Colorimetric determination of zinc in serum, plasma and urine

CLINICAL SIGNIFICANCE

Zinc intervenes in the function and structure of more than 70 enzymes involved in various metabolic processes such as synthesis or degradation of carbohydrates, lipids, proteins and nucleic acids.

Zinc deficiencies cause anemia, hepatosplenomegaly, delay in development, delayed healing of wounds and ulcerations, taste and smell alterations.

Decreases in Zinc concentration can be observed in physiological conditions such as the last months of pregnancy, the use of oral contraceptives and in pathological conditions such as myocardial infarction, alcoholic cirrhosis, malabsorption syndrome, lung infections, carcinomas and lymphomas.

TEST SUMMARY

Zinc reacts with the chromogen present in the reagent forming a coloured compound which colour intensity is proportional to the zinc concentration present in the

SAMPLES

Serum or plasma unhemolized. Use heparin salt as anticoagulant.

Urine 24 hours. Stability 8 days at 2-8°C.

REAGENTS

Borate buffer Reagent A: 0.37 M 8.2: Saliciladoxime 12.5 Dimetilgioxime 1.25 mM; surfactants and preservatives.

NITRO-PAPS; 0.4 mM, preservatives. Reagent B:

Standard: Zinc ion 200 μ g/dl (30.6 μ mol/l); stabilizers and preservatives.

MATERIAL REQUIRED BUT NOT SUPPLIED

Normal laboratory equipment. Spectrophotometer UV/VIS with thermostatation. Automatic Micropipette. Cuvette in optical glass or monouse in optical polystyrene. Distilled water.

PRECAUTIONS

Reagent may contain not reactive and conservative components. It is opportune to avoid contacts with the skin and do not swallow.

Perform the test according to the general "Good Laboratory Practice" (GLP) guidelines.

REAGENTS PREPARATION

Add 2 ml of Reagent B to a vial of Reagent A. Reagents are stable until expiration date on label. stored at 2-8°C.

Work Reagent is stable 15 days at 2-8°C.

Warning: do not contaminate reagents after the vials opening.

PROCEDURE

Kind of analysis: End point Reading time: 5 minutes Colour stability: 30 minutes Wavelength: 578 nm (570-582) Temperature: 20-25°C Lightpath 1 cm Blank Reagent Zero:

REAGENTS	BLANK	STANDARD	SAMPLE
Work Reagent	1 ml	1 ml	1 ml
Distilled Water	50 μl		
Standard		50 μl	
Sample			50 μl

Mix and read the absorbance against blank at 578 nm Colour is stable for 30 minutes.

CALCULATION

SERUM / PLASMA

 $Zn \mu g/dI =$ [A_(sample) / A_(standard)] x 200 Zn umol/l = [A_(sample) / A_(standard)] x 30.6

URINE

 $Zn \mu g/24h =$ [A_(sample) / A_(standard)] x 200 x dl urine

$Zn \mu mol/24h =$ [A_(sample) / A_(standard)] x 30.6 x Lt urine

EXPECTED VALUES SERUM / PLASMA

70 - 115 ua/dl (10.7 - 17.6 µM)

URINE

 $100 - 1000 \mu g/24h$ (15.3 - 153 µmol/24h)

Every laboratory should establish own reference intervals in accordance with own population.

- Strong lipemic sera can sometimes interfere in the analysis: is suggested to centrifuge or filtrate the sample with membrane 0.2 µm.
- Use glassware surely without Zinc traces
- The volumes could be proportionally changed.
- If the results are incompatible with clinical presentation, they have to be evaluated within a total clinical study.
- . Only for IVD use.

CALIBRATION/ QUALITY CONTROL

It is suggested to perform an internal quality control using control serum with known zinc values.

TEST PERFORMANCE

Precision			
Intra-assay (n = 21)	Mean (μg/dl)	SD (μg/dl)	CV%
Sample 1	94.14	2.220	2.36

Inter-assay (n = 21)	Mean (μg/dl)	SD (μg/dl)	CV%
Sample 1	94.48	2.502	2.65

Sensivity/limit of detection

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

Linearity

The method is linear up to 1000 µg/dl.

Methods comparison

A comparison with a commercial available product gave the following results in a comparison on 21 samples of serum:

Zinc LTA = xZinc Acid competitor = y n = 21

y = 0.96262x + 4.49686r = 0.99331.

A comparison with a commercial available product gave the following results in a comparison on 21 samples of urine:

Zinc LTA = xZinc Acid competitor = y

y = 0.99539x + 2.9706r = 0.99984

Interferences

No interference was observed in presence of: bilirubin ≤ 20 mg/dl

Haemoglobin can interfere with the analysis.

WASTE DISPOSAL

Product is intended for professional laboratories. Waste products must be handled as per relevant security cards and local regulations.

PACKAGING

CODE CC02750 (50 TESTS)

(liquid) Reagent A 5 x 8 ml Reagent B 1 x 10 ml (liquid) Standard 1 x 5 ml (liquid)

REFERENCES

Pasquinelli F., Diagnostica e Tecniche di Laboratorio. (pag.: 1103 – 1104) Rossini Editrice. (1984). Tetsuo Makino, Chimica Clinica Acta 197, 209-220

Maringoni A., Illuzzi R., ATB 1991 Abstract.

MANUFACTURER

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SYMBOLS

IVD Only for IVD use

LOT Lot of manufacturing

REF Code number

Storage temperature interval X

Expiration date

⚠ Warning, read enclosed documents

 \prod i Read the directions

Biological risk

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