

# CORMAY LIPOPROTEIN (a)



## DIAGNOSTIC KIT FOR DETERMINATION OF LIPOPROTEIN (a) CONCENTRATION

<b>Kit name</b>	<b>Kit size</b>	<b>Cat. No</b>
CORMAY LIPOPROTEIN (a)	1 x 75 ml	6-302

### INTRODUCTION

Lipoprotein(a) is a complex, cholesterol-carrying particle in the blood related to LDL which, when present at high levels, may be associated with the development of atherosclerosis and coronary heart disease, independent of LDL cholesterol and apoB. The structural component of Lp(a) distinguishing it from LDL is apolipoprotein(a), a large protein attached by disulfide bonding to the apoB-100 component of LDL. The similarity of the apo(a) sequence to those of plasminogen and hepatocyte growth factor suggests that the role of lipoprotein(a) in promoting the development of atherosclerosis may come from its capability to:

- 1) interfere in the breakdown of blood clots,
- 2) stimulate atherosclerotic cell proliferation.

Lp(a) levels is largely due to hereditary factors and considered to be useful in assessment of atherosclerosis risks.

### METHOD PRINCIPLE

When an antigen-antibody reaction occurs between Lp(a) in a sample and anti-Lp(a) antibody which has been sensitized to latex particles, agglutination results. This agglutination is detected as an absorbance change (700 nm), with the magnitude of the change being proportional to the quantity of Lp(a) in the sample. The actual concentration is then determined by interpolation from a calibration curve prepared from calibrators of known concentration.

### REAGENTS

<b>Package</b>	
1-Reagent	1 x 50 ml
2-Reagent	1 x 25 ml

### Reagent preparation and stability

The reagents are ready to use.

The reagents are stable up to the kit expiry date printed on the package when stored at 2-10°C. Protect from light!

### Concentrations in the test

suspension of latex particles sensitized with anti-Lp(a) antibodies (rabbit) (pH 7.3) 0.4 w/v%  
glycine buffer solution (pH 9.0)

### Warnings and notes

- Product for in vitro diagnostic use only.
- Reagent bottles should be shaken before use by gently inverting several times.
- After measurements are taken, reagent bottles should be capped and kept at 2-10°C. Care should be taken not to interchange the caps of reagent bottles.
- Reagents with different lot numbers should not be interchanged or mixed.
- The reagents contain sodium azide (< 0.1%) as a preservative. Avoid contact with skin and mucous membranes.

### ADDITIONAL EQUIPMENT

- automated clinical chemistry analyser capable of accommodating two-reagent assays;
- general laboratory equipment;

### SPECIMEN

Serum or plasma (Na-EDTA, K-EDTA, Na-Heparin, Li-Heparin).

It is recommended to perform the assay with freshly collected samples. If the test cannot be done immediately, the sample should be placed in a tightly sealable container and stored at -20°C. Repeated freezing and thawing should be avoided.

Samples which contain an excessive amount of Lp(a) should be diluted with physiological saline and re-tested.

### PROCEDURE

wavelength	700 nm
temperature	37°C

These reagents may be used in automatic analysers according to their service manual. These reagents may be used directly in HITACHI 911/912 analysers. Applications for analysers are available on request.

### REFERENCE VALUES

serum, plasma	< 30 mg/dl
---------------	------------

It is recommended for each laboratory to establish its own reference ranges for local population. Diagnosis should only be made after taking clinical symptoms and the results of other tests into consideration.

### QUALITY CONTROL

For internal quality control it is recommended to use the CORMAY Lp(a) CONTROL N (Cat. No 4-492) and CORMAY Lp(a) CONTROL P (Cat. No 4-493) with each batch of samples.

For the calibration of automatic analysers systems the CORMAY Lp(a) CALIBRATORS kit (Cat. No 4-281) is recommended. A calibration curve should be drawn, each time the test performed. The standard solutions should be measured at least twice.

### PERFORMANCE CHARACTERISTICS

These metrological characteristics have been obtained using an automatic analyser TBA-30R. Results may vary if a different instrument is used.

- **Analytical range:** 0.5 – 80 mg/dl.
- **Antigen excess:** up to 660 mg/dl.
- **Specificity / Interferences**  
Haemoglobin up to 500 mg/dl, bilirubin up to 30 mg/dl, triglycerides up to 6000 mg/dl, intralipid up to 5 g/l, RF up to 500 U/ml do not interfere with the test.

- **Precision**

Repeatability (within run) n = 10	Mean [mg/dl]	SD [mg/dl]	CV [%]
level 1	20.9	0.01	0.33
level 2	43.5	0.03	0.74

- **Method comparison**

A comparison between CORMAY reagent (y) and commercially available assay (x) using 25 samples gave following results:

$$y = 1.00x + 0.10 \text{ mg/dl};$$

$$R = 0.98 \quad (R - \text{correlation coefficient})$$

### WASTE MANAGEMENT

Please refer to local legal requirements.

## LITERATURE

1. Utermann G. et al.: Lp(a) Glycoprotein Phenotypes. Inheritance and relation to Lp(a)-lipoprotein concentrations in plasma., J. Clin. Invest., 80, 458 (1987).
2. McLaren J. W. et al.; cDNA sequence of human apolipoprotein(a) is homologous to plasminogen., Nature, 300, 132 (1987).
3. Neumeister B., Besenthal I., Liebich H.: Diagnostyka laboratoryjna., Urban & Partner, 126-127, (2001).

**Date of issue:** 10. 2004.

## MANUFACTURER

### P.Z. CORMAY

ul. Rapackiego 19, 20-150 Lublin, POLAND

P.O. Box 122, 20-954 Lublin 2

tel.: +48 (0) 81 749 44 00

fax: +48 (0) 81 749 44 34

<http://www.pzcormay.pl>

10/04/10/04