

5-TEST EP-LMWH Anti-Xa starter set in compliance with European Pharmacopoeia

REF 5D-90452

Complete set of individual reagents for the measurement of heparin in aqueous solutions using an anti-FXa chromogenic assay for pharmaceutical preparations in compliance with Eur.

Pharmacopoeia.

For Research Use Only.

Not for Use in Diagnostic Procedures.

Mixed storage.

INTENDED USE:

This Heparin Anti-FXa method can be used as an endpoint or kinetic chromogenic assay for measuring the concentration of heparin in the range from 0.025-0.2 IU/mL. This method is to be used for the determination of anti-FXa activity of Low Molecular Weight Heparin following the recommendations of the European Pharmacopoeia.

TEST PRINCIPLE:

Heparin is a sulphated polysaccharide with a high affinity for antithrombin. Antithrombin complexed with heparin has a fast and potent inhibitory activity for coagulation factors IXa, Xa and IIa (Thrombin). FXa in excess, is neutralized in proportion to the amount of heparin (Heparin – AT - complex). The remaining amount of FXa hydrolyses the chromogenic substrate and liberates the chromophoric group pNA. The colour is then read photometrically at 405 nm. There is an inverse relationship between the concentration of heparin and colour development measured at 405 nm.

Heparin + AT \rightarrow [AT Hep.]

 $[\mathsf{AT}\ \mathsf{Hep.}] + [\mathsf{FXa}\ (\mathsf{excess})] \ \to \ [\mathsf{FXa}\text{-}\mathsf{AT}\text{-}\mathsf{Hep.}] + [\mathsf{residual}\ \mathsf{FXa}]$

 $[residual FXa] + Substrate \rightarrow Peptide + pNA$

REAGENTS INCLUDED:

5-BUFFER USP/Ph.Eur. Tris-NaCl-BSA Buffer salts pH 7.4 Ref. 5D-80433

5-BUFFER USP/Ph.Eur. Tris-NaCl-BSA Buffer salts pH 7.4, 1000 mL 0.050 M Tris Buffer pH 7.4 at 25°C, 0.150 M NaCl, 1.0% (w/v) BSA **Kit content:** 1 Pouch

Reconstitution: dissolve pouch content in 1000 mL distilled water. Buffer stability after reconstitution: 4 weeks at 2-8°C when protected from any contamination.

5-BUFFER USP/Ph.Eur. Tris-NaCl-EDTA Buffer salts pH 8.4

5-BUFFER USP/Ph.Eur. Tris-NaCl-EDTA Buffer salts pH 8.4, 500 mL 0.050 M Tris Buffer pH 8.4 at 25°C, 0.175 M NaCl, 0.0075 M EDTA **Kit content:** 1 Pouch

Reconstitution: dissolve pouch content in 500 mL distilled water. **Buffer stability after reconstitution**: 4 weeks at 2-8°C when protected from any contamination.

5-ENZYME Factor Xa (Bovine)

Ref. 5D-60217

Lyophilized Bovine FXa

Kit content: 1 Vial with 30 μg lyophilized bovine FXa, stabilizers Reconstitution: dissolve vial content in 2 mL distilled water

Stock concentration: $15 \,\mu g/mL$

Working concentration: 3 µg/mL, stock solution diluted 1:5 in 5-

BUFFER 5D-80433

Reagent stability after reconstitution, excluding any contamination or evaporation, and stored in the original vial, is of:

• 3 months at 2-8°C.

7 days at room temperature (18-25°C).

• 6 months frozen at -20°C or less.*

5-PROTEIN Antithrombin (Human)

Ref. 5D-60104

Lyophilized Human Antithrombin III

Kit content: 2 vials with each 10 IU lyophilized human antithrombin,

stabilizers. Specific activity > 6 IU/mg

Reconstitution: dissolve vial content in 5 mL distilled water

Stock concentration: 2 IU/mL

Working concentration: 1 IU/mL; stock solution diluted 1:2 in 5-

BUFFER 5D-80433.

Reagent stability after reconstitution, excluding any contamination or evaporation, and stored in the original vial, is of:

1 month at 2-8°C.

• 72 hours at room temperature (18-25°C).

• 6 months frozen at -20°C or less.*

5-CHROM-65 Chromogenic Factor Xa Substrate

Ref. 5D-30807

Chromogenic Substrate for Factor Xa: Z-D-Arg-Gly-Arg-pNA·2HCl **Kit content:** 1 Vial with 25 mg (39 μ mol/vial) synthetic chromogenic Factor Xa Substrate, highly purified and stabilized. Mannitol is added as a bulking agent.

