

# IRON

Colorimetric Method

50 Tests

## PRINCIPLE :

The serum is deproteinized by trichloroacetic acid and the iron is dissociated from the protein transferrin by hydrochloric acid then reduced to ferrous by thioglycolic acid . The colored complex which the iron forms with bathophenanthroline is measured colorimetrically

## SAMPLE :

Serum . Samples free of hemolysis, can be stored for 24 hours at +2 to +8°C

## NORMAL VALUES :

Adults : 7.3 – 23.6 µmol/L (41-132 µg/dl)

## REAGENTS :

1.	Standard iron	35.8 µmol/L (200 µg/dL)
2.	Acids mixture Trichloroacetic acid Hydrochloric acid Thioglycolic acid	0.6 mol/L 2.1 mol/L 0.4 mol/L
3.	Chromogen Bathophenanthroline Sodium acetate	0.47 mmol/L 3.6 mol/L

## STABILITY :

The reagents are stable up to the expiry date specified when stored at +15 to +25 °C .

## PROCEDURE :

### Deproteinization :

Serum	0.5 ml
Reagent 2	0.5 ml

Mix, let stand for 10 min. then, centrifuge for 10 min. at 3000 rpm. Use the supernatant for the assay.

Pipette into test tubes ( FREE OF IRON ) :

	Blank ml	Standard ml	Sample ml
Supernatant	-	-	0.5
Reagent 1	-	0.25	-
Reagent 2	0.25	0.25	-
Dist. water	0.25	-	-
Reagent 3	0.50	0.50	0.50

Mix, incubate for 5 min. at 20 - 25°C. Measure the absorbances of the sample ( $A_{\text{Sample}}$ ) and the standard ( $A_{\text{standard}}$ ) against blank, at 535 nm. Color stable for one hour. Linearity up to 143 µmol/L (800 µg/dl)

## CALCULATION :

Iron Concentration

$$= \frac{A_{\text{Sample}}}{A_{\text{Standard}}} \times \text{Standard Conc.}$$

## QUALITY CONTROL :

For accuracy and reproducibility control:-  
Assayed Multi – Sera, Normal and Elevated

## REFERENCE :

Dreux, C. ( 1977 ) ; Ann. Biol. Clin. 35 : 275 .

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+15 to +25°C

50 Tests

In vitro diagnostic use

**CAT. NO.**

**IR 15 10**

## REAGENTS

<b>R1</b> Standard	<b>15 ml</b>
<b>R2</b> Acids Mixture	<b>25 ml</b>
<b>R3</b> Chromogen	<b>25 ml</b>

## CONTACTS

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