



Hemoglobin A1c Control Set

Intended Use

This product is for the purpose of monitoring accuracy and precision in the quantitative determination of human hemoglobin A1c (HbA_{1c}) in blood by automated immunoassay. For *in vitro* diagnostic use only.

Summary

Throughout the circulatory life of the red cell, Hemoglobin A1c is formed continuously by the adduction of glucose to the N-terminal of the hemoglobin beta chain. This process, which is non-enzymatic, reflects the average exposure of hemoglobin to glucose over an extended period. In a classical study, Trivelli et al¹ showed Hemoglobin A1c in diabetic subjects to be elevated 2-3 fold over the levels found in normal individuals. Several investigators have recommended that Hemoglobin A1c serve as an indicator of metabolic control of the diabetic, since Hemoglobin A1c levels approach normal values for diabetics in metabolic control.^{2,3,4}

Hemoglobin A1c has been defined operationally as the "fast fraction" hemoglobins (HbA_{1a}, A_{1b}, A_{1c}) that elute first during column chromatography with cation-exchange resins. The non-glycosylated hemoglobin, which consists of the bulk of the hemoglobin has been designated HbA₀. The Pointe Scientific procedure utilizes an antigen and antibody reaction to directly determine the concentration of the HbA_{1c}.

Controls should be included each time patients are assayed for HbA_{1c} to verify that the assay has worked correctly. The mean value of the controls were obtained by assaying representative samples of the entire lot.

Reagents

The lyophilized HbA_{1c} controls are hemolysates prepared from packed human erythrocytes. The controls provide two levels of HbA_{1c}, one level in the normal range and the other level in the elevated range. Stabilizers are added to maintain hemoglobin in the reduced state providing complete control of the HbA_{1c} procedure.

Reagent Preparation

Reconstitute vials with 0.5ml deionized water. Gently mix for 10 minutes. Observe for undissolved material. The reconstituted controls may be dispensed in 0.1ml aliquots, sealed tightly and frozen at -20°C.

Reagent Storage

1. Store controls at 2-8°C. Stable until expiration date if sealed tightly. PROTECT FROM LIGHT AND HEAT.
2. Reconstituted controls retain their assigned values for at least three months if frozen. If not frozen, the reconstituted controls are stable at least one month if stored at 2-8°C and sealed tightly.
3. Do not freeze and thaw more than once.
4. Do not store in a self-defrosting freezer.

Precautions

1. This product is for *in vitro* diagnostic use only.
2. Although this product has been tested and found non-reactive for Hepatitis B Surface Antigen (HbsAG) and HIV, no known test can offer assurance that products derived from human blood will not transmit disease. Therefore all human serum products and patient specimens should be handled in the same manner as an infectious agent.

3. Do not pipette by mouth. Avoid contact with skin and mucous membranes.

Procedure

The lyophilized HbA_{1c} controls should be assayed in the same manner as blood specimens including the hemolysate procedure. Follow the directions that accompany the instrument, reagent kit used in the assay, and the instrument application instructions for the reagent set.

Materials Provided

1. Normal level control – Level 1
2. Elevated level control – Level 2

Materials Required but not Provided

1. Hemoglobin A1c Reagent Set.
2. Pipette capable of accurately delivering 0.5 ml.
3. Deionized water

Limitations

Things to look for that might cause inaccurate results are improper pipetting, inadequate mixing and poorly calibrated instruments.

Expected Values

The assayed limits are to be used as a guide in determining the accuracy of the assay procedure. The assay results for the controls should fall within the stated expected range. If they do not, the assay should be repeated, checking closely for the factors mentioned in "Limitations".

The expected values are lot-specific.

Lot Number: 900604-006 Exp. Date: 12/210

Level I	4.1-6.1 %
Level II	10.2-13.2%

Symbols

	Biological risk
	CE Mark (98/79 CE regulation)
	in vitro medical device
	Batch Code
	Use by
	Storage temperature limits
	Read instruction for use
	Manufactured by UK

Gesán Production s.r.l.

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