(Arsenazo Method)



This reagent kit is for quantitative estimation of Calcium (Ca^{**}) in serum.

PRINCIPLE:

In an alkaline medium, calcium reacts with Arsenazo III & forms a purple coloured complex. Intensity of colour is measured at 650 nm & this corresponds to calcium concentration. The presence of 8-Hydroxy quinoline in the reagent prevents interference by magnesium.

Arsenazo + 2 Ca⁺⁺ → Arsenazo (Ca⁺⁺)

CLINICAL SIGNIFICANCE:

Clinical conditions like hyperparathyroidism, multiple myeloma, neoplasia of bone and parathyroidism, & conditions of rapid demineralization of bone results in elevated calcium values. Lower calcium values seen in case of hypoparathyroidism, tetany & occasionally with nephrosis and pancreatitis.

SPECIMEN COLLECTION AND STORAGE:

- 1. Fresh Unhemolysed Serum is the preferred specimen.
- 2. Heparinsed plasma may also be used.
- 3. Don't use E.D.T.A. plasma.

REAGENT: Also available as mono test packs.

	25 ml	25x1 ml
	No. of Bottles	
Reagent	1	25
Standard (10 mg/dl)	1	1

PREPARATION OF WORKING REAGENT:

Ready to use.

REAGENT STORAGE AND STABILITY:

The Reagent and Standard are stable at R.T. till the expiry date mentioned on the labels if external contamination of calcium is avoided.

GENERAL INSTRUMENT PARAMETERS:

Reaction Type : End Point
Standard Concentration : 10 mg/dl
Slope of Reaction : Increasing
Units : mg/dl

Wavelength : 650nm (630-670) nm

Incubation : 5 Minutes
Flowcell Temperature : 37° C
Reagent Volume : 1.0ml

Zero Setting : Reagent blank Sample Volume : 10 μl (0.01 ml) Path length : 1.0 cm

For 1 ml

Dispense	Blank	Std.	Test	
Reagent	1.0ml	1.0ml	1.0ml	
Standard	-	10 µl) μl	
Sample	-	-	10 µl	

Incubate for 5 min. at R. T. (25°-30°), then mix and read at 650 nm (630-670) nm or against Red filter

STABILITY OF REACTION MIXTURE:

The colour of the final reaction mixture is stable for one hour (when protected from light)

LINEARITY:

This method is linear for calcium values up to 20 mg/dl. For sample with values higher than 20 mg/dl, dilute the sample using normal saline and repeat the assay. Apply proper dilution factor while calculation.

CALCULATIONS:

	Abs. Test
Calcium Concentration (mg/dl) =	x 10
, ,	Abs. std

EXPECTED VALUES:

| Serum: adults | 8.4 to 11.5 mg/dl | Urine: | women | <250 mg/24 hours | <300 mg/24 hours |

Limitation interference:

No significant interference was observed from Bilirubin up to 20 mg/dl (Both conjugated and unconjugated Bilirubin) Hemoglobin up to 50 mg/dl,Lipemia as Triglycerides up to 2000 mg/dl, Ascorbic acid up to 50 mg/dl.

Measuring Range:

Measuring range: 0.2-25 mg/dl. Determine samples having higher concentrations manually dilute with 0.9% NaCl or distilled/deionized water (e.g. 1 + 1). Multiply the result by the appropriate dilution factor (e.g. 2). Analytical sensitivity (lower detection limit):Detection limit: 0.2 mg/dl.

NOTES:

- 1. All glassware and cuvettes should be rinsed with 30% Hydrochloric acid and again rinse with high quality distilled water thoroughly and repeatedly before use.
- If a larger volume of working reagent is required for the absorbance reading, requisite volumes can be taken in multiples keeping the same ratio of reagent to specimen or standard.
- 3. Dilute lipemic samples with normal saline (made in deionised water) and multiply with dilution factor.
- 4. Protect the kit from direct sunlight.
- 5. For accuracy of results, procedure has to be followed meticulously.
- 6. As with all diagnostic procedures the physician should evaluate data obtained by way of this kit in light of other clinical information.

BIBLIOGRAPHY:

- Young, S.D. Pestaner, L.C., and Gibbermon, V., Clin, Chem. Vol. 21, No.5, (1975).
- Tietz, N. W. Fundamentals of Clinical Chemistry, 2nd edition. W.B. Saunders Co., Toronto to (1976) page 901

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d.	$\overline{\mathbb{A}}$	Attention,see instructions for use	i	Consult Instructions For Use
a	IVD	For in vitro diagnostic use only	REF	Catalog #
in	1	Store at RT	LOT	Lot Number
11	8	Do not use if package is damaged	M	Date of Manufacturing
	3	Manufacturer		Use by