

## URIC ACID

Enzymatic, Colorimetric,

**Diagnostic reagent for determination of Uric acid concentration.**

Liquid, 2 Reagents  
For in Vitro Diagnostic Use

Store at 2°C - 8°C  
Do not freeze

| Cat. No. |           |         |
|----------|-----------|---------|
| A2340    | URIC ACID | 5X100ML |
| A2341    | URIC ACID | 5X50ML  |
| A2342    | URIC ACID | 5X25ML  |

### TEST PRINCIPLE

Uric acid in sample is oxidized to allantoin in presence of the enzyme uricase and H<sub>2</sub>O<sub>2</sub> is generated. The H<sub>2</sub>O<sub>2</sub> reacts with TOOS and 4-aminoantipyrine in the presence of peroxidase to form a violet dye. The intensity of color formed is proportional to the uric acid concentration and can be measured photometrically between 510 and 560 nm.

### REAGENTS COMPOSITION

Composition in the test: phosphate buffer pH 7.0 50 mM, TOOS 0.34 mM, 4-aminoantipyrine 0.3 mM, uricase 450 U/l, POD > 2500 U/l, surfactant.

### TEST PARAMETERS

|              |  |
|--------------|--|
| Method:      | Colorimetric, Endpoint, Trinder, Increasing Reaction, Enzymatic                    |
| Wavelength:  | Hg 546 nm (510 - 560nm)  |
| Temperature: | 37°C   |
| Sample:      | Serum, Plasma, Urine (dilute urine 1:10 with physiological saline) Don't use EDTA. |
| Linearity:   | up to 25 mg/dl   |

### REAGENTS PREPARATION

If reagents are mixed in reduced quantities, mix 9 parts of reagent 1 with 1 part of reagent 2. Stability of working reagent: ≥ 90 days at 2-8°C, away from light sources. Stability of unmixed reagents: up to expiration date on labels at 2-8°C; Stability since first opening of vials of unmixed reagents: ≥ 60 days at 2-8°C.

### PRECAUTIONS

Reagent may contain some non-reactive and preservative components. It is suggested to handle carefully it, avoiding contact with skin and swallow. Perform the test according to the general "Good Laboratory Practice" (GLP) guidelines.

### SPECIMEN

Serum, plasma heparinate. Oxalate, citrate and fluoride could yield a small decrease of uric acid. Urine. Uric acid is stable 5 days at 4-25°C. Dilute urine sample 1:10 with deionized water.

### TEST PROCEDURE

Bring reagent, standard and sample to room temperature or 37°C.

#### Sample Start

| Pipette into test tubes  | Reagent Blank Tube | Standard Tube | Sample Tube |
|--|--------------------|---------------|-------------|
| Working reagent  | 1000 µl            | 1000 µl       | 1000 µl     |
| Sample   | -                  | -             | 25 µl       |
| Standard   | -                  | 25 µl         | -           |
| Mix, incubate at 37°C for 5 minutes. Read absorbances of standard (As) and samples (Ax) against reagent blank. |                    |               |             |

### CALCULATION

Serum/plasma sample: uric acid mg/dl =  $A_x/A_s \times 5$  (standard value) Random urine sample: uric acid mg/dl =  $A_x/A_s \times 5 \times 10$  (standard value and dilution) 24 hours urine sample (uric acid mg/24h): uric acid mg/24h =  $A_x/A_s \times 5 \times 10 \times$  diuresis (dl) (standard value, dilution and diuresis in dl)

### UNIT CONVERSION

mg/dl  $\times$  59.4 = µmol/L

### LINEARITY

the method is linear up to 25 mg/dl.

If the value is exceeded, it is suggested to dilute sample 1+9 with saline and to repeat the test, multiplying the result by 10.

#### Sensitivity/limit of detection (LOD)

the limit of detection is 0.16 mg/dl.

### INTERFERENCES

no interference was observed by the presence of:  
hemoglobin ≤ 150 mg/dl  
bilirubin ≤ 12 mg/dl  
lipids interference is observed

#### Precision

|          | intra-assay (n=10) | mean (mg/dl) | SD (mg/dl) | CV%  |
|----------|--------------------|--------------|------------|------|
| sample 1 |                    | 4.60         | 0.04       | 0.90 |
| sample 2 |                    | 10.72        | 0.04       | 0.40 |
|          | inter-assay (n=20) | mean (mg/dl) | SD (mg/dl) | CV%  |
| sample 1 |                    | 4.57         | 0.04       | 0.90 |
| sample 2 |                    | 10.72        | 0.11       | 1.00 |

### Methods comparison

a comparison between Archem and a commercially available product gave the following results:

Uric acid T FL Archem = x

Uric acid competitor = y

n = 104

y = 0.976x - 0.026 mg/dl r<sup>2</sup> = 0.99

### EXPECTED VALUES\*

Serum/plasma samples:

Men: 3.5 - 7.2 mg/dl (0.21 - 0.42 mmol/l)

Women: 2.6 - 6.0 mg/dl (0.15 - 0.35 mmol/l)

24h urine: 250 - 750 mg/24h (1.50 - 4.50 mmol/l)

\* It is recommended that each laboratory establish its own normal range.

### QUALITY CONTROL

All control sera with uric acid values determined by this method can be used. We recommend:

"ARCON N", Assayed Control Serum Normal

**Cat.No. A3910**

"ARCON P", Assayed Control Serum Abnormal

**Cat.No. A3920**

### CALIBRATION

The assay requires the use of an Uric Standard or an Uric Calibrator. We recommend:

ARCHEM Standard

**Cat.No. A2340S (6 mg/dl)**

Any commercially available Standard or Calibrator suitable for this method may be used.

### AUTOMATION

Special adaptations for automatic analyzers can be made on request.

### REFERENCES

- Barham D., Trinder P. - Analyst, 97 142 (1972)
- Fossati P., Prencipe L., Berti G. - Clin. Chem. 26 277 (1980).
- Tamaoku K., Murao Y., Akiura K., Ohkura Y. - Anal. Ch. Acta, 136
- Tietz Textbook of Clinical Chemistry, Second Edition, Burtis-Ashwood (1994).
- HU Bergmeyer - Methods of enzymatic analysis, (1987).

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