

# **MAGNESIUM CALMAGITE**

# Colorimetric determination of magnesium in biological liquids

#### **TEST SUMMARY**

Magnesium reacts with Calmagite forming a pink complex

#### **SAMPLES**

Serum

Diluted Urine 1:5 with Distilled water. Stability: 7 days at 4°C.

# REAGENTS

Amino-metil-prophanol 1 M; Reagent A EGTA 0.8 mM; stabilizers and conservatives.

Calmagite 0.4 mM: surfactants. Reagent B

stabilizers and preservatives.

Standard Ion Magnesium 2 mEq/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. Physiologic solution.

#### **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**

Mix a part of Reagent A with a part of Reagent B. Reagents are stable until expiration date on label stored at 15-25°C.

Work Reagent is stable 30 days at 15-25°C.

# **PROCEDURE**

Colour stability:

Kind of analysis: **End Point** Reading Time: 2 minutes Wavelength: 520 nm(500-550) Temperature: R.T. Lightpath: 1 cm Zero: Blank Reagent

Reagents	Blank	Standard	Sample
Distilled water Standard Sample	10 μl  	 10 μl 	  10 ul
Work Reagent	1000 ul	1000 ul	1000 ul

120 minutes

# CALCULATION

# SERUM Magnesium (mEq/l)

(A sample / A standard) x 2

## SERUM Magnesium (mM)

(A sample / A standard) x 1

# URINE Magnesium (mEq/24h)

(A sample / A standard) x 2 x 5 x l of urine

#### **EXPECTED VALUES**

(0.65 - 1.05 mM) Serum: 1.3 - 2.1 mEq/l

Urine: 6.0 - 8.5 mEq/24h

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

#### NOTE

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. For this purpose the following control sera on human base are available on request:

**QN 0050 CH** 10 x 5 ml

Control Sera normal values

QP 0050 CH 10 x 5 ml

Control Sera pathological values

#### **TEST PERFORMANCE**

#### Precision

Intra-assay (n = 20)	Mean (mEq/l)	SD (mEq/l)	CV%
Sample 1	1.605	0.039	2.46
Sample 2	2.100	0.032	1.54

Inter-assay (n = 20)	Mean (mEq/l)	SD (mEq/l)	CV%
Sample 1	1.605	0.039	2.46
Sample 2	2.090	0.039	1.18

The method is linear up to 5 mEq/l (2.5 mM).

# Methods comparison

A comparison with a commercial available product gave the following results in a comparison on 50 samples:

Magnesium LTA = x Magnesium competitor = y

y = 0.91847x + 0.15r = 0.9594

# WASTE DISPOSAL

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

# **PACKAGING**

CODE CC02070	(600 TESTS)		
Reagent A	3 x 100 ml	(liquid)	
Reagent B	3 x 100 ml	(liquid)	
Standard	1 x 5 ml	(liquid)	

### REFERENCES

Maxwell H. e Coll. - Clin. Chem. 28/3; 520 (1982). Henry J.B. - Clinical Diagnosis and Management - 17<sup>th</sup> edition - Saunders Publisher (1984). Savory J. e coll. - Clin Chem 31/3, 487 - 488 (1985).

#### **MANUFACTURER**

LTA s.r.l.

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#### **SYMBOLS**

IVD Only for IVD use

LOT Lot of manufacturing

REF Code number

1 Storage temperature interval

Expiration date

Æ Warning, read enclosed documents

 $\square$ i Read the directions Biological risk

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