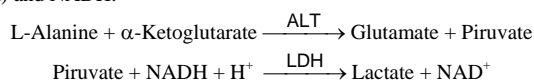


Store at +2 to +8°C

PRINCIPLE OF THE METHOD

Alanine aminotransferase (ALT) or Glutamate pyruvate transaminase (GPT) catalyses the reversible transfer of an amino group from alanine to α -ketoglutarate forming glutamate and pyruvate. The pyruvate produced is reduced to lactate by lactate dehydrogenase (LDH) and NADH:



The rate of decrease in concentration of NADH, measured photometrically, is proportional to the catalytic concentration of ALT present in the sample.

CLINICAL SIGNIFICANCE

The ALT is a cellular enzyme, found in highest concentration in liver and kidney.

High levels are observed in hepatic disease like hepatitis, diseases of muscles and traumatism, its better application is in the diagnosis of the diseases of the liver.

When they are used in conjunction with AST aid in the diagnosis of infarcts in the myocardium, since the value of the ALT stays within the normal limits in the presence of elevated levels of AST.

Clinical diagnosis should not be made on a single test result; it should integrate clinical and other laboratory data.

REAGENTS

- Reagent 1 (buffer)**
 TRIS pH 7.8 100 mmol/L
 Lactate dehydrogenase (LDH) 1200 U/L
 L-Alanine 500 mmol/L
- Reagent 2 (substrate)**
 NADH 0.18 mmol/L
 α -Ketoglutarate 15 mmol/L

PREPARATION

Working reagent (WR): Mix: 9 vol. (R1) Buffer + 1 vol. (R2) Substrate
 Stability: 21 days at +2 to +8°C or 72 hours at room temperature (+15 to +25°C).

STORAGE AND STABILITY

All the components of the kit are stable until the expiration date on the label when stored tightly closed at +2 to +8°C, protected from light and contaminations prevented during their use.

Do not use reagents over the expiration date.

Signs of reagent deterioration:

- Presence of particles and turbidity.
- Blank absorbance (A) at 340 nm \geq 1.00.

ADDITIONAL EQUIPMENT

- Spectrophotometer or colorimeter measuring at 340 nm.
- Thermostatic bath at 25°C, 30°C or 37°C (\pm 0.1°C)
- Matched cuvettes 1.0 cm light path.
- General laboratory equipment.

SAMPLES

Serum or plasma. Stability: 7 days at +2 to +8°C.

PROCEDURE

- Assay conditions:
 Wavelength: 340 nm
 Cuvette: 1 cm light path
 Temperature: 25°C / 30°C / 37°C
- Adjust the instrument to zero with distilled water or air.
- Pipette into a cuvette:

WR (mL)	1.0
Sample (μ L)	100
- Mix, incubate for 1 minute.
- Read initial absorbance (A) of the sample, start the stopwatch and read absorbance at 1 minute intervals thereafter for 3 minutes.
- Calculate the difference between absorbance and the average absorbance differences per minute (Δ A/min).

CALCULATIONS

Δ A/min x 1750 = U/L of ALT (T)

Units: One international unit (IU) is the amount of enzyme that transforms 1 μ mol of substrate per minute, in standard conditions. The concentration is expressed in units per litre of sample (U/L).

Temperature conversion factors

To correct results to other temperatures multiply by:

Assay temperature	Conversion factor to		
	25°C	30°C	37°C
25°C	1.00	1.32	1.82
30°C	0.76	1.00	1.39
37°C	0.55	0.72	1.00

QUALITY CONTROL

Control sera are recommended to monitor the performance of assay procedures with Contro N and Contro P.

If control values are found outside the defined range, check the instrument, reagents and technique for problems.

Each laboratory should establish its own Quality Control scheme and corrective actions if controls do not meet the acceptable tolerances.

REFERENCE VALUES

	25°C	30°C	37°C
Men	up to 22 U/L	29 U/L	40 U/L
Woman	up to 18 U/L	22 U/L	32 U/L

Normal newborns have been reported to show a reference range of up to double the adult, attributed to the neonate's hepatocytes. These values decline to adult levels by approximately 3 months of age.

These values are for orientation purpose; each laboratory should establish its own reference range.

PERFORMANCE CHARACTERISTICS

Measuring range: From detection limit of 3.9 U/L to linearity limit of 260 U/L.

If the results obtained were greater than linearity limit, dilute the sample 1/10 with NaCl 9 g/L and multiply the result by 10.

Precision:

	Intra-assay (n=20)		Inter-assay (n=20)	
	Mean (U/L)	SD	CV (%)	
Mean (U/L)	33.2	1.28	3.02	31.3
SD	1.00	1.47	3.02	1.29
CV (%)	3.02	1.14	3.02	1.57
				3.00
				1.22

Sensitivity: 1 U/L = 0.00052 A/min.

Accuracy: Results obtained using CHRONOLAB reagents did not show systematic differences when compared with other commercial reagents.

The results of the performance characteristics depend on the analyzer used.

INTERFERENCES

Anticoagulants currently in use like heparin, EDTA, oxalate and fluoride do not affect the results. Hemolysis interferes with the assay.

A list of drugs and other interfering substances with ALT determination has been reported by Young et. Al.

NOTES

CHRONOLAB has instruction sheets for several automatic analyzers. Instructions for many of them are available on request.

REFERENCES

- Abbott L. et al. Acid phosphatase. Kaplan A et al. Clin Chem The C.V. Mosby Co. St Louis. Toronto. Princeton 1984; 1079-1083.
- Young DS. Effects of drugs on Clinical Lab. Tests, 4th ed AACC Press, 1995.
- Young DS. Effects of disease on Clinical Lab. Tests, 4th ed AACC 2001.
- Burtis A et al. Tietz Textbook of Clinical Chemistry, 3rd ed AACC 1999.
- Tietz N W et al. Clinical Guide to Laboratory Tests, 3rd ed AACC 1995.

PACKAGING

Ref: 101-0437	Cont.: 4x250/4x30 ml
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