BIOPHEN™ Arixtra® Calibrator

CE

IVD

REF 222501

CAL1 | CAL2 | CAL3 | CAL4 3 x 1 mL

Human plasmas for the calibration of Arixtra® assays by the anti-Xa method.

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INTENDED USE:

The BIOPHEN™ Arixtra® Calibrator kit consists of lyophilized human plasmas, spiked with Arixtra® (Fondaparinux) at various concentrations, for the calibration of Arixtra® assays.
It is titrated and optimized for the assay of Arixtra® by anti-Xa chromogenic technique.

SUMMARY AND EXPLANATION:

Technical:
These calibrators are used to establish the calibration curve of Arixtra® in plasma by anti-Xa chromogenic assays (BIOPHEN™ Heparin 3 and 6 and BIOPHEN™ Heparin LRT).
Clinical:
Arixtra® is used as an anticoagulant for curative or preventive indications. Measuring the Arixtra® concentration in patients' plasma can be used for monitoring the therapy and adjusting drug dosage.

REAGENTS:

CAL1 **Calibrator 1:** Lyophilized human plasma containing no Arixtra®.
CAL2 **Calibrator 2:** Lyophilized human plasma containing a titrated quantity of Arixtra® of approximately 0.5 µg/mL.
CAL3 **Calibrator 3:** Lyophilized human plasma containing a titrated quantity of Arixtra® of approximately 1.0 µg/mL.
CAL4 **Calibrator 4:** Lyophilized human plasma containing a titrated quantity of Arixtra® of approximately 1.5 µg/mL.

Calibrator plasmas contain stabilizing agents.

The calibrator concentrations may vary slightly from one batch to another. For the assay, see the exact values indicated on the flyer provided with the kit used.

CAL1 | CAL2 | CAL3 | CAL4 **3 vials of 1 mL**

WARNINGS AND PRECAUTIONS:

- Some reagents provided in these kits contain materials of human origin. Whenever human plasma is required for the preparation of these reagents, approved methods are used to test the plasma for the antibodies to HIV 1, HIV 2 and HCV, and for hepatitis B surface antigen, and results are found to be negative. However, no test method can offer complete assurance that infectious agents are absent. Therefore, users of reagents of these types must exercise extreme care in full compliance with safety precautions in the manipulation of these biological materials as if they were infectious.
- Waste should be disposed of in accordance with applicable local regulations.
- Use only the reagents from the same batch of kits.
- Aging studies show that the reagents can be shipped at room temperature without degradation.
- This device of *in vitro* diagnostic use is intended for professional use in the laboratory.

REAGENT PREPARATION:

Gently remove the freeze-drying stopper, to avoid any product loss when opening the vial.

CAL1 | CAL2 | CAL3 | CAL4 Reconstitute the contents of each vial with exactly **1 mL of distilled water.**
Shake vigorously until complete dissolution while avoiding formation of foam and load it directly on the analyzer following application guide instruction.
For manual method, allow to stabilize for 10 minutes at room temperature (18-25°C), homogenize before use.

This plasmatic reagent can be more or less turbid after reconstitution. This turbidity is mainly due to plasma lipids that, after freeze-drying, become "less" soluble and may form a slight deposit. If necessary, let each vial stabilize 10 minutes at room temperature and shake before use.

STORAGE AND STABILITY:

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.

CAL1 | CAL2 | CAL3 | CAL4 Reagent stability after reconstitution, free from any contamination or evaporation, and stored closed, is of:

- **7 days** at 2-8°C.
- **48 hours** at room temperature (18-25°C).
- **Stability on board of the analyzer: see the specific application.**

REAGENTS AND MATERIALS REQUIRED BUT NOT PROVIDED:

Reagents:
- Distilled water.

Materials:
- Calibrated pipettes.

TRACEABILITY:

The Arixtra® calibration plasmas are titrated relative to an Internal Standard Reference, initially qualified against a fresh preparation of Arixtra®, diluted in normal human citrated plasma.

QUALITY CONTROL:

The BIOPHEN™ Arixtra® Calibrator kit is used to establish a calibration curve to measure Arixtra® levels in plasma by anti-Xa chromogenic methods, such as those provided by BIOPHEN™ Heparin 3 and 6 (221003/221006) and BIOPHEN™ Heparin LRT (221011/221013/221015) kits.
The calibrator target values are determined from multi-reagent (BIOPHEN™ Heparin 3 and 6, BIOPHEN™ Heparin LRT) and multi-instrument (Sysmex CS-series or equivalent) tests.
The use of quality controls serves to validate method compliance, along with between-series assay homogeneity for a given batch of reagents.
Include the quality controls with each series, as per good laboratory practice, in order to validate the test.
A new calibration curve should be established, preferably for each test series, and at least for each new reagent batch, or after analyzer maintenance, or when the measured quality control values fall outside the acceptable range for the method.

LIMITATIONS:

- If the calibrators are used under measurement conditions other than those validated by HYPHEN BioMed, the test results may vary. The laboratory is responsible for validating the use of these calibrators in its own analytical system.
- Any reagent presenting an unusual appearance or showing signs of contamination must be rejected.

REFERENCES:

1. Walenga J.M. *et al.* Development of a Synthetic Heparin Pentasaccharide: Fondaparinux. Turk J Haematol. 2002.
2. Alexander G.G.Turpie. Selective factor Xa inhibition with fondaparinux: from concept to clinical benefit. European Heart Journal Supplements. 2008.
3. Castellone D.A. and Van Cott E.M. Laboratory monitoring of new anticoagulants. Am.J.Hematol. 2010.

SYMBOLS:

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