

Boditech Ferritin Control

INTENDED USE

Boditech Ferritin Control is intended for *in vitro* diagnostic use in the quality control of Ferritin Assay Kit. **For *in vitro* diagnostic use only.**

INTRODUCTION

The use of Boditech Ferritin Control may be considered as an objective assessment of the precision of Ferritin Assay Kits and is an integral part of Good Laboratory Practices. Boditech Ferritin Control is provided in lyophilized form.

SAFETY PRECAUTIONS AND WARNINGS

- For *in vitro* diagnostic use only.
- Do not pipette by mouth.
- Exercise proper precautions that would be normally required for handling laboratory reagents.
- Boditech Ferritin Control should not be used past the expiration date.
- Boditech Ferritin Control is solely designed to be provided instrument-specific calibration curves of Boditech Readers and Ferritin Assay Kits.
- Human source materials from which Boditech Ferritin Control is derived were tested at a donor level for the Human Immunodeficiency Virus (HIV 1, HIV 2) antibody, Hepatitis B Surface Antigen (HBsAg) and Hepatitis C Virus (HCV) antibody, and found to be NON-REACTIVE. FDA-approved methods have been used to conduct these tests. However, since no method can offer complete assurance as to the absence of infectious agents, these human source materials and all patient samples should be handled as though capable of transmitting infectious diseases and should be disposed of as hazardous wastes.

STORAGE AND STABILITY

- Storage and stability condition of Boditech Ferritin Control.

Temperature	Unopened	Opened (after reconstitution)	
		+2 to +8 °C	+2 to +8 °C
Expiration date	Until expiration date on the label.	1 day	7 days

- Close the opened Boditech Ferritin Control bottle tightly after use.
- Once the Boditech Ferritin Control was frozen, it should be used ONE TIME ONLY for test, because repeated freezing and thawing can result in the change of test values.
- After use, any residual product should NOT BE RETURNED to the original vial.
- Bacterial contamination of reconstituted Boditech Ferritin Control will cause reductions in the stability of many components. If bacterial contamination is suspected, the vial should be discarded and a fresh vial needs to be reconstituted.

INSTRUCTIONS FOR USE

Boditech Ferritin Control is supplied in lyophilized form.

1. Carefully reconstitute each vial of lyophilized with exactly 0.5 mL of sterilized distilled water.
2. Close the bottle and allow to stand for 30 minutes before use. Ensure contents are completely dissolved by swirling gently. Avoid formation of foam. Do not shake.

Dispose of any discarded materials in accordance with the requirements of your local waste management authorities.

In the event of damage to the package, contact the **Boditech Med Inc.'s Technical Services**.

MATERIALS SUPPLIED

REF CFPO-99

Boditech Ferritin Control Box (2 vials)	
Boditech Ferritin Control level 1 (0.5 mL)	1
Boditech Ferritin Control level 2 (0.5 mL)	1
Instruction For Use	1
Control value & Barcode Sheet	1

For Technical Assistance

Boditech Med Inc.'s Technical Services at

Tel: +82 (33) 243-1400

E-mail: sales@boditech.co.kr



Boditech Med Inc.

43, Geodudanji 1-gil, Dongnae-myeon,

Chuncheon-si, Gang-won-do, Korea

Tel: +82 -33-243-1400 / Fax: +82 -33-243-9373

www.boditech.co.kr



Obelis s.a

Bd. Général Wahis 53,

1030 Brussels, BELGIUM

Tel: +(32) -2-732-59-54

Fax: +(32) -2-732-60-03

E-Mail: mail@obelis.net

