

# 5-TEST USP-LMWH Anti-Ila starter set in compliance with US Pharmacopoeia

**REF** 5D-90455

Complete set of individual reagents for the measurement of heparin and heparin-like anticoagulants in buffer using an anti-Flla chromogenic assay for pharmaceutical preparations in compliance with US Pharmacopoeia.

For Research Use Only.
Not for Use in Diagnostic Procedures.
Mixed storage.

#### **INTENDED USE**

This Heparin Anti-Flla method can be used as an endpoint or kinetic chromogenic assay for measuring the concentration of heparin in the range of 0.015-0.075 IU/mL. The method is to be used for anti-Flla activity of Heparin following the recommendations of the US 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 Ila (Thrombin). The Heparin Anti-Flla method is a method based on the inhibition of a constant amount of Thrombin (Flla) by the tested molecule in presence of exogenous antithrombin and the simultaneous hydrolysis of a thrombin-specific chromogenic substrate by remaining active thrombin. 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

 $\begin{array}{lll} \mbox{Heparin} + \mbox{AT} & \rightarrow \mbox{[AT Hep.]} \\ \mbox{[AT Hep.]} + \mbox{[Ila (excess)]} & \rightarrow \mbox{[FIIa-AT-Hep.]} + \mbox{[residual FIIa]} \\ \mbox{[residual FIIa]} + \mbox{Substrate} & \rightarrow & \mbox{Peptide} + \mbox{pNA} \end{array}$ 

#### **REAGENTS INCLUDED**

5-BUFFER USP Tris-NaCl-PEG-6000 Buffer salts pH 7.4

**Ref**. 5D-80432

5-BUFFER USP Tris-NaCl-PEG-6000 Buffer salts pH 7.4, 1000 mL 0.050 M Tris buffer pH 7.4 at  $25^{\circ}$ C, 0.150 M NaCl, 0.10% (w/v) PEG-6000

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 Tris-NaCl Buffer salts pH 7.4

**Ref**. 5D-80430

5-BUFFER USP Tris-NaCl Buffer salts pH 7.4, 1000 mL 0.050 M Tris buffer pH 7.4 at 25°C, 0.150 M NaCl

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

Ref. 5D-80431

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 Thrombin (Human)

Ref. 5D-60230

Lyophilized Human Thrombin Reagent **Kit content:** 2 Vials, 100 IU per vial

Reconstitution: dissolve vial content in 2 mL distilled water

Stock concentration: 50 IU/mL

Working concentration: 5 IU/mL (stock solution diluted 1:10 in 5-

BUFFER 5D-80432)

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

- 21 days at 2-8°C.
- 7 days at room temperature (18-25°C).
- 6 months frozen at -30°C or less.\*

#### 5-PROTEIN Antithrombin (Human)

Ref. 5D-60104

Lyophilized Human Antithrombin III Reagent

Kit content: 1 Vial, 10 IU per vial

Reconstitution: dissolve vial content in 2 mL distilled water

Stock concentration: 5 IU/mL

Working concentration: 0.5 IU/mL (stock solution diluted 1:10 in 5-

BUFFER 5D-80432)

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

- 1 month at 2-8°C.
- 72 hours at room temperature (18-25°C).
- 6 months frozen at -20°C or less.\*

## 5-CHROM-38 Chromogenic Factor IIa Substrate

Ref. 5D-30805

Sequence: D-Phe-Pip-Arg-pNA

Kit content: 1 Vial, 25 mg per vial/ 45 µmol/vial

Reconstitution: dissolve vial content in 15 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:

- 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 their labelled temperature. 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
- Water bath or heating block
- Plastic tubes or 96 well microplates

#### **TEST PROCEDURE**

Prepare 4 independent calibration curves of minimum 4 points spanning 0.015-0.075 IU/mL of your reference Heparin Preparation in 5-BUFFER 5D-80430. Use 5-BUFFER 5D-80430 as a blank for the reaction.

Prepare 4 independent dilutions of your sample in 5-BUFFER 5D-80430.

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

Add 100  $\mu$ L of preheated Human Thrombin solution and incubate 60 seconds at 37°C.

Add 250  $\mu$ L of preheated FlIa Chromogenic Substrate solution and incubate exactly for 240 seconds at 37°C.

Stop the reaction with 250 µL acetic acid solution.

Measure the absorbance at 405 nm.

Plot the absorbance versus log of heparin concentrations in International Units/mL. In some cases log transformation of absorbance may be needed to obtain linearity.

If necessary adjust the incubation time to give best does-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 US Pharmacopoeia guidelines for parallel-line assays.

Reagent	Tubes
Antithrombin III 0.5 U/mL preheated at 37°C	50 μL
Reference, test sample or blank	50 μL
Mix and incubate for 1 minute at 37°C	
Human Thrombin 5 IU/mL preheated at 37°C	100 μL
Mix and incubate for 1 minute at 37°C	
Chromogenic substrate 0.5 mM preheated at 37°C	250 μL
Mix and incubate at 37°C exactly for 4 minutes Stop the reaction by adding:	
Acetic acid 20%	250 μL
Mix and measure the absorbance at 405 nm against the corresponding blank.	

#### **ALTERNATIVE METHODS**

The assay can be miniaturized in 96 wells microplate.

Reagent	Microplate
Antithrombin III 0.5 U/mL preheated at 37°C	25 μL
Reference, test sample or blank	25 μL
Mix and incubate for 1 minute at 37°C	
Human Thrombin 5 IU/mL preheated at 37°C	50 μL
Mix and incubate for 1 minute at 37°C	
Chromogenic substrate 0.5 mM preheated 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 405 nm agains corresponding blank.	t the

Application protocols for automated analysers are available from info@5-diagnostics.com.

#### **ASSAY DETECTION RANGE**

0.015-0.075 IU/mL

#### **APPLICATIONS**

Measurement of the specific anti-Ila activity of heparin and heparinlike anticoagulants in purified milieu using a two-stage assay. This procedure is in compliance with the quality control of Heparin preparations listed in USP.

#### **REFERENCES**

US Pharmacopoeia



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