# NEO CHEM URIC ACID

(Uricase / PAP method)

KIT NAME	KIT SIZE
NEO CHEM - URIC ACID	2 x 25 ml

# INTRODUCTION

Uric acid is a product of purine catabolism. It is produced in the liver and excreted in the urine. Both, the amount of uric acid production and the efficiency of renal excretion, affect serum uric level. Elevated serum uric acid level is caused usually by gout, leukemia, diabetes mellitus, hyperfunction of parathyroid and thyroid, renal failure, renal calculosis. Urate concentration in serum depends on glomerular filtration, thus is useful for renal function monitoring.

# **METHOD PRINCIPLE**

nzymatic, colorimetric method with uricase and peroxidase. uric acid + 2  $H_2O + O_2$  <u>uricase</u> > allantoine +  $CO_2 + H_2O_2$ 

ADPS + 4-aminoantipyrine + 2  $H_2O_2$  POD> quinoneimine dye +4  $H_2O_2$  (coloured compound)

The colour intensity is proportional to the uric acid concentration.

#### KIT CONTENTS

Reagent Name	Pack Size
R1 - Uric Acid Reagent	2 X 25 ml
R2 - Standard	2 ml

Refer standard Concentration on the vial.

# WORKING REAGENT PREPARATION AND STABILITY

The reagents are to be stored at 2-8  $^{\circ}\text{C}$  . Do not freeze the reagents.

# CONCENTRATIONS IN THE TEST

 Buffer PIPES (pH7.8)
 > 150 mmol/l

 Chromogen
 1.0 mmol/l

 Ascorbate oxidase
 > 100 mmol/l

 Peroxidase (P0D)
 > 100 mmol/l

 Uricase
 > 100 mmol/l

Activators & stabilizers.

# ADDITIONAL EQUIPMENT

Automatic analyzer or photometer able to read at 505 nm (Hg 530-550 nm), Thermostat at 25°C or 37°C, General laboratory equipment.

#### **SPECIMEN**

Serum, heparinized plasma free from hemolysis. at -20°C.

Do not use EDTA and fluoride as anticoagulants

Specimen can be stored 3-5 days at 2-8°C or 6 months

Nevertheless it is recommended to perform the assay with freshly collected samples.

# **PROCEDURE**

These reagents may be used both for manual assay (Sample Start and Reagent Start method) and in several automatic analyzers. Programme Sheets are available on request.

Wavelength 505 nm
Temperature 25°C / 37°C
Cuvette 1 cm

# Pipette into the cuvettes:

F				
Reagent	Blank (B)	Standard (S)	Test (T)	
R1 Uric Acid Reagent	1000 μl	1000 μl	1000 μl	
Bring up the temperature of determination. Then add,				
Distilled water	20 μl			
R2 - Standard		20 μ1		
Sample			20 μl	



Mix well, incubate for 15 min. at 20-25°C or 10 min. at 37°C. Read the absorbance of test A(T) and standard A(S) against blank

#### CALCULATION

Uric acid concentration = A(T)/A(S)x standard concentration

#### REFERENCE VALUES

Female	2.5 - 6.8 mg/dl
Male	3.6 - 7.7 mg/dl

It is recommended for each laboratory to establish its own reference ranges for local population.

# **OUALITY CONTROL**

To Ensure adequate quality control, each run should include assayed normal and abnormal controls. If commercial controls are not available it is recommended that known value samples be aliquoted, frozen and used as controls.

# PERFORMANCE CHARACTERISTICS

Sensitivity / Limit of Quantitation: 0.2 mg/dl (11.9 µmol/l)

Linearity: up to 25 mg/dl

Specificity / Interferences

Haemoglobin up to 7.5 g/dl, ascorbate up to 62 mg/l, bilirubin up to 20 mg/dl and triglycerides up to 500 mg/dl do not interfere with the test.

# LITERATURE

- 1. Thefeld C. et al.: Dtsch. Med. Wschr. 98, 380-384 (1973).
- 2. Barham D., Trinder P.: Analyst 97, 142-145 (1972).
- Fossati P., Prencipe L., Berti G.: Clin. Chem. 26/2, 227-231 (1980).
- Henry R.J.: Clinical Chemistry, Harper & Row Publishers Inc., New York (1974).
- Kaplan L.A., Pesce A.J., ed. Chemistry Theory, Analysis, and Correlation, 3rd ed. St Louis, MO: Mosby, 501-2 (1996).
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# SYSTEM PARAMETER

Method	End Point
Wavelength	505 nm
Zero Setting	Reagent Blank
Temperature Setting	37° C
Incubation Temperature	37° C
Incubation Time	10 mins.
Delay Time	
Read Time	
No. of Reading	
Interval Time	
Sample Volume	0.020 ml (20 ul)
Reagent Volume	1.0 ml (1000 ul)
Standard Concentration	Refer Standard vial
Units	mg/dl
Factor	
Reaction Slope	Increasing
Linearity	25 mg/dl