COPPER ION



1. PERFORMANCE

Tube No.

1) Sampling method	Direct sampling method
	(Refer to Page 17)
2) Measuring range	: 1-100mg/l
3) Sampling time	: 1 to 2 minutes
4) Sample volume	: over 5 mℓ
5) Detectable limit	: 0.5mg/l
6) Shelf life	: 1 year
7) Operating temperature	: 0∼40 °C
8) Operating PH	: 2-11
9) Reading	: Direct reading from the scale
10) Colour change	: White→Orange

2. RELATIVE STANDARD DEVIATION

RSD-low: 10% RSD-mid.: 5% RSD-high: 5%

3. CHEMICAL REACTION

By reacting with Hydroxylamine sulphate, divalent Copper ion is reduced to monovalent Copper ion. This monovalent Copper ion is reacted with 2, 9-Diphenyl 1-4, 7-Dipheny 1-1, 10-phenanthroline and Chelate is produced.

 Cu^{2+} + NH₂OH + H₂SO₄→ Cu^{+} Cu^{+} + $C_{12}H_4N_2$ (CH₃)₂ (C₆H₅)₂→ Orange Chelate compound

4. CALIBRATION OF THE TUBE

CUPRIC SULPHATE STANDARD SOLUTION METHOD

5. INTERFERENCE AND CROSS SENSITIVITY

Substance	mg/ℓ	Interference	mg/ℓ	Coexistence
Ferric ion	20	Similar stain is produced.	Copper ion conc.×2	Higher readings are given.
Zinc ion		The accuracy of readings in not affected.	100	"
Chlorine ion		"		"
Manganous ion		11		"

(NOTE)

6. SAMPLING METHOD

- 1) Cut both ends of a fresh detector tube with a file.
- Squeeze the rubber bulb (an extra option), insert the tube end(B) into it as it is and immerse filled end(A) of the tube.
- 3) Put the thumb off the rubber bulb, and the sample solution shall rise up.
- 4) When the sample solution rises up to (C) of the tube, remove the tube from the rubber bulb and from thesample solution.
- 5) The concentration can be obtained directly from the reading value of scale printed on the tube.
- 6) When the concentration is over 100mg/l, dilute the sample solution and multiply the readings obtained by the dilution ratio.

