data



> 95 % as determined by reducing SDS-PAGE.

For Research Use Only

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017

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Background

Coagulation factor IX, also known as Christmas factor, Plasma thromboplastin component and PTC, is a secreted protein which belongs to the peptidase S1 family. Coagulation factor IX / F9 contains two EGF-like domains, one Gla (gammacarboxy-glutamate) domain and one?peptidase S1 domain. Coagulation factor IX / F9 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 Ca2+ons, phospholipids, and factor VIIIa. Defects in Coagulation factor IX / F9 are the cause of thrombophilia due to factor IX defect which is a hemostatic disorder characterized by a tendency to thrombosis. Defects in Coagulation factor IX / F9 are also the cause of recessive X-linked hemophilia B (HEMB) which also known as Christmas disease.

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