

# Calcium AS FS\*

**Diagnostic reagent for quantitative in vitro determination of calcium in serum, plasma or urine on photometric systems**

## Order Information

Cat. No.	Kit size
1 1130 99 10 021	R 5 x 25 mL + 1 x 3 mL Standard
1 1130 99 10 026	R 6 x 100 mL
1 1130 99 10 023	R 1 x 1000 mL
1 1130 99 10 704	R 8 x 50 mL
1 1100 99 10 030	6 x 3 mL Standard

## Summary [1,2]

Calcium plays an essential role in many cell functions: intracellularly in muscle contraction and glycogen metabolism, extracellularly, in bone mineralization, in blood coagulation and in transmission of nerve impulses. Calcium is present in plasma in three forms: free, bound to proteins or complexed with anions as phosphate, citrate and bicarbonate. Decreased total calcium levels can be associated with diseases of the bone apparatus (especially osteoporosis), kidney diseases (especially under dialysis), defective intestinal absorption and hypoparathyroidism. Increased total calcium can be measured in hyperparathyroidism, malignant diseases with metastases and sarcoidosis. Calcium measurements also help in monitoring of calcium supplementation mainly in the prevention of osteoporosis.

## Method

Photometric test using arsenazo III

## Principle

Calcium with arsenazo III at neutral pH yields a blue colored complex, whose intensity is proportional to the calcium concentration. Interference by magnesium is eliminated by addition of 8-hydroxyquinoline-5-sulfonic acid.

## Reagents

### Components and Concentrations

#### Reagent:

Phosphate buffer	pH 7.5	50 mmol/L
8-Hydroxyquinoline-5-sulfonic acid		5 mmol/L
Arsenazo III		120 µmol/L

#### Standard:

10 mg/dL  
(2.5 mmol/L)

### Storage Instructions and Reagent Stability

The reagent is stable up to the end of the indicated month of expiry, if stored at 2 – 8 °C and contamination is avoided. Do not freeze the reagent!

The standard is stable up to the end of the indicated month of expiry, if stored at 2 – 25 °C.

### Warnings and Precautions

- As calcium is an ubiquitous ion, essential precaution must be taken against accidental contamination. Only use disposable materials.
- Traces of chelating agent, such as EDTA can prevent the formation of the colored complex.
- The reagent contains sodium azide (0.95 g/L) as preservative. Do not swallow! Avoid contact with skin and mucous membranes.

- In very rare cases, samples of patients with gammopathy might give falsified results [7].
- Please refer to the safety data sheets and take the necessary precautions for the use of laboratory reagents. For diagnostic purposes, the results should always be assessed with the patient's medical history, clinical examinations and other findings.
- For professional use only!

### Waste Management

Please refer to local legal requirements.

### Reagent Preparation

The reagent and the standard are ready-to-use.

### Materials required but not provided

NaCl solution 9 g/L.

General laboratory equipment

## Specimen

Serum, heparin plasma or urine

Do not use EDTA plasma.

### Stability [5]

in Serum/Plasma:	7 days	at	20 – 25 °C
	3 weeks	at	4 – 8 °C
	8 months	at	–20 °C
in Urine:	2 days	at	20 – 25 °C
	4 days	at	4 – 8 °C
	3 weeks	at	–20 °C

Add 10 mL of concentrated HCl to 24 h urine and heat the specimen to dissolve calcium oxalate.

Discard contaminated specimens. Freeze only once!

## Assay Procedure

**Application sheets for automated systems are available on request.**

Wavelength	650 nm, Hg 623 nm (630 – 670 nm)
Optical path	1 cm
Temperature	20 – 25 °C / 37 °C
Measurement	against reagent blank

	Blank	Sample or standard
Sample or standard	-	10 µL
Dist. water	10 µL	-
Reagent	1000 µL	1000 µL
Mix, incubate for 5 min. and read absorbance against reagent blank.		

