CORMAY ALPHA 1-MICROGLOBULIN 500

DIAGNOSTIC KIT FOR DETERMINATION OF α 1-MICROGLOBULIN CONCENTRATION

Kit name	Kit size	Cat. No
CORMAY ALPHA 1-MICROGLOBULIN 500	1 x 1000 ml	6-327

INTRODUCTION

α 1-microglobulin (αMi) is a low molecular weight glycoprotein (24-33 kD) which was initially isolated from the urine of patients with renal tubular disorders in 1975. It is mainly synthesized in the liver and is widely distributed in various body fluids.
The measurement of αMi in serum and urine has been considered to be useful for the diagnosis of functional renal disorders, the assessment of the progress and prognosis of diseases.

METHOD PRINCIPLE

When an antigen-antibody reaction occurs between αMi in a sample and anti-αMi antibody which has been sensitized to latex particles, agglutination results. This agglutination is detected as an absorbance change (572 nm), with the magnitude of the change being proportional to the quantity of αMi in the sample. The actual concentration is then determined by interpolation from a calibration curve prepared from calibrators of know concentration.

REAGENTS

Package

1-Reagent	1 x 500 ml
2-Reagent	1 x 500 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. On board stability of the reagents depends on type of analyser used for analysis. Protect from light and avoid contamination!

Concentrations in the test

suspension of latex particles sensitized with (rabbit) anti-αMi antibodies glycine buffer solution	0.25 w/v%

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 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, plasma or urine.
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.
Nevertheless, it is recommended to perform the assay with freshly collected samples.

PROCEDURE

wavelength	572 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 [3]

serum, plasma	10.0 – 30.0 mg/l
urine	1.0 – 5.0 mg/l

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 control serum for determination of αMi with each batch of samples, eg.: ROCHE or BIORAD.
For the calibration of automatic analysers systems the CORMAY ALPHA 1-MGLOB CALIBRATORS (S) (Cat. No 4-286) for serum samples and the CORMAY ALPHA 1-MGLOB CALIBRATORS (U) (Cat. No 4-285) for urine samples is recommended.
Calibration stability depends on type of analyser used for analysis. The calibration curve should be prepared with change of reagent lot number or as required e.g. quality control findings outside the specified range.

PERFORMANCE CHARACTERISTICS

These metrological characteristics have been obtained using an automatic analyser HITACHI 917. Results may vary if a different instrument is used.

- **Analytical range:** 1.5 – 200 mg/l (serum)
  0.3 – 50.0 mg/l (urine).
  For higher concentration dilute the sample with 0.9% NaCl and repeat the assay. Multiply the result by dilution factor.

- **Specificity / Interferences**
  Haemoglobin up to 0.5 g/dl, NH_4Cl do 400 mg/dl, bilirubin up to 31 mg/dl do not interfere with the test in urine.

- **Precision**

Repeatability (run to run) n = 20	Mean [mg/l]	SD [mg/l]	CV [%]
level 1	0.5	0.0	3.97
level 2	1.6	0.0	1.81
level 3	13.9	0.1	0.42

- **Method comparison**

A comparison between CORMAY reagent (y) and commercially available assay (x) using 55 serum samples gave following results:
y = 1.00x + 2.83 mg/l;
R = 1.00 (R – correlation coefficient)

A comparison between CORMAY reagent (y) and commercially available assay (x) using 55 urine samples gave following results:
y = 1.00 x - 0.52 mg/l;
R = 1.00 (R – correlation coefficient)

WASTE MANAGEMENT
Please refer to local legal requirements.

LITERATURE

1. Galvin J. P. et al.: Particle enhanced photometric immunoassay systems., Clin. Lab. Assays (Pap. Annu. Clin. Lab. Assays Conf.), 4th, 73 (1983).
2. Singer J. M. et al.: The latex fixation test. I. Application to the serologic diagnosis of rheumatoid arthritis, Amer. J. Med., 21, 888 (1956).
3. Yoshihisa Ito: a1-microglobulin (protein HC), Nippon Rinsho, 47, 176 (1989).

Date of issue: 01. 2013.

MANUFACTURER

PZ CORMAY S.A.
22 Wiosenna Street,
05-092 Łomianki, POLAND
tel.: +48 (0) 22 751 79 10
fax: +48 (0) 22 751 79 14
http://www.cormay.pl

01/13/01/13