abcam

Product datasheet

Human Factor XI ELISA Kit ab108834

1 References 画像数 1

製品の概要

製品名 Human Factor XI ELISA Kit

検出方法 Colorimetric

再現性

 サンプル
 N
 平均値
 SD
 CV%

 Overall
 4.9%

Inter-Assay(日差再現性)

Intra-Assay(同時再現性)

サンプル	N	平均値	SD	CV%
Overall				10%

サンプルの種類 Serum, Plasma

アッセイタイプ Sandwich (quantitative)

検出感度 0.69 ng/ml

検出範囲 1.563 ng/ml - 50 ng/ml

添加回収試験 96 % **全工程の試験時間** 4h 00m

ステップ Multiple steps standard assay

種交差性 交差種: Human

製品の概要 Abcam's Factor XI Human *in vitro* ELISA (Enzyme-Linked Immunosorbent Assay) kit is designed

for the quantitative measurement of Factor XI levels in human plasma and serum samples.

A Factor XI specific antibody has been precoated onto 96-well plates and blocked. Standards or test samples are added to the wells and subsequently a Factor XI specific biotinylated detection antibody is added and then followed by washing with wash buffer. Streptavidin-Peroxidase Complex is added and unbound conjugates are washed away with wash buffer. TMB is then used to visualize Streptavidin-Peroxidase enzymatic reaction. TMB is catalyzed by Streptavidin-Peroxidase to produce a blue color product that changes into yellow after adding acidic stop solution. The density of yellow coloration is directly proportional to the amount of Factor XI captured in plate.

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The entire kit may be stored at -20°C for long term storage before reconstitution - Avoid repeated freeze-thaw cycles.

試験プラットフォーム Microplate

製品の特性

保存方法 Store at -20°C. Please refer to protocols.

内容	1 x 96 tests
100X Streptavidin-Peroxidase Conjugate	1 x 80µl
10X Diluent M Concentrate	1 x 30ml
20X Wash Buffer Concentrate	2 x 30ml
50X Biotinylated Human Factor XI Antibody	1 x 120µl
Chromogen Substrate	1 x 7ml
Factor XI Microplate (12 x 8 well strips)	1 unit
Factor XI Standard	1 vial
Sealing Tapes	3 units
Stop Solution	1 x 11ml

機能 Factor XI triggers the middle phase of the intrinsic pathway of blood coagulation by activating

factor IX.

組織特異性 Isoform 2 is produced by platelets and megakaryocytes but absent from other blood cells.

関連疾患 Defects in F11 are the cause of factor XI deficiency (FA11D) [MIM:612416]; also known as

plasma thromboplastin antecedent deficiency or Rosenthal syndrome. It is a hemorrhagic disease characterized by reduced levels and activity of factor XI resulting in moderate bleeding symptoms, usually occurring after trauma or surgery. Patients usually do not present spontaneous bleeding

but women can present with menorrhagia. Hemorrhages are usually moderate.

配列類似性 Belongs to the peptidase S1 family. Plasma kallikrein subfamily.

Contains 4 apple domains.

Contains 1 peptidase S1 domain.

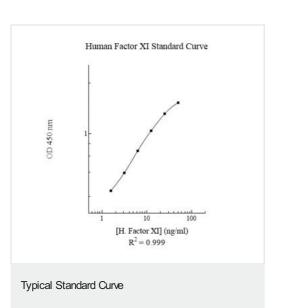
翻訳後修飾 Activated by factor XIIa (or XII), which cleaves each polypeptide after Arg-387 into the light chain,

which contains the active site, and the heavy chain, which associates with high molecular weight

(HMW) kininogen.

細胞内局在 Secreted.

画像



Representative Standard Curve using ab108834.

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