ORDER INFORMATION

CODE: DL2701 - R1 - 1 x 10 ML + R2 - 1 x 2 ML DL2702 - R1 - 2 x 10 ML + R2 - 2 x 2 ML DELTA LIPASE

(Methyl Resorufin)

INTENDED USE:

This reagent kit is intended for **"in vitro"** quantitative determination of LIPASE activity in serum & plasma.

CLINICAL SIGNIFICANCE:

Lipases are glycoproteins with a molecular weight of 47000 Daltons. Lipase hydrolyzes the ester linkages. Specifically, lipase catalyzes the partial hydrolysis of dietary triglycerides in the intestine to the 2-monoglyceride intermediate, with the production of long chain fatty acids.

PRINCIPLE:

The chromogenic lipase substrate 1,2-O-dilauryl-rac-glycero-3-glutaric acid-(6-methylresorufin)-ester is cleaved by the catalytic action of alkaline lipase solution to form 1,2-O-dilauryl-rac-glycerol and an unstable intermediate glutaric acid-(6-methylresorufin)-ester. This decomposes spontaneously in alkaline solution to form glutaric acid and methylresorufin. The color intensity of the red dye formed is directly proportional to lipase activity and can be determined photometrically.

REAGENT COMPOSITION:

Reagent 1: Buffer Reagent Reagent 2: Substrate Reagent

Lipase Calibrator Concentration: Printed on the Vial.

SAMPLES:

Li-, Na- vagy NH₄-heparin plasma, serum.

EDTA-, oxalate-, fluoride or citrated plasma lead to decreased resultits.

WORKING REAGENT PREPARATION & STABILITY:

Reagent 1 & Reagent 2 are Ready to use and are stable up to the expiry date stated on the label.

LIPASE Calibrator: Dissolve with distilled water (Qty Printed on the vial). Cap and mix gently to dissolve contents. Reconstituted calibrator is stable for 7 days at 2 to 8°C or 3 months at –20°C; aliquots into small volumes and freeze.

GENERAL SYSTEM PARAMETERS:

Reaction type Fixed Time
Wave length 580 nm
Light Path 1 Cm
Reaction Temperature 37°C

Blank / Zero Setting With Distilled Water

Low Normal at 37°C 13 U/I High Normal at 37°C 60 U/I Linearity 250 U/I

ASSAY PROCEDURE:

	Calibrator	Sample
Reagent 1	1000 μl	1000 μl
Calibrator	10 μl	
Sample		10 μΙ

Mix and incubate for 60 second and then add

Reagent 2	200 μl	200 μl
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Mix and after 60 second incubation, measure the change in absorbance for 60 seconds at 37° C.

Determine the Absorbance.

CALCULATION:

Lipase Activity (U/I) = $\frac{\Delta \text{ Abs. of Sample}}{\Delta \text{ Abs. of Calibrator}} X \text{ Conc. of Calibrator}$

LINEARITY:

Reagent is Linear up to 250 U/I.

Dilute the sample appropriately and re-assay if Lipase Activity exceeds 250 U/I.

REFERENCE NORMAL VALUE:

Serum Lipase activity 13-60 U/I

It is recommended that each laboratory should assign its own normal range.

QUALITY CONTROL:

For accuracy it is necessary to run known controls with every assay.

LIMITATION & PRECAUTIONS:

- 1. Storage conditions as mentioned on the kit to be adhered.
- 2. Do not freeze or expose the reagents to higher temperature as it may affect the performance of the kit.
- 3. Before the assay bring all the reagents to room temperature.
- 4. Avoid contamination of the reagent during assay process.
- 5. Use clean glassware free from dust or debris.

BIBLIOGRAPHY:

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