



BIOPHEN™ UFH Control Plasma

REF 223101 C1 C2 6 vials x 1 mL

BIOPHEN™ UFH Control C1

REF 224101 C1 12 vials x 1 mL

BIOPHEN™ UFH Control C2

REF 223901 C2 12 vials x 1 mL



English, revision: 11-2023

INTENDED USE:

For quality control of Unfractionated Heparin (UFH) assays, using a quantitative automated method.
This device of *in vitro* diagnostic use is intended for professional use in the laboratory.

SUMMARY AND EXPLANATION:

Technical:

These controls are proposed for the quality control of UFH anti-Xa chromogenic assays in plasma (BIOPHEN™ Heparin LRT).

Clinical:

Measuring the heparin concentration in patients' plasma may be used for monitoring the therapy and adjusting drug dosage.

REAGENTS:

- C1 Lyophilized human plasma containing approximately 0.20 IU/mL of UFH.
- C2 Lyophilized human plasma containing approximately 0.50 IU/mL of UFH.

Controls plasmas contain stabilizing agents.

The control 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.

The product is classified as non-hazardous and is not subject to labeling according to EC Regulation No. 1272/2008 [CLP].

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.
- Any serious incident that has occurred in relation to the device shall be reported to the manufacturer and the competent authority of the Member State in which the user and/or the patient is established.
- Summary of Safety and Performance (SSP) is available in the European database on medical devices (see Eudamed public website: <https://ec.europa.eu/tools/eudamed> or on request to HYPHEN BioMed).

REAGENT PREPARATION:

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

C1 C2 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.

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.

C1 C2 Reagent stability after reconstitution, free from any contamination or evaporation, and stored closed, is of:

- 7 days at 2-8°C.
- 60 days frozen at -20°C or less*
- Stability on board of the analyzer: see the specific Application Guide.

*Thaw only once, as rapidly as possible at 37°C and use immediately.

REAGENTS AND MATERIALS REQUIRED BUT NOT PROVIDED:

- Laboratory material.

TRACEABILITY:

Lot to lot variability measured on 3 lots is: %CV ≤ 10%. Controls are traceable to the WHO (World Health Organization) International Standard of reference for UFH. Certificate of traceability and uncertainty is available on the HYPHEN BioMed website:

Uncertainty			
C1	± 0.02 IU/mL	C2	± 0.03 IU/mL

QUALITY CONTROL:

For quality control of UFH assays anti-Xa chromogenic methods, with the BIOPHEN™ Heparin LRT (221011/221013/221015) kits. The target values are determined from multi-reagent and multi-instrument 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. If the controls fall outside of the acceptance range, the series of assays must be invalidated and the analyses repeated. Check all system parameters before repeating the series.

LIMITATIONS:

- If the controls 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 controls in its own analytical system.
- Any reagent presenting no limp appearance or showing signs of contamination must be rejected.

REFERENCES:

- Gray E. *et al.* Heparin and low-molecular-weight heparin. *Thromb Haemost.* 2008.

e-IFU (other languages) are available on www.hyphen-biomed.com. For customer support or Application Guides, please contact your local provider or distributor (see www.hyphen-biomed.com).

Changes compared to the previous version.

The following symbols may appear on the product labeling:

REF Catalogue number	LOT Batch code	IVD <i>In-vitro</i> diagnostic medical device
Rx Numerical < x > identification of reagent	See instructions for use	WHO STD WHO standard code
Temperature limitation	Manufacturer	YYYY-MM-DD Use by
CE XXXX CE marking of conformity with notified body ID number.	Reconstitution volume	CONTENTS Contents
Cx Numerical < x > identification of control	i-MA See instructions in Method Application guide	CONTAINS Contains
EXP Expiration date	Contains sufficient for <n> tests	UNIT Measurement unit
TARGET VALUE Target Value	Keep away from sunlight and heat	CALx Numerical < x > identification of calibrator
UDI Unique Device Identifier	Contains biological material of animal origin	Contains human blood or plasma derivatives
DANGER Danger	WARNING Warning	UK CA UKCA marking of conformity
Biological risks	ACCEPTANCE RANGE Acceptance range	