IRON CRX

FE 0100 CH 2 x 50 ml FE 0400 CH 4 x 100 ml

INTENDED USE

Reagent for quantitative in vitro determination of iron in biological fluids.

SUMMARY OF TEST

Serum iron concentration connotes the Fe(III) bound to serum transferrin and does not include the iron contained in serum as free hemoglobin.

PRINCIPLE OF THE METHOD

Serum iron reacts with chromazurol-B and CTMA-Br to form a complex deeply blue colored. The absorbance measured at 630 nm is directly proportional to the amount of iron in the sample.

KIT COMPONENTS

For in vitro diagnostic use only.

The components of the kit are stable until expiration date on the label.

Keep away from direct light sources.

FE CRX R1 0100: 2 x 50 ml (liquid) blue cap 0400: 4 x 100 ml (liquid) blue cap

Composition: chromazurol-B 0.13 mM, CTMA-bromide

Standard: iron(III) solution 200 µg/dl - 5 ml

Store all components at 15-25°C.

0.82 mM, acetate buffer pH 4.75.

MATERIALS REQUIRED BUT NOT SUPPLIED

Current laboratory instrumentation. Spectrophotometer UV/VIS with thermostatic cuvette holder. Automatic micropipettes. Glass or high quality polystyrene cuvettes. Saline solution.

REAGENT PREPARATION

Reagent R1: ready to use.

Stability of reagent: up to expiration date on labels at $15-25^{\circ}\mathrm{C}$

Stability since first opening of vials: preferably within 60 days at 15-25°C.

PRECAUTIONS

FE CRX R1: Warning. Causes serious eye irritation (H319). Causes skin irritation (H315). Harmful to aquatic life with long lasting effects (H412).

Wear protective gloves / eye protection / face protection (P280). If eye irritation persists: Get medical advice / attention (P337+P313). Wash with water thoroughly after handling (P264). Avoid release to the environment (P273).

Standard: It is not classified as hazardous

SPECIMEN

Serum. Samples are stable 4 days at 2-8°C. Separate serum/plasma from clot within 1 hour.

TEST PROCEDURE

Wavelenght: 630 nm (allowed 620 \div 640 nm) Lightpath: 1 cm Temperature: 25, 30 or 37°C

Mix, incubate at 25, 30 or 37°C for 5 minutes.

Read absorbances of standard (Ac) and samples (Ax) against reagent blank.

RESULTS CALCULATION

serum/plasma sample:

iron
$$\mu$$
g/dl = $\frac{Ax}{Ac}$ x 200 (standard value)

EXPECTED VALUES

men 59 - 158 μg/dl (10.6 - 28.3 μmol/l) women 37 - 145 μg/dl (6.60 - 26.0 μmol/l)

Each laboratory should establish appropriate reference intervals related to its population.

QUALITY CONTROL AND CALIBRATION

It is suggested to perform an internal quality control. For this purpose the following human based control sera are available:

QUANTINORM CHEMA - MULTINORM CHEMA

with normal or close to normal control values QUANTIPATH CHEMA - MULTIPATH CHEMA with pathological control values.

Please contact Customer Care for further information.

TEST PERFORMANCE

Linearity

the method is linear up to 500 µg/dl.

If the limit value is exceeded, it is suggested to dilute sample 1+9 with distilled water and to repeat the test, multiplying the result by 10.

Sensitivity/limit of detection (LOD)

the limit of detection is 12 µg/dl.

Interferences

no interference was observed by the presence of:

hemoglobin interferes bilirubin ≤ 50 mg/dl lipids interferes

Precision

intra-assay (n=10)	mean (μg/dl)	SD (μg/dl)	CV%
sample 1	111.70	2.31	2.10
sample 2	172.10	2.28	1.30
inter-assay (n=20)	mean (μg/dl)	SD (μg/dl)	CV%
sample 1	111.79	2.66	2.40
sample 2	170.58	2.53	1.50

Methods comparison

a comparison between Chema and a commercially available product gave the following results:

Iron CRX Chema = x Iron competitor = y n = 95

 $y = 1.08x - 8.71 \mu g/dl$ $r^2 = 0.97$

WASTE DISPOSAL

This product is made to be used in professional laboratories.

P501: Dispose of contents according to national/international regulations.

REFERENCES

Garcic A. - Clin. Chim. Acta 94, 115 (1979)

Tietz Textbook of Clinical Chemistry, Second Edition, Burtis-Ashwood (1994).

MANUFACTURER

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SYMBOLS

IVD in vitro diagnostic medical device

LOT batch code

REF catalogue number

consult instructions for use