LI@UIZYME

GAMMA GT

(SASZ Method)

Code	Product Name	Pack Size
LS020A	Liquizyme Gamma GT	25 ml

Intended Use

Diagnostic reagent for quantitative *in vitro* determination of GGT in human serum and plasma.

Clinical Significance

Although GGT is present in a variety of tissues, the serum enzyme appears to be primarily from the hepato-biliary system. Consequently, GGT is elevated in all forms of liver disease or damage. It is clinically useful in detecting obstructive jaundice, cholangitis and cholecystitis. Elevated levels are also observed with drug use (alcohol, sedatives, anticonvulsants and tranquilizers).

Principle

GGT present in the sample catalyzes the transfer of the glutamyl group from the substrate γ -glutamyl-3-carboxy-4-nitroanilide to glycylglycine forming glutamyl glycylglycine and 5-amino-2-nitrobenzoate.

L-γ-glutamyl-3- GGT L-γcarboxy-4-nitronilide → glutamylglycylglycine +
+ glcylglycine 5-amino 2nitrobenzoate

The rate of formation of 5-amino-2-nitrobenzoate is proportional to the activity of GGT present in the sample and can be measured kinetically at $400-420\,\mathrm{nm}$.

Reagent Composition Reagent 1: Buffer Reagent

Tris buffer (ph 8.25) : >125 mmol/LGlycyl Glycine : >125 mmol/L

Reagent 2: Substrate Reagent

L-γ-Glutamyl-3-carboxy-4-nitronilide <20 mmol/L

Reagent Preparation

Reagents are liquid, ready to use.

Stability And Storage

The unopened reagents are stable till the expiry date stated on the bottle and kit label when stored at $2-8^{\circ}$ C.

Materials Required But Not Provided

- Clean & Dry container.
- Laboratory Glass Pippetes or Micropioettes & Tips
- Colorimeter or Bio-Chemistry Analyzer.

Working Reagent Preparation

Mix 4 portion of reagent R1 with 1 portion of reagent R2.

Stability:

 $14 \, days$: at $2 - 8^{\circ}C$



Specimen Collection And Handling

Use serum, plasma (EDTA).

It is recommended to follow NCCLS procedures (or similar standardized conditions).

Stability In Serum / Plasma:

Unit Conversion

 $U/I \times 0.017 = \mu mol/I$

Expected Values

At 37°C

Male : 10 - 45 U/L Female : 5 - 32 U/L

It is recommended that each laboratory verify this range or derives reference interval for the population it serves.

Performance Data

Data contained within this section is representative of performance on Beacon system.

Data obtained in your laboratory may differ from these values.

Intra-assay precision	Mean	SD	CV
Within run (n=20)	(U/L)	(U/L)	(%)
Sample 1	55.59	1.49	2.68
Sample 2	176.45	2.74	1.55
Inter-assay precision	Mean	SD	CV
Run to run (n=20)	(U/L)	(U/L)	(%)
Sample 1	87.68	2.93	3.35

Comparisor

A comparison between Beacon GGT (y) and a commercially available test (x) using 20 samples gave following results:

y = 1.036 x - 1.979 U/L

r = 0.997

Interferences

Following substances do not interfere:

haemoglobin up to 5 g/l, bilirubin up to 40 mg/dl, triglycerides up to $2000\,\text{mg/dl}$.

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Warning And Precautions

For in vitro diagnostic use. To be handled by entitled and professionally educated person.

Waste Management

Please refer to local legal requirements.

Assay Procedure

: 405 nm Wavelength Cuvette : 1 cm

Addition Sequence	Volume	
Working Reagent	1000 μΙ	
Sample	50 μΙ	

Mix and read the initial absorbance after 60 sec at 37°C and repeat the reading after every 1,2 and 3 minutes. Calculate the mean absorbance change per minute. ($\Delta A/min$).

Calculation

Using factor:

GGT activity (U/L) ΔA/min x 2210

Applications for automatic analysers are available on request.

Assay Parameters For Photometers

Kinetic
405
50
1000
60
60
3
2210
37
Increasing
10
45
1.68
500
Water
U/L

References

- 1. Szasz G., Weimann G. Suhler F., Wahlefrld A.W., Presijn J. P. : Z Klin. Chem. Klin. Biochem. 12, 228 (1994).
- 2. Persijn & van der Slik W.: J. Clin. Chem. Clin. Biochem. 14, 421-427 (1976).
- 3.Tietz Textbook of Clinical Chemistry and Molecular Diagnostics. Burtis, C.A., Ashwood, E.R., Bruns, D.E.; 5th $edition, WB\,Saunders\,Comp., 2012.$

Symbols Used On Labels

REF

Catalogue Number



Manufacturer

Lot Number

 \mathbf{i}

See Instruction LOT for Use



Storage Temperature



CONT

Expiry Date

Content



In Vitro Diagnostics





BEA/24/GGT/LS/IFU-01 08/01/2022