



PRESTIGE 24i LQ UA PLUS

DIAGNOSTIC KIT WITH ASCORBATE OXIDASE FOR DETERMINATION OF URIC ACID CONCENTRATION

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 urate 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 and in urine depends on glomerular filtration, thus is useful for renal function monitoring.

METHOD PRINCIPLE

Enzymatic, colorimetric method with uricase and peroxidase.

uric acid + 2
$$H_2O + O_2$$

allantoine + $CO_2 + H_2O_2$

ADPS + 4-aminoantipyrine + 2 H_2O_2 — point quinoneimine dye + 4 H_2O

(coloured compound)

The colour intensity is proportional to the uric acid concentration.

REAGENTS Package

Tuchage	Cat. No 4-209	Cat. No 4-409
	(24-TRAY)	(36-TRAY)
1-Reagent	6 x 40 ml	8 x 23 ml
2-Reagent	6 x 12.5 ml	8 x 7.5 ml

The reagents when stored at 2-8°C are stable up to expiry date printed on the package. Stability on board of the analyser at 2-10°C: Prestige 24i - 12 weeks, Biolis 24i Premium - 12 weeks. Protect from light and avoid contamination!

Concentrations in the test

buffer PIPES (pH 7.0)	100 mmol/l
4-aminoantipyrine	0.78 mmol/l
ADPS	0.67 mmol/l
ferricyanide potassium	3.8 µmol/l
peroxidase (POD)	> 38.34 µkat/l
uricase	> 1.65 µkat/l
ascorbate oxidase	> 66.7 µkat/l
sodium hydroxide	< 1 %

Warnings and notes

- Product for in vitro diagnostic use only.
- The reagents contain sodium azide (< 0.1%) as a preservative. Avoid contact with skin and mucous membranes.
- 1-Reagent meeting the criteria for classification in accordance with Regulation (EC) No 1272/2008.



H315 Causes skin irritation. H319 Causes serious eye irritation.

P280 Wear protective gloves/protective clothing/eye protection/face protection.

P302+P352 IF ON SKIN: Wash with plenty of soap and water. P305+P351+P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

SPECIMEN

24- hours urine, serum, heparinized plasma free from hemolysis.

Do not use EDTA, fluoride and oxalate as anticoagulants!

Urine preparation: To prevent precipitation of salts of uric acid, 10 ml of NaOH (500 g/L) should be added to the collection bottle before collection of a 24-hour specimen. Urine should be diluted with distilled water in the ratio of 1 to 4 (multiply the result by 5).

Serum and plasma can be stored 3-5 days at 2-8°C or 6 months at -20°C. 24-hours urine samples can be stored approximately 3 days at room temperature.

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

PROCEDURE

These reagents may be used in automatic analysers analysers Prestige 24i, Biolis 24i, Sapphire 400 and Prestige 24i Premium, Biolis 24i Premium, Sapphire 400 Premium.

- 1-Reagent and 2-Reagent are ready to use.
- 1-Reagent put on basic position in reagent tray.
- 2-Reagent put on start position in reagent tray.

For reagent blank deionized water is recommended.

REFERENCE VALUES 5

serum / plasma	mg/dl	µmol/l
females	2.5 - 6.8	149 - 405
males	3.6 - 7.7	214 - 458
24-hours urine	mg/24h	mmol/24h
	250 - 750	1.49 - 4.46

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

QUALITY CONTROL

For internal quality control it is recommended to use the CORMAY SERUM HN (Cat. No 5-172) and CORMAY SERUM HP (Cat. No 5-173) for determination in serum or CORMAY URINE CONTROL LEVEL 1 (Cat. No 5-161) and LEVEL 2 (Cat. No 5-162) for determination in urine with each batch of samples.

For the calibration of automatic analysers systems the CORMAY MULTICALIBRATOR LEVEL 1 (Cat. No 5-174; 5-176) and LEVEL 2 (Cat. No 5-175; 5-177) is recommended.

The calibration curve should be prepared every 12 weeks (Prestige 24i, Biolis 24i Premium), 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 the automatic analyser Prestige 24i. Results may vary if a different instrument or a manual procedure is used.

