

AXIS-SHIELD Homocysteine Control Kit



REF FHCY200

REF B08177

ENGLISH

Intended Use

The Axis-Shield Homocysteine Control Kit is intended to be used as an assayed quality control serum when used for the quantitative measurement of total L-homocysteine in human serum or plasma.

Summary and Principle

The use of quality control material is indicated as an objective assessment of the precision of methods and techniques in use and is an integral part of good laboratory practises. Three levels of control are available to allow performance monitoring within the clinical range.

Contents

3 vials (1.5 mL each) contain L-homocysteine in processed human serum in the following concentration ranges:


HOMOCYSTEINE ASSAY	PRODUCT CODE	CONTROL L	CONTROL M	CONTROL H
		RANGE (µmol/L)	RANGE (µmol/L)	RANGE (µmol/L)
Axis EIA Homocysteine Assay	FHCY100	5.6-8.4	10.0-15.0	20.0-30.0
Axis-Shield Liquid Stable (LS) 2-Part Homocysteine Reagent Assay	FHRWR100 / 200 / 1000	3.6-9.4	8.7-15.7	19.1-32.0
	FHRWAU100 / 200 / 1000			
	B08176			
3-Reagent Homocysteine Assay for SYNCHRON UniCel DxC	B08175			

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Precautions

IVD For *In Vitro* Diagnostic Use.

Caution: Federal law restricts this device to sale by or on the order of a physician

 The controls contain human sourced and potentially infectious components. Components sourced from human blood have been tested and found to be non-reactive for Hepatitis B Surface Antigen (HBsAg), HIV-1 Antigen (HIVAg) or HIV-1 RNA, HCV antibody, HIV-1/2 antibody, HTLV-1/2 antibody and Hepatitis B core Antibody (HBc). No known test method can offer complete assurance that products derived from human sources will not transmit infection. Therefore all human sourced materials should be considered potentially infectious. It is recommended that these materials be handled in accordance with Biosafety Level 2¹ or local/national guidelines on laboratory safety procedures. Controls contain <0.10% sodium azide as preservative.

Storage and Stability

The Axis-Shield Homocysteine Control Kit must be stored refrigerated (2-8°C). The controls are stable until the expiration date when stored and handled as directed. Do not use past expiration date.

Procedure and Handling

Each control should be treated as a patient sample and run in accordance with the instructions accompanying the instrument, kit or reagent being used.

Before sampling, allow the control to reach room temperature (18-25°C) and swirl gently before use to ensure homogeneity. Return the control to storage at 2-8°C immediately after use.

Dispose of any discarded materials in accordance with local waste management regulations.

Assignment of values

Individual laboratory means should fall within the corresponding acceptable range. Variations over time and between laboratories may be caused by differences in laboratory technique, instrumentation and reagents, or by test method modifications. It is recommended that each laboratory establishes its own mean and acceptable ranges and use those provided only as a guide.

Limitations

This product must not be used past the expiration date.






If there is evidence of microbial contamination or excessive turbidity in the product discard the vial.

This product is not intended for use as a calibrator.

Technical Support

For Technical Support please contact your local distributor or representative.

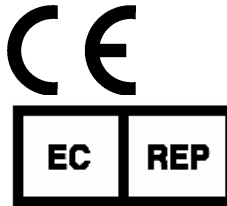
¹ US Department of Health and Human Services, *Biosafety in Microbiological and Biomedical Laboratories*, Fifth Edition. Washington, DC: US Government Printing Office December 2009.

REF	Product Code
IVD	<i>In Vitro</i> Diagnostic Medical Device
LOT	Lot number
	Consult Instructions For Use
Rx Only	For Prescription Use Only
	Manufacturer
	Use by
	Store at 2-8°C
	CAUTION, consult accompanying documents
CONTROL KIT	Control Kit
CONTROL L	Control Low, Medium, High (L, M, H)

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