CORMAY ALPHA 1-MICROGLOBULIN

DIAGNOSTIC KIT FOR DETERMINATION OF α 1-MICROGLOBULIN CONCENTRATION

Kit name	Kit size	Cat. No
CORMAY ALPHA 1-MICROGLOBULIN	1 x 100 ml	6-307

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 50 ml
2-Reagent	1 x 50 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 (rabbit) anti-αMi antibodies	0.25 w/v%
glycine buffer solution	

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.
Samples which contain an excessive amount of αMi should be diluted with physiological saline and re-tested.

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

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. 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 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).

- **Specificity / Interferences**
  Haemoglobin up to 500 mg/dl, NH_4Cl do 400 mg/dl, bilirubin up to 31 mg/dl do not interfere with the test in urine.

- **Precision**

Repeatability (within 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)

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).

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