- Sensitivity (serum / plasma): 0.16 mg/dl (9.52 µmol/l).
 Sensitivity (urine): 0.18 mg/dl (10.71 µmol/l).
- Linearity (serum / plasma): up to 32 mg/dl (1903 µmol/l). Linearity (urine): up to 67 mg/dl (3985 µmol/l). For higher concentration of uric acid in serum or plasma, dilute the sample with 0.9% NaCl and repeat the assay. Multiply the result by dilution factor.

Specificity / Interferences

Haemoglobin up to 1.25 g/dl, ascorbate up to 62 mg/l, bilirubin up to 20 mg/dl and triglycerides up to 1000 mg/dl do not interfere with the test.

Precision

Repeatability (run to run)	Mean	SD	CV
n = 20	[mg/dl]	[mg/dl]	[%]
level 1	4.81	0.03	0.61
level 2	9.18	0.06	0.70

Method comparison

A comparison between uric acid values determined at Prestige 24i (y) and at COBAS INTEGRA 400 (x) using 60 serum samples gave following results: y = 0.9909 x + 0.1649 mg/dl;

R = 0.9970 (R – correlation coefficient)

A comparison between uric acid values determined at Prestige 24i (y) and at ADVIA 1650 (x) using 84 urine samples gave following results: y = 0.9161 x + 0.4401 mg/dl;

R = 0.9939 (R – correlation coefficient)

These metrological characteristics have been obtained using the automatic analyser Biolis 24i Premium. Results may vary if a different instrument or a manual procedure is used.

Sensitivity (serum / plasma): 0.21 mg/dl (12.49 µmol/l). Sensitivity (urine): 0.71 mg/dl (42.23 µmol/l). Linearity (serum / plasma): up to 29 mg/dl (1725 µmol/l). Linearity (urine): up to 67 mg/dl (3985 µmol/l).
 For higher concentration of uric acid in serum or plasma, dilute the sample with 0.9% NaCl and repeat the assay. Multiply the result by dilution factor.

Specificity / Interferences

Haemoglobin up to 1.25 g/dl, ascorbate up to 62 mg/l, bilirubin up to 20 mg/dl and triglycerides up to 1000 mg/dl do not interfere with the test.

Precision

Repeatability (run to run)	Mean	SD	CV
n = 20	[mg/dl]	[mg/dl]	[%]
level 1	4.95	0.04	0.73
level 2	8.67	0.14	1.63
Reproducibility (day to day)	Mean	SD	CV
n = 80	[mg/dl]	[mg/dl]	[%]
level 1	4.74	0.23	4.75
level 2	8.85	0.19	2.20

A comparison between uric acid values determined at Biolis 24i Premium (y) and at COBAS INTEGRA 400 (x) using 41 serum samples gave following results:

y = 0.9804 x + 0.0771 mg/dl;R = 0.9971 (R - correlation coefficient)

A comparison between uric acid values determined at Biolis 24i Premium (y) and at ADVIA 1650 (x) using 83 urine samples gave following results: y = 0.9154 x + 0.8018 mg/dl;

R = 0.9953 (R – correlation coefficient)

WASTE MANAGEMENT

Please refer to local legal requirements.

LITERATURE

- 1. Thefeld C. et al.: Dtsch. Med. Wschr. 98, 380-384 (1973).
- 2. Barham D., Trinder P.: Analyst 97, 142-145 (1972).
- 3. 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).
- Tietz N.W., ed. Clinical Guide to Laboratory Tests, 3rd ed. Philadelphia, PA: WB Saunders, 624, (1995).

APPLICATION for Prestige 24i, Biolis 24i and Sapphire 400

Item name	48	UA plu	IS								
Data infor	mation			Calibr	ation						
Units mg/dl				Type							
Decimals 1			Standard								
				#1	*	* #4					
Analysis				#2	*						
Туре		END		#3			#6				
Main W.Le	ength1	546									
Sub W.Ler		700		Norma	al Rang	е					
Method	0	Uricase	e			Ma	le	F	emale		
			-		Lo		High	Low			
Corr				Serum	3		7.7	2.5	6.8		
	Slope Inter										
	000	X+ 0.	.000	Plasma	ı <u>3</u>	.6	7.7	2.5	6.8		
				CSF							
				Dialysi	is						
				Other							
Aspiration Kind	ı Doub	le	Data Rea	a Process d			Absor	·hance	Limit		
	2040			Start	En	d					
Ī	Volume		Mair		54	-	Hig		3.000		
Sample	4		Sub	30	31		8				
Reagent1	200	μl									
Reagent2	50		Facto	r		Endpoint Limit			2.000		
			Blank	correction	correction 1.0000 Linear Check (%)						
									•		
Third Mix.	-			ition							
R1 Blank Water-Blank Dilue				ient	10	D:Dil	2				
Monitor			Pro	zone Chec	k						
0 Level Po	int 1				Start		End L		mit (%)		
Span	3.0	000	First	t							
			Seco	ond					Low		
			Thir	ď					Low		