Reconstitution: dissolve vial content in 13 mL distilled water

Stock concentration: 3 mM

Working concentration: 0.5 mM, stock solution diluted 1:6 in 5-

BUFFER 5D-80431

Reagent stability after reconstitution, excluding any contamination or evaporation, and stored in the original vial, is of:

• 3 months at 2-8°C

7 days at room temperature (18-25°C).

Do not freeze.

STORAGE CONDITIONS:

Unopened reagents must be stored in their original packaging at $2-8^{\circ}\text{C}$. They are then stable until the expiration date printed on the label

Stability of diluted reagents should be checked in the working conditions of the laboratory user.

*Thaw only once, as rapidly as possible at 37°C, adapting the incubation period to the volume of reagent. The stability of the thawed reagent should be checked under laboratory work conditions.

OTHER REAGENTS AND MATERIAL REQUIRED BUT NOT PROVIDED:

Reagents:

Distilled water

• Glacial acetic acid 20 % V/V

 USP, EP or International Standards from NIBSC, Internal Reference preparations

Materials:

 Spectrophotometer or automatic instrument for chromogenic assays

Stopwatch

Calibrated pipettes

Calibrated water bath or heating block

Plastic tubes or 96 well microplates

TEST PROCEDURE:

Prepare 4 independent calibration curves of minimum 4 points spanning 0.025 to 0.2 IU/mL of your reference Heparin Preparation in Buffer 5D-80433. Use Buffer 5D-80433 as a blank for the reaction.

Prepare 4 independent dilutions of your sample in Buffer 5D-80433.

Add 50 μL of Antithrombin III solution to 50 μL of sample or calibrator or blank. Mix gently and incubate 60 seconds at 37°C in a water bath or heating block.

Add 100 μL of Bovine Factor Xa solution and incubate 60 seconds at 37°C.

Add 250 μ L of FXa Chromogenic Substrate solution pre-warmed at 37°C and incubate for precisely 240 seconds at 37°C.

Stop the reaction with 375 μL acetic acid solution.

Measure the absorbance at 405 nm.

Plot the absorbance versus log of heparin concentrations in International Units/mL.

If necessary adjust the incubation time to give best dose-response curve

Determine the slope for the regression line of both reference and sample curves to calculate the potency.

Follow statistical analysis of results of biological assays and tests in compliance with Pharmacopoeia guidelines for parallel-line assays.

Reagent	Tube	
Antithrombin III 1 IU/mL	50 µL	
Reference, test sample or blank	50 μL	
Mix and incubate for 1 minute at 37°C		
Bovine Factor Xa 3 ug/mL	100 μL	
Mix and incubate for 1 minute at 37°C		
Chromogenic substrate 0.5 mM pre-warmed at 37°C	250 μL	
Mix and incubate at 37°C exactly for 4 minutes Stop the reaction by adding:		
Acetic acid 20%	375 µL	
Mix and measure the absorbance at 405nm against the corresponding blank.		

ALTERNATIVE METHODS

The assay can be miniaturized in 96 wells microplate.

Reagent	Microplate	
Antithrombin III 1 IU/mL	25 µL	
Reference, test sample or blank	25 µL	
Mix and incubate for 1 minute at 37°C		
Bovine Factor Xa 3 ug/mL	50 μL	
Mix and incubate for 1 minute at 37°C		
Chromogenic substrate 0.5 mM pre-warmed at 37°C	125 µL	
Mix and incubate at 37°C exactly for 4 minutes Stop the reaction by adding:		
Acetic acid 20%	25 µL	
Mix and measure the absorbance at 405nm against the corresponding blank.		

Note: Reagent concentrations may be adapted in order to obtain higher OD values. Please contact <u>info@5-diagnostics.com</u>.

Application protocols for automated analysers are available from $\underline{info@5-diagnostics.com}.$

ASSAY DETECTION RANGE:

0.025-0.2 IU/mL

APPLICATIONS:

Measurement of the specific anti-FXa activity of heparin in purified milieu using a two-stage assay. This procedure is in compliance with the quality control of Heparin preparations listed in European Pharmacopoeia.

REFERENCES:

European Pharmacopoeia 8.0:2.7.5. Assay of Heparin



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