

**ZYMUTEST Factor IX****REF** RK032A

96 tests

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ELISA kit for Factor IX

FOR RESEARCH USE ONLY.

DO NOT USE IN DIAGNOSTIC PROCEDURES.

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**INTENDED USE:**

The ZYMUTEST Factor IX kit is a one step ELISA method for measuring Factor IX (FIX) in human plasma, or in any biological medium where FIX is present.

**This kit is for research use only and must not be used for patient diagnosis or treatment.**

**SUMMARY AND EXPLANATION:**

Factor IX (FIX) is a vitamin K dependent glycoprotein which participates in the middle phases of blood coagulation. When activated by Factor XI, in the presence of calcium, Factor IXa (FIXa) forms an active complex with FVIII, in the presence of calcium and phospholipids, the activated Factor IX (FIXa) forms, with the Factor VIII, an activator complex of Factor X in Factor Xa.

**PRINCIPLE:**

First, the immunoconjugate, a rabbit polyclonal antibody specific for FIX coupled to horse radish peroxidase (HRP), is introduced into the microwells of ELISA plate coated with a polyclonal antibody specific for FIX. Then, the diluted tested sample is immediately introduced, and the immunological reaction starts. The FIX present in the specimen, binds onto the polyclonal antibody coated solid phase through one epitope, and fixes the polyclonal antibody coupled to HRP through free epitopes. Following a washing step, the peroxidase substrate, 3,3', 5,5' - Tetramethylbenzidine (TMB), in presence of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), is introduced in the microwells and a blue colour develops. When the reaction is stopped with Sulfuric Acid, a yellow colour is obtained. The amount of colour developed is directly proportional to the concentration of human FIX in the tested sample.

**REAGENTS:**

- COAT**: Micro ELISA plate, containing 12 strips of 8 wells, coated with a rabbit polyclonal antibody specific for human FIX, stabilised and packed in an aluminium pouch hermetically sealed in presence of a desiccant. Contains BSA.
- SD**: 2 vials containing 50 mL of **Sample Diluent**, ready to use. Contains BSA.
- CAL**: 3 vials of **FIX Calibrator (calibrator plasma)**, lyophilised. Each vial must be restored with 2 mL of Sample Diluent to obtain a plasma containing a concentration "C" (expressed in %) of human FIX. This concentration "C" is greater than or equal to 100%, according to the lot. This calibrator is related to the NIBSC international standard. Contains BSA.
- CI**: 1 vial containing 0.5 mL of lyophilised **Plasma FIX Control I High** (human plasma). Contains BSA.
- CII**: 1 vial containing 0.5 mL of lyophilised **Plasma FIX Control II Low** (human plasma). Contains BSA.
- IC**: 3 vials of **Anti-(h)-FIX-HRP immunoconjugate**, a polyclonal antibody coupled to HRP, lyophilised. Contains BSA.
- CD**: 1 vial of 25 mL of **Conjugate Diluent**, ready to use. Contains BSA.
- WS**: 1 vial of 50 mL of 20 fold concentrated **Wash Solution**.
- TMB**: 1 vial of 25 mL peroxidase substrate: **3,3', 5,5' - Tetramethylbenzidine** containing hydrogen peroxide. Ready to use.
- SA**: 1 vial of 6 mL of **0.45M Sulfuric acid (Stop solution)**. Ready to use.

The exact concentration of controls and calibrator, and the acceptable interval concentration for the controls are indicated on the flyer provide in the kit. Concentrations vary from lot to lot. For the assay, refer to the values provided on the flyer of the kit.

Reagents **SD**, **CD** and **WS** contain 0.05% Kathon CG and reagent SA contains sulfuric acid, see WARNINGS AND PRECAUTIONS.

