Document No.: INS-AACO-EN

Revision Date : December 19, 2018 (Rev. 03)



Boditech HbA1c Control

INTENDED LISE

Boditech HbA1c Control is intended for in vitro diagnostic use in the quality control of HbA1c Assay Kit. For in vitro diagnostic use only.

INTRODUCTION

The use of Boditech HbA1c Control may be considered as an objective assessment of the precision of HbA1c Assay Kits and is an integral part of Good Laboratory Practices. Boditech HbA1c Control is provided in lyophilized form.

COMPONENTS

Boditech HbA1c Control consists of 'Boditech HbA1c Control level 1', 'Boditech HbA1c Control level 2', 'Instruction for Use' and 'Barcode Sheet'.

- The control contains HbA1c stock and Horse serum
- Each control vial packed in a box.

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 HbA1c Control should not be used past the expiration date.
- Boditech HbA1c Control is solely designed to be provided instrumentspecific calibration curves of Boditech Readers and HbA1c Assay Kits.
- Human source materials from which Boditech HbA1c 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 HbA1c Control.

	Unopened	Opened (after reconstitution)	
Temperature	+2 to +8 °C	+2 to +8 °C	-20 to -80 °C
Expiration date	Until expiration date on the label.	7 days	20 days

- Close the opened Boditech HbA1c Control bottle tightly after use.
- Once the Boditech HbA1c 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
- Bacterial contamination of reconstituted Boditech HbA1c 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 HbA1c Control is supplied in lyophilized form.

- 1. Carefully reconstitute each vial of lyophilized with exactly 0.5 mL of sterilized distilled water.
- 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.

Please refer to package inserts of the test cartridges for detailed test

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-96

Boditech HbA1c Control Box (2 vials) Boditech HbA1c Control level 1 (0.5 mL) 1 Boditech HbA1c Control level 2 (0.5 mL) 1 Instruction For Use 1 Control value & Barcode Sheet

OUALITY CONTROL

- Quality control tests are a part of the good testing practice to confirm the expected results and validity of the assay and should be performed at
- The control tests should be performed immediately after opening a new test lot to ensure the test performance is not altered.
- Quality control tests should also be performed whenever there is any question concerning the validity of the test results.

For Technical Assistance

Boditech Med Inc.'s Technical Services at

Tel: +82 (33) 243-1400 E-mail: sales@boditech.co.kr



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양식-GE02-15 (Rev.04) 1 / 1