# ALKALINE PHOSPHATASE

METHOD - PNPP-DEA METHOD PRODUCT CODE - DA01

# INSTRUCTIONS FOR USE



INTENDED USE: Test for estimation of Alkaline Phosphatase activity in serum / plasma using PNPP kinetic method.

#### SUMMARY AND PRINCIPLE

Serum Alkaline Phosphatase levels are of interest in the diagnosis of hepatobiliary disorders and bone disease associated with increased osteoblastic activity. Moderate elevations of Alkaline Phosphatase may be seen in Hodgkin's disease, congestive heart failure, ulcerative colitis, regional enteritis, and intra-abdominal bacterial infections. Alkaline Phosphatase is a reagent set for determination of Alkaline Phosphatase activity based on kinetic method using P-nitrophenylphosphate. Alkaline Phosphatase is a single reagent system with one step reconstitution.

PNPP

Alkaline Phosphatase

p-nitrophenol + phosphate

# KIT COMPONENTS

Reagent 1: Substrate Reagent

Reagent 2: Diluent

### REAGENT PREPARATION, STORAGE & STABILITY

Reconstitute each substrate tablet with diluent as per the instruction indicated on the substrate bottle. The working reagent is stable for 21 days at 2- 8  $^{\circ}$ C. The reagent kit should be stored at 2- 8  $^{\circ}$ C and is stable till the expiry date indicated on the label.

### PRECAUTIONS & HANDELING

The reagents/samples should be handled by qualified personnel only. Discard reagent/sample as per good laboratory practices and local regulatory requirements. Read the instructions given on the labels and instructions for use carefully before using the kit. The kit is intended for in-vitro diagnostic use only. Don't freeze the reagent. Do not shake the reagent vigorously. Discard the reagent if the absorbance of the reagent exceeds 0.700 O.D. against D/W at 405 nm. Contamination of the reagent should be avoided.

# **TEST PARAMETERS**

Name	ALK.PHOS.
Reaction Type	Kinetic (个)
Wavelength	405 nm
Flow Cell Temp.	37 °C
Blank setting	Reagent
Blank abs. limit	< 0.700
Linearity	2000 IU/L

Reagent Vol	1000 µl
Sample Vol	20 µl
Temperature	37 °C
Delay Time	60 sec.
Read Time	30 sec.
Factor	2720
Standard Conc.	1 <del>4</del> 27

# MATERIALS REQUIRED BUT NOT PROVIDED

Test tubes, Micropipette with tips, Analyzer, Controls, Incubation chamber.

### SPECIMEN COLLECTION & PRESERVATION

Blood should be collected in a clean dry container. Heamolysed specimen should be avoided as it may falsely elevate results. EDTA, Citrate and Oxalate inhibit Alkaline Phosphatase activity and should not be used as anti-coagulant. Alkaline Phosphatase in the serum/plasma is stable for 4 days when stored at 2- 8  $^{\rm OC}$  and several months when stored at -10° C.

#### COMPONENTS OF REAGENT

Component	Concentration
Diethanolamine Buffer, pH 9.8	1 Mol/l
p-nitrophenyl phosphate	10 mmol/l
Magnesium chloride	0.5 mmol/l
Stabilizers and inactive ingredients.	

# **ASSAY PROCEDURE**

	Blank	Test
Reagent	1000 μΙ	1000 µl
Serum / Plasma	-	20 µl
Mix the reagent and sample in t the stop watch.	the above-mentioned ra	tio and start
the stop water.		

Record absorbance at 60<sup>th</sup>, 90<sup>th</sup>, 120<sup>th</sup> & 150<sup>th</sup> sec. (30 sec. interval).

# CALCULATION

Calculate average Abs/min = Abs / 30 sec x 2 Alkaline Phosphatase (IU/L) = Abs / min x 2720

### REFERENCE VALUES FOR NORMAL PEOPLE

	25 °C	30 °C	37 °C
Adult (>15 years)	60-170 IU/L	70-207 IU/L	108-306 IU/L
Children (<15 years)	150-450 IU/L	195-585 IU/L	210-810 IU/L

# **ACTIVITY COMPARISON CHART**

Temperature of Assay	Corresponding Activity (IU/L) at		
	25 °C	30 °C	37°C
25 °C	1.00	1.30	1.80
30 °C	0.75	1.00	1.35

### PERFORMANCE CHARACTERISTICS

**Measuring Range:** The assay is linear between 27 - 2000 IU/L. If the Alkaline Phosphatase value exceeds linearity limit (above 2000 IU/L), dilute the specimen suitably with normal saline and repeat the assay. In that case, assay value should be multiplied with the dilution factor to obtain correct Alkaline Phosphatase value of the specimen.

**Interference:** There is no significant interference in samples containing Bilirubin upto 20 mg/dL and Haemoglobin upto 400 mg/dL.

**Precision:** Precision studies has been carried out using quality control sera as shown below:

(n=10)	Within Run		Between Run			
Specimen Material	Mean (IU/L)	SD (IU/L)	CV %	Mean (IU/L)	SD (IU/L)	CV %
Low Value Serum	105.4	0.52	0.5	110	0.92	0.8
High Value Serum	406.7	0.82	0.2	394.9	0.74	0.2

Note: We recommend all the laboratories to establish its own accuracy and precision data.



# **QUALITY CONTROL**

Inclusion of a normal value and abnormal value chemistry control serum in each test run ensures optimum quality control. Consistent use of same type and methodology of control serum provides between run precision and accuracy data for Alkaline Phosphatase. We recommend to produce such data on daily basis for greater accuracy in assay system which include reagents, instrument, apparatus and operator.

# **PRECAUTIONS**

- Discard the reagent if its absorbance exceeds 0.700 at 405 nm against distilled water.
- If Alkaline Phosphatase activity exceeds 2000 IU/L then dilute the specimen suitably with normal saline & repeat the assay. In such case the results obtained should be multiplied by dilution factor to obtain the correct Alkaline Phosphatase activity.

### **BIBLIOGRAPHY**

- Henry R.J., "Enzymes" in Clinical Chemistry Principle and Techniques, Harper & row Publishers, New York, 815 (1974).
- 2. Young D.S. et. al, Clin Chem.18,1041, (1972)

Symbol	Explanation	Symbol	Explanation
•	Manufactured By	IVD	In Vitro Diagnostic Use
LOT	Lot Number	[]i	Read Instructions Before Use
REF	Catalogue Number	1	Storage Temperature
۳	Manufacturing Date	$\sum$	Number of Tests / Volume
$\square$	Expiry Date	(2)	Do Not Reuse
涨	Protect from Sunlight	<b>†</b>	Keep Dry