Item name	48	UA plus			
Auto Reri	un SW		Auto Reru	n Condition (A	bsorbance)
ON			Absorbance	Range	
			-	Lower	OFF
Auto Reru	un Range	(Result)		Higher	OFF
	ON	ON			
	Lower	Higher	Prozone Rat	nge	OFF
Serum	0.16	23	-		
Urine	0.18	67			
Plasma	0.16	32			
CSF					
Dialysis					
Other					

APPLICATION for Prestige 24i Premium, Biolis 24i Premium and Sapphire 400 Premium

Sappinre													
Item No	48	Item	Name	UA	plus							Opti	cal
- ·						<i></i>							
Data inf	ormati	on				Calib		tion					
Units			mg/dl			Type Linear2							
Decimal	s		1			Std sa	am	ple coi	nc.				
						Blank		0	#1	*		#2	*
Analysis	5					#3			#4	1		#5	
Туре		l	END m	ethod		#6							•
Main W	ave Len	oth	546nm					r					
		-	700nm										
Sub Way	ve Leng	ui											
Method			Uricase										
Correla	tion												
			Int	roort									
	Slope	1		ercept									
Y=	1	X	+ 0										
Y . X Y	40	Υ.		T T 1		1						0	1
Item No.	48	Item	Name	UA	plus	1				L		Optic	al
Achiroti	ion					n.	ate	Proce	ee				
Aspirati Kind		սիեշ					ata ead		55	6	start	1	End
	D0	uble				K	ad	\vdash	Mair	-			End
Vol.		• •	1 /	11 1	r	4		F	Main	_	51	_	52
	Kind	Vo			Units	1			Sub		30		31
Sam	•	4	-	-	μl	l							
	gent 1	16	-	-	μl	Al	os.I	Limit	Lov	v	-		ligh
Rea	gent 2	4	0 1	0	μl			[-0.1		~	3	
									-				
Blank v								ection		•	_		
Water E	Blank							corre					
								Point I			2		
Reaction	n <u>M</u> oni	tor				Li	nea	ar Che	ck (%)				
0 Level			1			1							
Span			3			Pr	oz	one Cl	heck				
~						'	Start End Limit (%)					t (%)	
Third m	ivina					Fi	First						
OFF	uaing						c01			-			Low
OFF						36	<i>ι</i> υ	uu					LUW
Item No.	48	Item	Name	IIA	plus	1				1		Optic	•a1
nem ivo.	-10	neifi	vanie	UA	Pius	L				L		opin	aı
Normal	Range					Pani	c R	ange					
i mali		ale	Fe	emale		Panic Range Male Female						ale	
		High		Hig	rh			Low		igh	Lo		High
Serum					<u> </u>	Serur	n	LOW	n	igii		, vv	ringii
	3.6	7.7	2.5	6.	0				_				
Urine				-	_	Urine			_				
Plasma	3.6	7.7	2.5	6.	8	Plasn	na						
CSF						CSF							
Dialysis						Dialy							
Other						Other							
Item No.	48	Item	Name	UA	plus					T		Optic	al
						-				-			
Auto Re	erun SV	V											dition
ON									(Ab	sorb	ance	e)	
								-	Lo	wer	()FF	
Auto Re	run Ra	nge (Conc.)						Hi	gher	()FF	
	First	0. (Low			High							
	Dil	Re	Value	Dil	Re	Value	Г	Dil	Ant	0 R4	run	Cor	dition
Serum	~ 11	ne	0.21			29	Ľ			zon		001	anon
Urine							-	_			-,		
			0.71			67	_	_	U	r			
Plasma			0.21		<u> </u>	29							
CSF		L			L					lutio			
Dialysis					<u> </u>				10	0:Dil	2		
Other				1	1								

Date of issue: 06. 2015.

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 <u>http://www.cormay.pl</u>