CE IVD

Precision in Hemostasis

Behnk Elektronik

BE Factor XII Deficient Plasma FXII

Immuno-depleted plasma for the determination of Factor XII activity in human plasma

REF 771612: DP (6 x 1 mL)

| INTENDED USE

This reagent is designated for professional use in laboratory (semi-automated or automated method). It allows the quantitative determination of Factor XII activity in citrated human plasma to assess the status of coagulation factors normally found in blood.
This test is realized with Behnk reagents as follows:
REF 771200, REF 771201: BE APTT K Kaolin + CaCl
REF 771250, REF 771251: BE APTT SL Silica + CaCl
REF 771700: BE Owren Buffer (Plasma dilution buffer)

| PRINCIPLE [1]

The test is based on the measurement of clotting time in the presence of cephalin and activator with a method in which all factors are present in excess (supplied by Factor XII Deficient Plasma) except Factor XII, which is derived from the sample to be tested.

GENERALITIES [2] [3] [4] [5]

Factor XII is involved at different stages:

- In the endogenous coagulation pathway
- In relation with Kallikreins in case of inflammation
- In fibrinolysis

There are pathological changes in the FXII in the following cases:

- In congenital deficits (autosomal recessive), the rate of Factor XII varies from 15 % to 80 % in heterozygotes and less than 1 % in Homozygotes.

Factor XII deficiency is not accompanied by haemorrhagic syndromes, which suggests that there is another mechanism substituting to the activation of Factor XII. It has not been demonstrated that this deficit increases risks of thrombosis.

REAGENTS

DP **FXII** Deficient Plasma FXII

Freeze dried citrated plasma without Factor XII, removed by selective immune adsorption.

Human Origin

According to 1272/2008 regulation, this reagent is not classified as dangerous.

| SAFETY CAUTIONS [13] [14]

- Refer to current Material Safety Data Sheet available on request or on www.behnk.de
- Each donor unit used to manufacture this product was tested and found non-reactive for HbsAg, antibody to Hepatitis C and antibody to HIV-1/HIV-2.
- However, no test method can offer complete assurance that infectious agents are absent. All specimens or reagents from biological origin should be handled as potentially infectious, in accordance with good laboratory practices using appropriate precautions.
- Waste disposal: Respect legislation in force in the country.

| Any serious incident that has occurred in connection with the device is notified to the manufacturer and the competent authority of the Member State in which the user and/or patient is based.

PREPARATION OF REAGENTS

Open the vial carefully and add exactly the volume of demineralised water stated on the label.
Cap the vial and let stand for 15 minutes at room temperature.
MXII gently by swirling and inverting before use, to homogenise the content.

| STABILITY AND STORAGE

Unopened vials, stored away from light at 2-8 °C are stable until the expiry date stated on the label.
Stability after reconstitution:

- 2-8 °C 8 hours
- On board Stability (OBS)* 4 hours
- 15-25 °C 4 hours

* 18-22 °C

Do not use any reagent after expiry date.

SAMPLES COLLECTION AND HANDLING [6] [7]

Plasma from careful venipuncture with anticoagulant ratio of 1/10 (sodium citrate solution 0.109 M). MXII immediately the blood with anticoagulant.
Avoid drawing with a syringe that could result in the formation of micro-clots.
Centrifuge for 10 minutes at 3000 g and extract supernatant.
Stability:

- 4 h at 2-25 °C
- 15 days at -20 °C, 1 month at -80 °C (if quickly frozen. Defrost at 37 °C until complete defrosting).

LIMITS [8] [9]

Anticoagulants present in the specimen may interfere with the Factor XII activity in the specimen.
The presence of Lupus anticoagulants may lead to an underestimation of Factor XII activity in the specimen.
For a more comprehensive review of factors affecting this assay, refer to the publication of Young D.S.

MATERIAL REQUIRED BUT NOT PROVIDED

Basic medical analysis laboratory equipment
Automated or semi-automated coagulation analyzer
Demineralised water

| REFERENCE RANGE [10] [11] [12]

Plasma (adult): Usually 60-150 %

In the new born, the Factor XII level is lower (approx. 50% of adult values).
After strenuous physical exercises, Factor XII can rise to 200-300 %.

Each laboratory should establish its own normal ranges for the population that it serves.

QUALITY CONTROL

REF 773100: BE Trol 1; REF 773101: BE Trol 2
Controls are required for checking the accuracy and reproducibility of the results.
The control intervals should be adapted to each laboratory's individual requirements.
Values obtained should fall within the defined limits.
Follow the applicable government regulations and local guidelines for quality control.

PROCEDURE

| Semi-automated systems
Pre-incubate CC reagent ($CaCl_2$ 0.025 M) of the APTT reagent 15 min to reach a temperature of 37 °C.
Dilute specimens and controls: 1/10 in BE Owren Buffer.
Calibrators: prepare dilutions as indicated in § Calibration.

- Diluted specimen (calibrators, controls, plasmas): 100 µL
- Deficient Plasma: 100 µl
- APTT reagent (mXII before use) 100 µl

Incubate for 180 sec at 37 °C

- CC reagent (37 °C): 100 µL

The automatic countdown timer will start immediately after CC reagent addition and stop when the clot is formed.
Automated method on Behnk Thrombolyzer series
Refer to the full detailed application specific to the automated system.

Note:

- Performances and stability data have been validated on Thrombolyzer Compact X (available on request).
- With manual procedure and on other automated coagulation analyzer, performances and stability data must be validated by user.
- Other validated applications or proposal applications are available on request.

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CALIBRATION

Use REF 775100: BE Cal Ref
Reference plasma traceable to WHO SSC/ISTH Secondary Coagulation Standard NIBSC code: SSCLOT4.
Semi-automatized method:
Prepare a calibration curve with dilution 1/10, 1/20, 1/40 and 1/80 in BE Owren Buffer. Measure in triplicate the clotting time of each level.
Automated method on Behnk Thrombolyzer series:
Perform a calibration with BE Cal Ref using automatic dilutions indicated in the specific application.

CALCULATION

Results are expressed in % of Deficient Factor according to the calibration curve.

PERFORMANCES

The studies were performed on Thrombolyzer Compact X.

Precision:

Within run N = 20	Level 1	Level 2
Mean (%)	143	84
S.D. (%)	5.7	5.0
C.V. %	4.0	5.9

Between run N = 20	Level 1	Level 2
Mean (%)	104	54
S.D. (%)	7.2	6.4
C.V. %	7.0	11.7

Detection limit: equivalent to 4 % of Factor XII

Measuring Range: from 25 % (QL) to 125 %

Interferences (APTT Silica, sec):

Turbidity	No interference up to 731 mg/dL of Triglycerides
Bilirubin	Positive interference from 133 µmol/L
Hemoglobin	No interference up to 261 µmol/L

Other substances may interfere with the results (see § Limits)

Calibration Stability: Make a new calibration when changing reagent batch, if quality control results are found out of the established range and after maintenance operations.

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| = Significant modifications

IFU_771612-EN_V02_20230923

Manufacturer	Expiry date	In Vitro Diagnostic	Temperature limitation	Catalogue number	See insert	Batch number	Store away from light	Sufficient for	Dilute with	Demineralized water	Biological hazard

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