**WARNINGS AND PRECAUTIONS:**

- Biological products must be handled with all necessary precautions and considered as being potentially infectious.
- Waste should be disposed of in accordance with applicable local regulations.
- Use only the reagents from the same batch of kits. Do not mix reagents from different kit batches when performing an assay; they are optimized for each batch of kits.
- Handle the reagents with care to avoid contamination during use. If possible, avoid reagent evaporation during use by limiting the liquid-air exchange surface.
- To preserve reagent stability, seal the vials after use with their respective caps.
- Aging studies, conducted over a 3-week period at 30°C, show that the reagents can be shipped at room temperature over a short period of time, without degradation.
- The human plasma used to prepare the calibrator and controls I and II has been tested by recorded methods and is certified free of HIV antibodies, Hbs Antigen and HCV antibodies.
- The bovine plasma used to prepare the BSA has been tested by recorded methods and is certified free of infectious agents, in particular the causative agent of bovine spongiform encephalitis.
- Sulfuric acid, although diluted to 0.45M is caustic. As for any similar chemical, handle Sulfuric acid with great care. Wear protection glasses and gloves when handling. Avoid any skin and eye contact.
- For *in vitro* use.

**CD** **SD** **WS**:

H317 : May cause an allergic skin reaction.

**REAGENT PREPARATION AND STABILITY:**

Bring the kit at room temperature, at least 30 min before use. Store the unused reagents at 2-8°C. Vials are closed under vacuum. Remove carefully the stopper of lyophilized reagents, in order to avoid any loss of powder when opening the vials.

When appropriately used and stored, according to the recommended protocol and cautions, the kit can be used over a 1 month period, and strip by strip, if required.

- COAT (Micro ELISA plate)**: Open the aluminium pouch and take off the required amounts of 8 well strips for the test series. When out of the pouch, the strips must be used within 30 minutes. Unused strips can be stored at **2-8°C for 4 weeks** in their original aluminium pouch, in presence of the desiccant, hermetically closed and protected from any moisture, and stored in the provided plastic microplate storage bag (minigrip).
- SD (Sample Diluent)**: Ready to use. This reagent contains 0.05% Kathon CG. Stability of reagent, provided that any contamination or evaporation is avoided, kept in its original vial is:
  - 4 weeks** at 2-8°C.
- CAL (Factor IX Calibrator)**: Reconstitute each vial with 2 mL of "Sample Diluent", shake vigorously until fully dissolved, in order to obtain a plasma containing a FIX concentration "C", in % (**already diluted 1:50**). Stability of reconstituted reagent, provided that any contamination or evaporation is avoided, kept in its original vial is:
  - 24 hours** at room temperature (18-25°C).
  - 72 hours** at 2-8°C.
- CI (FIX Control I (human plasma, high))**: Reconstitute each vial with 0.5 mL of distilled water, shake vigorously until fully dissolved. Stability of reconstituted reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 24 hours** at room temperature (18-25°C).
  - 72 hours** at 2-8°C.
  - 2 months** frozen at -20°C or below.
- CII (FIX Control II (human plasma, low))**: Reconstitute each vial with 0.5 mL of distilled water, shake vigorously until fully dissolved. Stability of reconstituted reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 24 hours** at room temperature (18-25°C).
  - 72 hours** at 2-8°C.
  - 2 months** frozen at -20°C or below.
- IC (Anti-(h)-FIX-HRP immunoconjugate)**: Reconstitute each vial with 2 mL of **Conjugate Diluent** at least 15 min before use. Let the pellet to be completely dissolved before use, and shake the vial in order to homogenize the content. Stability of reconstituted reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 24 hours** at room temperature (18-25°C).
  - 4 weeks** at 2-8°C.
- CD (Conjugate Diluent)**: Ready to use. This reagent contains 0.05% Kathon CG. Stability of reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 4 weeks** at 2-8°C.
- WS (Wash Solution)**: Incubate, if necessary, the vial in a water bath, at 37°C, until complete dissolution of crystals. Shake the vial and dilute the amount required **1:20** in distilled water (the 50 mL contained in the vial allow to prepare 1 liter of Wash Solution). Stability of the wash solution, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 4 weeks** at 2-8°C.
 Stability of the dilute wash solution, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 7 days** at 2-8°C.
 This reagent contains 0.05% Kathon CG.
- TMB**: Ready to use. Stability of reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 4 weeks** at 2-8°C.
- SA (Stop Solution)**: Stop solution containing 0.45M sulfuric acid, ready to use. See CAUTIONS AND WARNINGS. Stability of reagent, provided that any contamination or evaporation is avoided, kept in its original vial:
  - 4 weeks** at 2-8°C.

**STORAGE CONDITIONS:**

Unopened reagents should be stored at 2-8 °C in their original packaging. Under these conditions, they can be used until the expiry date printed on the kit.