## Calculation

With standard or calibrator

$$\text{Calcium [mg/dL]} = \frac{A_{\text{Sample}}}{A_{\text{Std / Cal}}} \times \text{Conc. Std / Cal [mg/dL]}$$

### Conversion factor

$$\text{Calcium [mg/dL]} \times 0.2495 = \text{Calcium [mmol/L]}$$

$$\text{Calcium/U [mg/24 h]} \times 0.025 = \text{Calcium/U [mmol/24 h]}$$

## Calibrators and Controls

For calibration of automated photometric systems the DiaSys TruCal U calibrator is recommended. This method has been standardized against the reference method Atomic Absorption Spectrometry (AAS). For internal quality control DiaSys TruLab N and P or TruLab Urine controls should be assayed. Each laboratory should establish corrective actions in case of deviations in control recovery.

	Cat. No.	Kit size
TruCal U	5 9100 99 10 063	20 x 3 mL
	5 9100 99 10 064	6 x 3 mL
TruLab N	5 9000 99 10 062	20 x 5 mL
	5 9000 99 10 061	6 x 5 mL
TruLab P	5 9050 99 10 062	20 x 5 mL
	5 9050 99 10 061	6 x 5 mL
TruLab Urine Level 1	5 9170 99 10 062	20 x 5 mL
	5 9170 99 10 061	6 x 5 mL
TruLab Urine Level 2	5 9180 99 10 062	20 x 5 mL
	5 9180 99 10 061	6 x 5 mL

## Performance Characteristics

### Measuring range

The test has been developed to determine calcium concentrations within a measuring range from 0.04 – 20 mg/dL (0.01 – 5 mmol/L). When values exceed this range samples should be diluted 1 + 1 with NaCl solution (9 g/L) and the result multiplied by 2.

### Specificity / Interferences

No interference was observed by ascorbic acid up to 30 mg/dL, bilirubin up to 40 mg/dL, hemoglobin up to 500 mg/dL, lipemia up to 2,000 mg/dL triglycerides and magnesium up to 15 mg/dL. Strontium salts in medicine may lead to strongly increased calcium values. For further information on interfering substances refer to Young DS [6].

### Sensitivity / Limit of Detection

The lower limit of detection is 0.04 mg/dL (0.01 mmol/L).

### Precision (at 20 – 25 °C)

Intra-assay precision n = 20	Mean [mg/dL]	SD [mg/dL]	CV [%]
Sample 1	8.79	0.09	1.04
Sample 2	12.5	0.15	1.20
Sample 3	14.0	0.24	1.73

Inter-assay precision n = 20	Mean [mg/dL]	SD [mg/dL]	CV [%]
Sample 1	8.82	0.18	2.01
Sample 2	12.3	0.11	0.90
Sample 3	13.7	0.26	1.92

### Method Comparison

A comparison between DiaSys Calcium FS (y) and a commercially available test (x) using 70 samples gave following results:

$$y = 1.02 x - 0.20; r = 0.999$$

## Reference Range

Serum/Plasma [2]:

$$8.6 - 10.3 \text{ mg/dL} \quad (2.15 - 2.57 \text{ mmol/L})$$

Urine [1]: Women < 250 mg/24 h (6.24 mmol/24 h)  
Men < 300 mg/24 h (7.49 mmol/24 h)

Each laboratory should check if the reference ranges are transferable to its own patient population and determine own reference ranges if necessary.

## Literature

1. Thomas L. Clinical Laboratory Diagnostics. 1<sup>st</sup> ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998. p. 231-241.
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3. Michaylova V, Ilkova P. Photometric determination of micro amounts of calcium with arsenazo III. Anal Chim Acta 1971;53:194-8.
4. Bauer PJ. Affinity and stoichiometry of calcium binding by arsenazo III. Anal Biochem 1981;110:61-72.
5. Guder WG, Zawta B et al. The Quality of Diagnostic Samples. 1<sup>st</sup> ed. Darmstadt: GIT Verlag; 2001. p. 20-1 and p. 50-1.
6. Young DS. Effects of Drugs on Clinical Laboratory Tests. 5th ed. Volume 1 and 2. Washington, DC: The American Association for Clinical Chemistry Press 2000.
7. Bakker AJ, Mücke M. Gammopathy interference in clinical chemistry assays: mechanisms, detection and prevention. ClinChemLabMed 2007;45(9):1240-1243.

## Manufacturer



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