

# GAMMA-GT kinetic-szasz/persijn method



#### PRODUCT CODE CZ007

#### INTENDED USE

This reagent is intended for *in vitro* quantitative determination of Gamma-GT in serum or plasma.

## CLINICAL SIGNIFICANCE

Gamma-glutamyl transferase ( $\gamma$ -GT) is a cellular enzyme with wide tissue distribution in the body, primarily in the kidney, pancreas, liver and prostate. Measurements of gamma-glutamyl transferase ( $\gamma$ -GT) activity are used in the diagnosis and treatment of hepatobiliary diseases such biliary obstruction, cirrhosis or liver tumors. Clinical diagnosis should not be made on a single test result; it should integrate clinical and other laboratory data.

#### PRINCIPLE

Gamma-GT catalyzes the transfer of glutamic acid to acceptors like glycylglycine in this case. This process releases 5-amino-2nitrobenzoate, which can be measured at 405 nm. The increase in absorbance at this wavelength is directly related to the activity of gamma-GT.

γ-GT

L-Gamma-glutamyl-3-carboxy-4-nitranilide+Glycylglycine \_\_\_\_\_\_ L-Gamma-Glutamyl-Glycylglycine + 5-Amino-2-nitrobenzoate

## REAGENT COMPOSITION

REAGENT 1 (BUFFER)	
Tris pH 8.25	100 mmol/L
Glycylglycine	100 mmol/L
REAGENT 2 (SUBSTRATE)	
L-γ-glutamyl-3-carboxy-4-nitroanilide (Glupa-C)	4mmol/L

#### REAGENT PREPARATION SUBSTRATE START

R1 and R2 are ready-to-use and stable upto the expiry date if

contamination is avoided and stored at 2-8°C and protect from light.

#### SAMPLE START

Mix 4 parts of R1 + 1 Part of R2 = Mono reagent Stability of mono reagent: 4 Weeks at 2-8°C, 5 days at 15-25°C, Protect from light.

#### SPECIMEN

Serum, EDTA plasma, avoid hemolysis.

## PRECAUTION

- 1- The reagents contain sodium azide as preservative. Do not swallow and avoid contact with skin and mucous membranes.
- 2- Avoid contamination, use clean laboratory wares. Avoid direct exposure of working reagent to light.



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:	405 nm
:	1 cm light path
•	25°C/ 30°C/37°C

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# PROCEDURE

SUBSTRATE START	
Reagent 1 Buffer	1000 µL
Sample	100 µL
Mix, incubates for approx 1 min, then add,	
Reagent 2 Substrates	250 μL
SAMPLE START	
Mono reagent (R1+R2)	1000 µL
Sample	100 µL

## **READING FOR BOTH**

Mix and read absorbance after 1 min and start stop watch. Read absorbance again after 1, 2 and 3 min.

## CALCULATION

SUBSTRATE START	
405 nm	ΔA / min X 1421
SAMPLE START	
405 nm	ΔA / min X 1158

## LINEARITY

up to 700 U/L, the sample should be diluted 1 + 5 with 0.9 % NaCl solution, if  $\Delta A/min$  exceeds 0.200, multiply the result by 6.

#### NORMAL RANGE

	25°C	30°C	37°C
Women	4-18 U/L	5-23 U/L	7-32 U/L
Men	6-28 U/L	8-38 U/L	11-50 U/L
Each laboratory should	d establish reference	ranges for its o	wn patients'

population.

## QUALITY CONTROL

All control sera with values determined by this method can be used.

## SYMBOL ON LABELS

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Symbols	Signify	Symbols	Signify
REF	Catalogue Number	SIZE	Pack Size
$\Box$	Expiry Date	VOL	Volume
ł	Storage Condition	LOT	Lot Number
Ĩ	Instruction for Use	IVD	In Vitro Diagnostics
$\sim\sim$	Manufacturing Date		Manufacturer
$\overline{\Sigma}$	Number of Tests	2	For Single Use Only
EC REP	EC Representative	(6	European conformity

## BIBILOGRAPHY

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- 3- Szasz G. Gamma-Glutamyltranspeptidase. In: Bergmeyer Hu. Methoden Der Enzymatischen Analyse. Weinheim: Verlag Chemie, 1974. P. 757.



MDSS GmbH Schiffgraben 41 30175 Hannover, Germany Doc.No.: IFU-CH-007 Rev.: 05 Page **1** of **1**