**REAGENTS AND MATERIALS REQUIRED BUT NOT PROVIDED:****Reagents:**

- Distilled water.

**Materials:**

- 8-channel pipettes** allowing dispensing volumes of 50-300 µL.
- Pipettes** at variable volumes from 0 to 20 µL, 20 to 200 µL and 200 to 1000 µL.
- Micro ELISA plate washing equipment and shaker.**
- Micro ELISA plate reader** with a wavelength set up at 450 nm.

## SPECIMEN COLLECTION AND PREPARATION:

Specimens should be prepared and stored in accordance with applicable local guidelines.

### • Specimens:

Human plasma obtained from anticoagulated blood (trisodium citrate), EDTA collected human plasma may also be used. The storage conditions are the same with citrated plasma.

### • Collection:

The blood (9 volumes) should be carefully collected onto the trisodium citrate anticoagulant (1 volume) (0.109 M) by clean venipuncture. Discard the first tube.

### • Centrifugation:

Within 2 hours, use a laboratory-validated method to obtain platelet-poor plasma, for example at least 15 minutes at 2500g at room temperature (18-25°C) and allow the plasma to settle in a plastic tube.

### • Plasma storage:

- 24 hours at room temperature (18-25°C).
- 6 months at -20°C.

Frozen plasma specimens should be thawed rapidly at 37°C, then shaken thoroughly and tested immediately. Resuspend any precipitate by shaking vigorously immediately after thawing and before use.

## PROCEDURE:

### Assay procedure:

1. Plasma Controls I and II must be tested diluted fifty fold (1:50), with Sample Diluent.

2. The plasma or specimen must be tested **diluted at 1:50** in the Sample Diluent. For expected FIX concentrations higher than "C" in %, dilute at 1:100 (D=100), or more. For low or very low FIX concentrations lower dilutions can be used.

3. Using the **Plasma FIX Calibrator**, with a FIX concentration "C" (greater than or equal to 100%, according to the lot), provided in the kit, prepare the following standard solutions:

FIX concentration (%)	C	C/2	C/4	C/10	C/20	0
Vol. of Plasma FIX calibrator	1 mL	0.5 mL	0.25 mL	0.1 mL	0.05 mL	0 mL
Vol. of Sample diluent	0 mL	0.5 mL	0.75 mL	0.9 mL	0.95 mL	1 mL

Mix for homogenization.

The standard dilutions are stable for **8 hours** at room temperature (18-25°C).

4. Remove the required number of strips from the aluminium pouch and put the strips in the frame provided. In the different wells of the micro ELISA plate introduce the reagents and perform the various assay steps as indicated on the following table:

Reagent	Volume	Procedure
Conjugate anti (h)-FIX-HRP. (Restored with 2 mL of Conjugate Diluent)	50 µL	Introduce the Anti-(h)-FIX- HRP immunoconjugate in the micro ELISA plate wells.
FIX calibrator or tested sample or sample diluent (blank)	200 µL	Introduce <b>immediately</b> the calibrator solutions or the tested samples in the corresponding micro ELISA plate well (a).
<b>Mix gently on a plate shaker or manually.</b> <b>Incubate for 1 hour at room temperature (18-25°C)</b>		
Wash Solution [WS] (20 fold diluted in distilled water before use)	300 µL	Proceed to 5 successive washings (b).
[TMB]/H <sub>2</sub> O <sub>2</sub> Substrate	200 µL	Immediately after the washing, introduce the substrate into the wells (b). <b>Note:</b> The substrate distribution, row by row, must be accurate (c, d).
<b>Incubate for exactly 5 minutes at room temperature (18-25 °C) (d).</b>		
0.45 M Sulfuric Acid [SA] (5)	50 µL	Following exactly the same time intervals than for the addition of substrate, stop the colour development by introducing the 0.45M sulfuric acid (c).
<b>Wait for 10 minutes in order to allow the colour to stabilize and measure absorbance at 450 nm (A450). Subtract the blank value (e) if necessary.</b>		

