COD 11592 50 mL	COD 11593 200 mL	COD 11598 500 mL
STORE AT 2-8°C		
Reagents for measurement of ALP concentration Only for in vitro use in the clinical laboratory		

ALKALINE PHOSPHATASE (ALP) - AMP





ALKALINE PHOSPHATASE (ALP) - AMP 2-AMINO-2-METHYL-1-PROPANOL BUFFER (IFCC)

PRINCIPLE OF THE METHOD

Alkaline phosphatase (ALP) catalyzes in alkaline medium the transfer of the phosphate group from 4-nitrophenylphosphate to 2-amino-2-methyl-1-propanol (AMP), liberating 4-nitrophenol. The catalytic concentration is determined from the rate of 4-nitrophenol formation, measured at

CONTENTS

	COD 11592	COD 11593	COD 11598
A. Reagent	1 x 40 mL	1 x 160 mL	4 x 100 mL
B. Reagent	1 x 10 mL	1 x 40 mL	2 x 50 mL

COMPOSITION

- Reagent: 2-Amino-2-methyl-1-propanol 0.4 mol/L, zinc sulfate 1.2 mmol/L, N-hydroxyethylethylenediaminetriacetic acid 2.5 mmol/L, magnesium acetate 2.5 mmol/L, pH
- B. Reagent: 4-Nitrophenylphosphate 60 mmol/L.

STORAGE

Store at 2-8°C

Reagents are stable until the expiry date shown on the label when stored tightly closed and if contaminations are prevented during their use

Indications of deterioration:

Reagents: Presence of particulate material, turbidity, absorbance of the blank over 1.200 at 405 nm (1 cm cuvette).

WARNING AND PRECAUTIONS

Exercise the normal precautions required for handling all laboratory reagents. Safety data sheet available for professional user on request. Disposal of all waste material should be in accordance with local guidelines. Any serious incident that might occur in relation to the device shall be reported to BioSystems S.A.

REAGENT PREPARATION

Working Reagent:

- Cod. 11592 and 11593: Transfer the contents of one Reagent B vial into a Reagent A bottle. Mix gently. Other volumes can be prepared in the proportion: 4 mL Reagent A + 1 mL Reagent B. Stable for 2 months at 2-8°C
- Cod. 11598: Transfer 25 mL of one Reagent B vial into a Reagent A bottle. Mix gently. Other volumes can be prepared in the proportion: 4 mL Reagent A + 1 mL Reagent B. Stable for 2 months at 2-8°C.

ADDITIONAL EQUIPMENT

- Analyzer, spectrophotometer or photometer with cell holder thermostatable at 37°C and able to read at 405 nm
- Cuvettes with 1 cm light path.

SAMPLES

Serum and plasma collected by standard procedures.

Alkaline phosphatase in serum or plasma is stable for 7 days at 2-8°C. Heparin may be used as anticoagulant.

PROCEDURE

- 1. Bring the Working Reagent and the instrument to reaction temperature
- 2. Pipette into a cuvette: (Note 1)

Working Reagent	1.0 mL
Sample	20 µL

- 3. Mix and insert the cuvette into the photometer.
- 4. Record initial absorbance and at 1 minute intervals thereafter for 3 minutes.
- Calculate the difference between consecutive absorbances, and the average absorbance difference per minute (ΔA/min).

CALCULATIONS

The ALP catalytic concentration in the sample is calculated using the following general formula:

$$\Delta A/\min x \frac{Vt \times 10^6}{\epsilon \times I \times Vs} = U/L$$

The molar absorbance (ε) of 4-nitrophenol at 405 nm is 18450, the lightpath (I) is 1 cm, the total reaction volume (Vt) is 1.02, the sample volume (Vs) is 0.02, and 1 U/L are 0.0166 µkat/L. The following formulas are deduced for the calculation of the catalytic concentration:

ΔA/min	x 2764 = U/L x 46.08 = μkat/L
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REFERENCE VALUES

Reaction temperature	men	women
37°C, up to ²	115 U/L = 1.92 μKat/L	105 U/L = 1.75 μKat/L

Concentrations in growing children are higher and highly variable. These ranges are given for orientation only; each laboratory should establish its own reference ranges.

It is recommended to use the Biochemistry Control Serum level I (cod. 18005, 18009 and 18042) and II (cod. 18007, 18010 and 18043) to verify the performance of the measurement

Each laboratory should establish its own internal Quality Control scheme and procedures for corrective action if controls do not recover within the acceptable tolerances.

METROLOGICAL CHARACTERISTICS

- Detection limit: 1.0 U/L = 0.017 μkat/L
- Linearity limit: 1200 U/L = 20 μkat/L. For higher values dilute sample 1/2 with distilled water and repeat measurement.
- Repeatibility (within run):

Mean Concentration	CV	n
61 U/L = 1.02 μkat/L	1.0 %	20
244 U/L = 4.07 μkat/L	0.7 %	20

Reproducibility (run to run):

Mean Concentration	CV	n
61 U/L = 1.02 μkat/L	3.4 %	25
244 U/L = 4.07 μkat/L	1.1 %	25

- Trueness: Results obtained with this reagent did not show systematic differences when compared with reference reagents. Details of the comparison experiments are available on
- Interferences: Lipemia (triglycerides < 10 g/L) and bilirubin (< 20 mg/dL) do not interfere. Hemoglobin (> 2.5 g/L) interfere. Other drugs and substances may interfere

These metrological characteristics have been obtained using an analyzer. Results may vary if a different instrument or a manual procedure are used.

DIAGNOSTIC CHARACTERISTICS

Alkaline phosphatase catalyzes the hydrolysis of organic phosphate monoesters at alkaline pH. The enzyme is present in practically all tissues of the body, especially at or in the cell membranes, and it occurs at particularly high concentrations in placenta, intestinal epithelium, kidney tubules, osteoblasts and liver.

The form present in the sera of normal adults originates mainly in the liver and bone.

Elevated serum ALP is found in patients with bone disease associated with increased osteoblastic activity (Paget's disease, primary and secondary hyperparathyroidism, bone tumors, rickets, osteomalacia, bone fractures) and also in patients with hepatobiliary disease (obstructive jaundice, hepatitis, hepatotoxicity caused by drugs, liver cancer). Physiological changes, such as bone growth and pregnancy, may cause increases in ALP levels4.5

Clinical diagnosis should not be made on the findings of a single test result, but should integrate both clinical and laboratory data

NOTES

1. These reagents may be used in several automatic analysers. Instructions for many of them are available on request.

BIBLIOGRAPHY

- 1. IFCC primary reference procedures for the measurement of catalytic activity concentrations of enzymes at 37 °C. Part 9. Reference procedure for the measurement of catalytic concentration of alkaline phosphatase. Clin Chem Lab Med 2011; 49:1439-1446.
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