

CALCIUM SYSTEM PACK

(ARSENazo III METHOD)

B Auto 400, Unicorn 480, Bonavera Chem 400, Beaconic B400 & Beaconic chem 400 (Fully Auto Biochemistry Analyzer)

Code	Product Name	Pack Size
UNI11	Calcium System Pack	4x50 ml

INTENDED USE

Diagnostic reagent for quantitative *in vitro* determination of Calcium in human serum.

CLINICAL SIGNIFICANCE

Calcium has numerous function within the body, not only as a structural factor in bones and teeth, but also in normal neuromuscular function and the clotting of blood.

Hypercalcemia may develop in patients with Paget's disease of bone and hyperparathyroidism. The cause of hypercalcemia in malignancy is an increased bone resorption either caused by metastasis or by humoral factors produced by the tumor cell.

In Rickets, Coeliac diseases, idiopathic steatorrhea, osteomalacia, tropical sprue and following surgical resection of the small intestine, serum calcium is often moderately reduced, usually in association with low plasma protein concentration.

PRINCIPLE

Arsenazo III combines with calcium ions at pH 6.5 to form a colored chromophore, the absorbance of which is measured at 650 nm (650-660nm) and is proportional to calcium concentration.

Arsenazo III has a high affinity ($k^{\circ} = 1 \times 10^7$) for calcium ions and shows no interference from other cations normally present in serum, plasma or urine.

REAGENT COMPOSITION

Reagent 1: Calcium Arsenazo (III) Reagent

Arsenazo III < 0.10 mmol/L

Buffer > 50 mmol/L

REAGENT PREPARATION

Reagents are liquid. ready to use.

STABILITY AND STORAGE

The unopened reagents are stable till the expiry date stated on the bottle and kit label when stored at 2 - 8°C.

On board Stability: Min 30 days if refrigerated (+8 - +14°C) and not contaminated.

SPECIMEN COLLECTION AND HANDLING

Use unheamolyse serum.

It is recommended to follow NCCLS procedures (or similar standardized conditions).



BEACON

Stability in serum

7 days at +20 - +25°C

3 weeks at +4 - +8°C

CALIBRATION

Calibration with the Beacon Multicalibrator is recommended.

QUALITY CONTROL

It's recommended to run normal and abnormal control sera to validate reagent performance.

UNIT CONVERSION

mg/dL x 0.25 = mmol/L

EXPECTED VALUES

Serum:

8.5- 11.0 mg/dL

It is recommended that each laboratory verify this range or derives reference interval for the population it serves.

PERFORMANCE DATA

Data contained within this section is representative of performance on Beacon System. Data obtained in your laboratory may differ from these values.

Limit of quantification: 0.6 mg/dL

Linearity: 15 mg/dL

Measuring range: 0.6 - 15 mg/dL

PRECISION

Intra-assay precision Within run (n=20)	Mean (mg/dL)	SD (mg/dL)	CV (%)
Sample 1	8.46	0.10	1.23
Sample 2	12.39	0.13	1.05

Inter-assay precision Run to run (n=20)	Mean (mg/dL)	SD (mg/dL)	CV (%)
Sample 1	8.26	0.08	1.00

COMPARISON

A comparison between Calcium System Pack (y) and commercially available test (x) using 20 samples gave following results:

$y = 1.0174x - 0.1521$ mg/dL

$r = 0.999$

INTERFERENCES

Following substances do not interfere:

haemoglobin upto 10 g/l bilirubin up to 40 mg/dl, triglycerides up to 500 mg/dl.

BEACON DIAGNOSTICS PVT. LTD. 424, NEW GIDC, KABILPORE, NAVSARI - 396 424. INDIA

WARNING AND PRECAUTIONS

For *in vitro* diagnostic use. To be handles by entitled and professionally educated person.

Reagents of the kits are not classified like dangerous.

MSDS will be provided on request.

WASTE MANAGEMENT

Please refer to local legal requirements.

Parameter For B Auto 400, Unicorn 480, Bonavera Chem 400, Beaconic B400 & Beaconic chem 400 (Fully Auto Biochemistry Analyzer)

Test Name	CALCIUM
Full Name	CALCIUM
PRI Wave	630 nm
SEC Wave	-
Assay/Point	1 Point End
Start	-
End	13
Decimal	2
Unit	mg/dL
Linearity Range Low	0.6
Linearity Range High	15
Sample Volume	4 µl
Reagent 1 (R1) Volume	200 µl
Reagent 1 (R2) Volume	-
Substrate Depleted/Abs.limit	-
Linearity	15 mg/dL
Out Of Linearity Range	-
Calibration Type	2 Point linear
Points	2
Blank Type	Reagent
Concentration Blank	0.00
Concentration Std	Refer calibrator value sheet

NOTE

The program is made as per the in house testing, it can be modified as per requirements.

Clinical diagnosis should not be made on findings of a single test results, but both clinical and laboratory data.

REFERENCES

1. Bishop , M. C. Dubeb - on Laufen, J.L., Burtis, Carl Aa and Ashwood, Tietz 110,61.

Symbols Used On Labels



Catalogue
Number



Manufacturer



See Instruction
for Use



Lot Number



Content



Storage Temperature



Expiry Date



In Vitro Diagnostics

BEA/24/CAL/UN/IFU Ver: 02
28/01/2026