### Remarks:

- Distribute controls and tested specimen as rapidly as possible (≤10 minutes), in order to obtain a homogeneous immunological kinetics for antibodies binding. A too long delay between the distribution of the first and the last wells may induce an influence of immunological kinetics and produce wrong results.
- Never let the wells of ELISA plate empty between the addition of the reagents or following the washing step in order to preserve insoluble proteins. The next reagent must be added within 3 minutes, in order to prevent the plate from drying, which could damage the immobilized components. If necessary, keep the plate filled with Wash Solution and empty it just before the introduction of the next reagent. The washing instrument must be adjusted in order to wash the plates gently, and to avoid a too drastic emptying, which could lower plate reactivity.
- For addition of the [TMB] substrate, the time interval between each row must be accurate and exactly determined. It must be the same when stopping the reaction.
- Avoid letting the plate in the bright sunlight during incubations and more particularly during colour development. A micro-ELISA plate shaker can be used. An incubation temperature of (18-25°C) must be respected. Results are affected by a too high (>25°C) or too low (<18°C) temperature, and measured OD at 450 nm are then too high or too low. It has to be considered when analyzing the results. In the same way, if a microplate shaker is used, it should be used only at the beginning of each step (sample introduction, immunoconjugate introduction, stop solution introduction), for 1 to 2 minutes, in order to obtain a good homogeneity. OD 450 values generated in the assay are significantly increased if shaking is used throughout the incubation steps.
- For bichromatic readings, a reference wavelength at 690 nm or at 620 nm can be used.

## TWO STEP METHOD:

- The assay of FIX can also be performed with a "two step" method. The calibration curve is from 0 to C% (as for the one step method). The "FIX calibrator" [CAL] being reconstituted with 2 mL of Sample Diluent [SD].
- The immunoconjugate [IC] must be reconstituted with 7.5 mL of Conjugate Diluent [CD].
- Tested plasma must be assayed at a fifty fold (1:50) dilution or at higher dilutions in Sample Diluent [SD], if required.
- In each microwell of ELISA plate, introduce 200 µL of the calibration solution (prepared as for the one step method) or 200µL of the diluted 1:50 tested plasma (or more if necessary). Following a 1 hour incubation at room temperature (18-25°C), wash the plate and introduce 200 µL/well of immunoconjugate [IC]. Incubate 1 hour at room temperature (18-25°C), wash the plate, and introduce [TMB] substrate (200 µL/well). Stop the colour development developed for **exactly 5 min** with 50 µL of 0.45M sulfuric acid [SA] per well and OD at 450 nm is measured. Washing and operating cautions, as well as results interpretation, are the same as recommended for the one step method.

## RESULTS:

- For the manual endpoint method, plot the calibration curve, with the **OD 405 nm** along the Y-axis and the FIX concentration, expressed as %, along the X-axis by choosing the "best fit" interpolation mode (refer to the flyer in the kit).
- The concentration of FIX, in the test specimen, at the standard dilution (D:50), and expressed in %, is directly inferred from the calibration curve.
- For higher or lower dilution (ie. D), the concentration measured must be multiplied by the complementary dilution factor (D:50 ; i.e., **x2 for D:100**).
- For **controls I and II**, the concentrations are directly deduced from the calibration curve.
- Alternatively, an ELISA software (i.e., Dynex, Biolise, etc...) can be used for the calculation of concentrations.

**The results obtained should be for research use only and must not be used for patient diagnosis or treatment.**

## LIMITATIONS:

- To ensure optimum test performance and to meet the specifications, the technical instructions validated by HYPHEN BioMed should be followed carefully. The laboratory is responsible for validating any changes made to these instructions for use.
- Any reagent presenting an unusual appearance or showing signs of contamination must be rejected.
- Any suspicious samples or those showing signs of activation must be rejected.
- Any plasma displaying a coagulum or showing signs of contamination must be rejected.

## PERFORMANCE:

Detection threshold ≤ 5%.

Intra-assay: 3-8%.

Inter-assay: 5-10%.

No significant heparin interference up to 2 IU/mL.

## REFERENCES:

- Taran LD, "Factor IX of the blood coagulation system: a review", Biochemistry (Mosc.), 62(7):685-93, 1997.
- Amiral J. *et al.* "Application of enzyme immunoassays to coagulation testing", Clin Chem, 30(9):1512-6, 1984.

## SYMBOLS:

Symbols used and signs listed in the ISO 15223-1 standard, see Symbol definitions document.

Changements par rapport à la précédente version.