



CREATININE LR liquid reagent

REF C2701220 1036 Test

CE C2701220A 1036 Test

IVD For in vitro medical device

"Intra-Assay" precision (within-Run)

Determined on 20 samples for each control (N-H) (Normal-High).

Results:

MEAN [mg/dl] N = 1.15 H = 3.87

S.D. N = 0.05 H = 0.13

C.V.% N = 4.6 H = 3.4

"Inter-Assay" precision (between-Run)

Determined on 20 samples for each control (H-N). Results:

MEAN [mg/dl] N = 1.15 H = 3.86

S.D. N = 0.05 H = 0.08

C.V.% N = 4.19 H = 2.02

Correlation

A study based comparing this method with a similar method on 20 samples has given a correlating factor $r = 0.98$

$$y = 0.98x + 0.0775$$

Interferences

No interference was observed by the presence of:

Bilirubin ≤ 15 mg/dl.

Triglycerides ≤ 1500 mg/dl

Hemoglobin ≤ 150 mg/dl

Quality Controls

It's necessary, each time the kit is used, to make the quality controls and to check that values obtained are within the acceptance range provided in the insert. Each laboratory should establish its own mean and standard deviation and adopt a quality control program to monitor laboratory testing.

Bibliography

Henry, R.J.: Clinical Chemistry: Principles and Techniques, Harper & Row.N.Y., p. 287, (1964).

Young et al., Clinical Chemistry, 18, (1972).

Newman, D.J., Price, C.P.: Renal function and nitrogen metabolites. In: Burtis CA, Ashwood ER, editors. Tietz Textbook of Clinical Chemistry. 3rd ed. Philadelphia: W.B. Saunders Company; p. 1204, (1999). Kaplan, L.A., Pesce, A.J.: "Clinical Chemistry", Mosby Ed. (1996).

Symbols

CE CE Mark (98/79 CE regulation)

IVD in vitro medical device

LOT Batch Code

Hourglass icon Use by

Thermometer icon Storage temperature limits

Book icon Read instruction for use

Factory icon Gesán Production srl

Use

Kit for measurement of creatinine in serum, plasma and urine Colorimetric method Jaffé without deproteinization

Summary

Creatinine measurements are used in the diagnosis and treatment of renal diseases and in monitoring renal dialysis.

Principle

Fixed time analysis. Creatinine reacts in alkaline environment with picrate to give a coloured compound whose intensity is proportional to the creatinine concentration in the sample.

Reagents

R1	Litium hydroxide	120.0 mmol/l
	Boric acid	80.0 mmol/l
R2	Picric acid	67.0 mmol/l

Reagent Preparation

Reagents are liquid and ready to use. About using as monoreagent ("sample starter" procedure) pour the content of R2 vial into the R1 vial. For minor use mix the reagents R1 and R2 in equal parts.

Storage and Stability

- Store the kit at 15-25°C
- After opening, the vials R1, R2 are stable 90 day if recapped immediately and protected from contamination, evaporation, direct light, and stored at the correct temperature.
- Working solution stability (R1+ R2): 7 days at 15-25°C.

Precaution In Use

The product is not classified as dangerous (DLg. N. 285 art. 28 l. n. 128/1998). However the reagent should be handled with caution, according to good laboratory practice.

Waste Management

Please refer to the local legal requirements.

Specimen Collection and Preparation

- Serum or plasma. Urine.
- Do not use samples with haemolysis.
- The creatinine is stable in the samples up to 24 hours at 2-8°C.

Note

- The kit, according to this method, must be used in manual procedures. About automatic using follow specific applications.
- Avoid direct light, contamination and evaporation.
- The volumes in the procedure can be changed proportionally.
- In case of complaint or quality control request, refer to the lot number on the package or the lot number on the singles vials.

Procedure

Wavelength λ : 510 (500-550) nm
 Working Temperature 37°C
 Optical Path 1 cm
 Reaction "fixed time"

Monoreagent Procedure "sample starter"

	Blank	STD	Sample
Working Reagent	1000µl	1000µl	1000µl
Distilled Water	100 µl	-	-
Sample	-	-	100 µl
Standard	-	100 µl	-

Mix and after 30" at 37°C measure the absorbance of sample (E1C) and standard (E1STD) against the reagent blank. After other 2' make the second reading (E2C), (E2STD).

- Bireagents Procedure "substrate starter"

	Blank	STD	Sample
Reagent R1	500µl	500µl	500µl
Distilled Water	100 µl	-	-
Sample	-	-	100 µl
Standard	-	100 µl	-

Mix, incubate at 37°C for 1', then add :
 Reagent R2 500 µl 500 µl 500 µl
 Mix and after 30" at 37°C measure the absorbance of sample (E1C) and standard (E1STD). After other 2' make the second reading (E2C), (E2STD).

Calculation

$$\text{Creatinine [mg/dl]} \text{ o } [\mu\text{mol/l}] = \frac{(E2C - E1C)}{(E2STD - E1STD)} \times \text{Conc. STD}$$

The reagent performances are related to 37°C, 1 cm and 510 nm.

Conversion Factor

$$\text{Creatinine [mg/dl]} \times 88.4 = \text{Creatinine } [\mu\text{mol/l}]$$

Reference Values

Serum - plasma
Man 0.9 - 1.3 mg/dl (80 - 115 µmol/l)
Woman 0.6 - 1.1 mg/dl (53 - 97 µmol/l)
Urine 0.80 - 1.80 g/24h

Reference values are considered indicative since each laboratory should establish reference ranges for its own patient population. The analytical results should be evaluated with other information coming from patient's clinical history.

ANALYTICAL PERFORMANCES

Linearity

Reaction is linear up to a concentration of 25 mg/dl (2210 µmol/l) with a range of 0.3-25 mg/dl (27-2210 µmol/l). Samples with values exceeding 25 mg/dl must be diluted with saline solution. Multiply, then, the result for diluting factor.

Analytical sensitivity

The test sensitivity in terms of detection limit is 0.3 mg/dl (27 µmol/l).

Creatinine LR

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