

GAMMA GT SL

Kinetic Method (Szasz-Tris)
Liquid reagents ready to use

REF. 0011 5x 10 ml
REF. 4196 2x 50 ml
REF. 4197 2x100 ml



INTENDED USE

Quantitative determination of glutamyltransferase (γ -GT) in serum.

PRINCIPLE

The γ -GT, in the presence of glycyl-glycine, splits the L- γ -glutamyl-3-carboxy-4-nitroanilide (carboxi-glupa) in L- γ -glutamyl-glycyl-glycine and 5-amino-2-nitrobenzoate. The absorbance change in time unit measured at 405 nm is proportional to the enzyme activity in the sample.

SAMPLE

Serum, plasma EDTA. Avoid hemolysis.

The γ -GT in serum is stable one week at 2-25°C. Store at -20°C for prolonged periods.

KIT COMPONENTS

Reagent (A) γ -GT Volume = 40/80 ml	Tris buffer Glycyl-glycine	10 mmol/l 125 mmol/l
Reagent (B) γ -GT Volume = 20 ml	Buffer L- γ -glutamyl-3-carboxy-4-nitroanilide	100 mmol/l 6 mmol/l

The reagents are stable until the expiration date indicated on the label if stored at 2-8°C and protected from light. Do not freeze. Once opened reagents are stable for 2 months at 2-8°C if contamination is avoided.

Keep bottles closed when not in use.

REAGENT PREPARATION

Liquid Reagents, bring to room temperature (15-25°C) before use.

For use as **monoreagent**: add a part of Reagent (B) to 4 parts of Reagent (A).

The working solution (A+B) is stable 5 days at 15-25°C and 3 weeks at 2-8°C.

PRECAUTIONS AND WARNINGS

Reagents may contain some non-reactive and preservative components. It is suggested to handle carefully it, avoiding contact with skin and swallow.

Use the normal precautions required in the laboratory.

Dispose of waste according to local laws.

PROCEDURE

Wavelength:	405 nm
Lightpath:	1 cm
Temperature:	37°C
Reading:	against distilled water
Method:	Increasing kinetic

Use as monoreagent:

pipette:

Working solution (A+B)	1000 μ l
sample	100 μ l

Mix, incubate at 37°C for 1 minute, read the initial absorbance against water. Make 3 readings at a distance of 60 seconds. Calculate the average value of the absorbance variations per minute. ($\Delta A/min$).

Use as bireagent:

pipette:

Reagente (A)	1000 μ l
sample	100 μ l

mix and after 1 minute add:

Reagent (B)	250 μ l
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Mix, incubate at 37°C for 1 minute, read the initial absorbance against water. Make 3 readings at a distance of 60 seconds. Calculate the average value of the absorbance variations per minute. ($\Delta A/min$).

This method describes the manual procedure to use the kit.

For automated procedure, ask for specific applications.

RESULTS CALCULATION

Perform calculation in Units per litre, multiplying the $\Delta A/min$ by the factor as it is indicated:

Monoreagent: activity in U/L: $\Delta A/min \times 1158$

Bireagent: activity in U/L: $\Delta A/min \times 1421$

EXPECTED VALUES

Men: ≤ 50 U/L

Women: ≤ 35 U/L

Each laboratory should establish appropriate reference intervals related to its population.

QUALITY CONTROL

You must perform the controls at each kit's use and verify that the values obtained are within the reference range reported in the operating instructions. For this purpose we recommend the use of control sera: PRECISENORM (REF.6000) and PRECISEPATH (REF.6001).

PERFORMANCE

Sensitivity: the sensitivity of the method is: 2 U/L

Linearity: the method is linear up to 450 U/L. For higher values, dilute the sample 1:10 and multiply the result by 10.

Precision intra-assay:

	Level 1	Level 2	Level 3
Mean (U/l)	2140	147.1	307.0
DS	0.819	1.792	1.414
CV %	3.83	1.22	0.46

Precision inter-assay:

	Level 1	Level 2	Level 3
Mean (U/l)	21.21	152.5	322.5
DS	0.619	1.958	1.958
CV %	2.92	1.28	0.61

Interferences: bilirubin does not interfere up to 30 mg/dl. Ascorbic acid does not interfere up to 50 mg/dl. The presence of hemolysis causes an underestimation proportional to the degree of hemolysis.

The commonly used anticoagulants inhibit the activity of γ -GT.

Anticonvulsants (phenytoin, barbiturates) cause a false increase in γ -GT.

Correlation against a reference method: $Y = 1.0399x + 2.1657$ $r = 0.9994$

REFERENCES

1. Szasz G., Clin. Chem. 22,2051 (1976).
2. Vassault, A. et al. Ann. Biol. Clin., 44,686 (1986).
3. Young, D.S., et al., Clin. Chem. 21:1D (1975).