

Albumin

BCG Monoreagent



Cat.No	Package Size
103 016	9 x 10 mL/ Standard
103 005	2 x 50 mL/ Standard
103 000	4 x 100 mL/ Standard
103 004	2 x 500 mL/ Standard
103 002	(Hit I) 4 x 50 mL
103 003	Hit II) 4 x 100 mL
103 009	Hit II 6 x 100 mL
103 006	(AU) 8 x 70 mL

METHOD / TESTPRINCIPLE

Colorimetric determination of serum albumin by the photometric measurement of the bromocresol green (BCG) reaction with albumin

REAGENT

Composition (concentrations in the test)

Reagent :

Succinate buffer (pH 4.20)	140	mmol/L
Bromocresol green	0.26	mmol/L
Detergent	2.0	g/L

Standard : Std

Bovine albumin	4,0	g/dL
	40,0	g/L

Stability and Storage

When stored at 2-25° C and protected from light, the reagents are stable until the expiry date printed on the labels.

Preparation and Stability

The reagent is ready for use.

PRECAUTIONS

- For *in vitro* diagnostic use only.
- The standard contains < 0,95 g/L sodium azide. Avoid contact with skin and/or mucous membranes.

SAMPLES

Serum / Plasma

REFERENCE VALUES

<u>Ambulatory patients</u>	<u>Patients at rest</u>
3.8 - 5.5 g/dL	3.5 - 5.2 g/dL
38 - 55 g/L	35 - 52 g/L

Note: It is recommended for each laboratory to establish and maintain its own reference values. The given data here are only an indication.

ASSAY PROCEDURE

This reagent can be used manually (see method below) and on most analyzers. Applications are available on request.

Wavelength	:	578 / 623 nm
Temperature	:	37°C
Cuvette	:	1 cm light path

Read against Reagent Blank (RB)

	RB	Standard	Sample
Reagent	1 mL	1 mL	1 mL
Standard	-	10 µL	-
Sample	-	-	10 µL

Mix and read the absorbance (A) after 5 minutes incubation.

The final color is stable for at least 15 minutes.

CALCULATION

$$\frac{A_{\text{Sample}}}{A_{\text{Standard}}} \times C \quad \begin{array}{l} \text{g/dL} \Rightarrow C = 4.0 \\ \text{g/L} \Rightarrow C = 40 \end{array}$$

C = standard concentration.

CALIBRATORS & CONTROLS

For the calibration of automated analyzers Greiner Multicalibrator is recommended, for quality control use Greiner normal and abnormal controls Unitrol I and Unitrol II.

PERFORMANCE DATA

- Analytical range

The reagent is linear up to 6 g/dL (60 g/L).

- Detection limit

Determined according to SFBC protocol (Vassault and Coll.) protocol, the detection limit is 0.03 g/dL (0,3 g/L).

- Precision

Within-run reproducibility

N=11

	Mean	SD	CV	Unit
Control 1	3,43	0,029	0,85%	g/dL
Control 2	3,41	0,030	0,88%	g/dL
Patient	3,84	0,027	0,70%	g/dL

Between-run reproducibility

N=11

	Mean	SD	CV	Unit
Control 1	3,50	0,025	0,71%	g/dL
Control 2	3,38	0,033	0,98%	g/dL
Patient	3,79	0,039	1,03%	g/dL

- Correlation

A comparative study has been performed between the Greiner method and another commercial reagent on 34 human serum samples. The parameters of linear regression are as follows:

$$y = 1.008 x + 0.077 \quad \text{mg/dL} \quad r = 0.991$$

INTERFERENCES

- Ascorbic Acid: no interference until 50 mg/dL
- Bilirubin: no interference until 40 mg/dL,
- Hemoglobin: no interference until 500 mg/dL
- Triglycerides: no interference until 1000 mg/dL

BIBLIOGRAPHY

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4. Vassault, A. Grafmeyer, D. Naudin, et al., Protocole de validation de techniques. (Document B, stade 3) Ann. Biol. Clin., (1986), 44, 686.
5. Johnson, A. M., Rohlf, E. M., Silverman, L. M. Proteins, in Tietz Fundamentals of Clinical Chemistry, 5th edition, Burtis, C.A. et Ashwood, E.R. (W.B. Saunders eds. Philadelphia USA). 2001, 325.

SYMBOLS USED

IVD

For *in vitro* diagnostic medical use

LOT

Batch Code



Use by



Temperature limitation