

Colorimetric Nitro-PAPS method. Kit for measurement of zinc concentration in serum, plasma and urine.



1 485

1,603



1 03

0,73

10600

5x8 mL (R1) 1x10 mL (R2) 1x5 mL (STD)



## PRINCIPLE OF THE METHOD

Zinc reacts with NTRO-PAPS yielding at room temperature an intensely colourted complex which intensity is proportional to the concentration of the Zinc present in the sample. The method doesn't need de-proteinization, neither sample blank.

#### **TEST PREFORMANCE**

Precision				
Intra-assay (n=10)	Mean (μg/dL)	SD (µg/dL)	CV (%)	
Sample 1	148,5	2,456	1,65	
Sample 2	224,4	1,831	0,82	
Inter-assay (n=30)	Mean (μg/dL)	SD (μg/dL)	CV (%)	

Method comparison

Sample 1

Sample 2

Results obtained on the same samples with an equivalent method have done the following

143 3

218,7

y= 0,9931x-1,0654 R = 0.9996

Sensitivity/limit of detection

The method is able to discriminate until 3 µg/dL.

The method is linear up to 1000  $\mu g/dL$ .

#### SAMPLE

Serum, plasma with heparin as anticoagulant. Urine 24 hours.

- Avoid use of hemolyzed sample.
- Zinc in the sample is stable for 8 days at +2-8°C.
- Strong lipemic serum can interfere with the measurement . Centrifuge or filtrate the sample before testing

## REFERENCE VALUE

	SERUM AND PLASMA					
	70 – 115	μg/dL		10.7 – 17.6	μmol/L	
_	URINE					
Ĺ	100 - 1000	μg/24h		15.3 – 153	μmol/24h	Ī

## **REAGENTS - INITIAL CONCENTRATION**

Reagent 1	Borate buffer 0,37 M pH 8.2 Salicilaldoxime 12,5 mM Dimetilglioxime 1.25 mM Surfactants and preservatives.	5x8mL	REF: 1060001
Reagent 2	NITRO-PAPS; 0,4 mM Preservatives.	1x10mL	REF: 1060002
Standard	Standard Zinc ion 200 ug/dL	1x5 mL	REF: 1060003

## REAGENTS -STORAGE AND STABILITY

Kit: Store at +2-8°C. Stable until the expiry date shown on the label.

Working Reagent: Store at +2-8°C.

Stable 15 days

Reagents are stable after opening until the expiry date shown on the Opened reagents:

bottles when are protected from direct light, tightly closed, and stored

at reported temperature

## **REAGENTS - PREPARATION**

Reagents are liquid ready to use.

To prepare the Working Reagent add 2 mL of Reagent 2 to a vial of Reagent 1.

Alternately mix 4 volumes of Reagent 1 with 1 volume of Reagent 2 depending on the number of sample,

## **REAGENTS - PRECAUTION AND WARNING**

- This method describes the manual use of this kit. For use with automatic analyzer see the specific applications.
- Quality control data sheet of the reagents are available upon request. Refer to the batch number on the label

## ADDITIONAL EQUIPMENT

Spectrophotometer Cuvettes with 1 cm lightpath.

## SAFETY PRECAUTIONS

Reagent 1

- Causes serious eve damage [H318]

The product is not classified as dangerous

Danger

Standard

- Harmful to aquatic life with long lasting effects.[H412]
- Contact with acids liberates very toxic gas.[EUH032]

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

IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. [P305 + P351 + P3381

- Immediately call a POISON CENTER or doctor/physician. [P310]

Reagent 2 The product is not classified as dangerous

# WASTE MANAGEMENT

Please refer to local legal requirements

#### ANALYTICAL PROCEDURE

Wavelength  $\lambda = 578(520 \text{ or } 570) \text{ nm}$ 

Lightpath 1 cm

Temperature Room temperature (+20-25°C) Measurement Against reagent blank end point (increase)

Allow reagents to reach working temperature before using

	DEAIN	STANDARD	SAIVIFLL	
Working reagent	1000	1000	1000	μL
Mix thoroughly and incubate for 1 minute at room temperature (+20-25°C).				
Measure the absorbance of	the Sample	and the Standard ag	ainst the blank	(ABS1).
Distilled water	50	-	-	μL
Sample	-	-	50	μL
Standard	-	50	-	μL

Mix thoroughly and incubate for 5 minutes at room temperature (+20-25°C). Measure the absorbance of the Sample and the Standard against the blank (ABS2). The colour is stable for at least 30 minutes at (+20-25°C), protected from direct light

## **CALCULATIONS**

Serum, plasma or Urine:

$$\label{eq:Zinc} \mbox{Zinc ($\mu g\slashed{g}\slashed{dL}) or ($\mu g\slashed{g}\slashed{dL}$) or ($\mu g\slashed{g}\slashed{dL}$) or ($\mu g\slashed{g}\slashed{dL}$) and $(\mu g\slashed{dL}$) or ($\mu g\slashed{dL}$$$

Conversion factor:

Zinc ( $\mu$ g/dl) x 0.1529 = Zinc ( $\mu$ mol/l)

## QUALITY CONTROL

Each laboratory should estabilish its own internal Quality control scheme and procedures fpr corrective action if controls do not recover within the acceptable tolerances.

## **BIBLIOGRAPHY**

- Pasquinelli F., Diagnostica e Tecniche di Laboratorio, (pag..: 1099-1102) Rossini Ed. (1984).
   TETSUO MAKINO, Chimica Clinica Acta 197, 209-220 (1991)
- 3. MARINGONI A., ILLUZZI R., ATB 1991 Abstract.

## **SYMBOLS**



Read use istruction



Manufactured by

SAMDIE



Size/number of tests



In Vitro Diagnostic **Medical Device** 



CE mark declaring compliance with the In Vitro Diagnostics Medical Device Directive, 98/79/EC



Catalogue number



Store at +15-25°C



Batch number



Store at +2-8°C



LOT

Vial content