

EZ2302 Total Copper & Copper (II) Analyser

Method and reagent sheets

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1. Legal information

Manufacturer: AppliTek NV/SA

Distributor: Hach Lange GmbH

The translation of the manual is approved by the manufacturer.

2. Analytical specifications

Please refer also to the respective technical datasheet at Hach Support Online.

| Copper – All specifications | | | | | |
|-----------------------------|--|--|--------------------------|-------------------|--|
| Analysis method | Colo | Colorimetric measurement at 546 nm using bicinchoninate method | | | |
| Parameter | Tota | Il Cu & Cu (II) | | | |
| | Standard measurement cycle time: 30 minutes | | | | |
| Cycle time | Inte | nal dilution: + 5 min. | | | |
| | Exte | ernal dilution: + 5 – 10 min. | | | |
| Limit of detection (LOD) | ≤3 | ug/L | | | |
| Precision/Repeatability | Bett | er than 2% full scale range for sta | ndard test solutions | | |
| Cleaning | Auto | omatic; frequency freely programm | nable | | |
| Calibration | Automatic, 2-point; frequency freely programmable | | | | |
| Validation | Automatic; frequency freely programmable | | | | |
| | Acidity, metal ions like aluminium (III) [(Al) ³⁺)] > 10 mg/L, cyanide [(CN) ⁻)], hardness, | | | | |
| Interferences | $ \text{iron(III)}[(\text{Fe})^{3+}) > 10 \text{ mg/L}, \text{ nickel(II)}[(\text{Ni})^{2+}] \text{ and silver(II)}[(\text{Ag})^{+})]. \text{ Large amounts of colour}$ | | | | |
| | and | turbidity interfere. Fats, oil, proteir | ns, surfactants and tar. | | |
| Measuring ranges | % o | f range - Dilution | Low range (mg/L) | High range (mg/L) | |
| | Α | 10% of standard range | 0.003 | 0.30 | |
| | В | 25% of standard range | 0.02 | 0.75 | |
| | C 50% of standard range | | 0.02 | 1.50 | |
| | 0 standard range | | 0.03 | 3.0 | |
| | 1 | internal MP dilution (factor 4) | 0.2 | 12 | |
| | 3 | internal MP dilution (factor 10) | 0.5 | 30 | |

3. Analysis method

Summary

The determination of Copper (II) and total Copper is based on two methods, combined in one analyser.

The Copper (II) concentration is determined in the 'Cu' method. The total Copper concentration is determined in the 'Total Cu' method. The concentration of all parameters is determined alternately in the 'Main'-method.

The calibration for Copper (II) is performed in the 'Cu' method. The calibration for total Copper is performed in the 'Total Cu' method.

Remark

The methods cannot be started at the same time.

3.1 Copper (II)

Summary

The determination of the copper concentration in water is based on the reaction of copper ions with 2,2-biquinoline in an alkaline solution to an intense coloured pink complex. The absorption is measured at 546 nm.

Analysis steps

The analysis vessel is cleaned and filled with fresh sample. After sampling acid solution is added and the initial absorbance value is measured at 450 nm. This measurement is performed to correct for any colour contribution of the sample itself. Next, the colour solution is added and after respecting a stirring period – performed to obtain complete colour development – the final absorbance value is determined. With the obtained absorbance values, the copper concentration can be calculated according to Beer's law.

Calibration

The calibration procedure measures a REF1 Cu solution (channel 9, REF1 valve) and a REF2 Cu solution (channel 10, REF2 valve) to adapt the slope and offset factors by means of a two point calibration.

3.2 Total Copper

Summary

The determination of the copper concentration in water is based on the reaction of copper ions with 2,2-biquinoline in an alkaline solution to an intense coloured pink complex. The absorption is measured at 546 nm. . Prior to the total copper analysis, the sample is digested by use of an acid solution.

Analysis steps

The sample is mixed with acid solution and heated to 120 °C (or up to 150 °C – programmable) in an oven during several minutes (standard 10 minutes; programmable up to 60 minutes). After digestion, the sample is cooled and transferred into the analysis vessel. The initial absorbance value is measured at 546 nm. This measurement is performed to correct for any colour contribution of the sample itself. Next, the colour solution is added and after respecting a stirring period – performed to obtain complete colour development – the final absorbance value is determined. With the obtained absorbance values, the copper concentration can be calculated according to Beer's law.

Calibration

The calibration procedure measures a REF1 Cu solution (channel 9, REF1 valve) and a REF2 Cu solution (channel 10, REF2 valve) to adapt the slope and offset factors by means of a two point calibration.

4. Reagents

A CAUTION



Chemical exposure hazard. Obey laboratory safety procedures and wear all of the personal protective equipment appropriate to the chemicals that are handled. Read the safety data sheet from the supplier before bottles are filled or reagents are prepared. For laboratory use only. Make the hazard information known in accordance with the local regulations of the user.

A CAUTION



Chemical exposure hazard. Dispose of chemicals and wastes in accordance with local, regional and national regulations.

4.1 Reagent overview and consumption

In the tables below, the products that are needed to prepare the reagents are listed. The product name, the formula, the molecular weight, the CAS No. and the amount needed to prepare 1 liter of the reagents is given. Check the consumption of the reagents (28 days) to adapt the volumes needed.

| Product | Consumption | Consumption/28 days A rata 1 analysis/30 min | Recommended containers | |
|----------------------------------|---------------------|---|------------------------|--|
| Acid solution (1M) (Total Cu) | ~ 1.0 mL / analysis | ~ 1.4 L | Plastic – 2.5 L | |
| Acid solution (0.25M) (Cu) | ~ 1.0 mL / analysis | ~ 1.4 L | Plastic – 2.5 L | |
| Colour solution (Total Cu) | ~ 1.0 mL / analysis | ~ 2.7 L | Plastic – 5 L | |
| Colour solution (Cu) | ~ 1.0 mL / analysis | ~ 2.7 L | Plastic – 5 L | |
| REF1 solution (Total Cu + Cu) | ~ 1 L / calibration | 1 | Plastic – 1 L | |
| REF2 solution (Total Cu + Cu) | ~ 1 L / calibration | 1 | Plastic – 1 L | |

4.2 DI-water overview and consumption

| | Rinse water (mL/analysis) Type I | | | n water sis) Type I | Total (mL/analysis) | Consumption/28 days A rata 1 analysis/30 min |
|---|----------------------------------|---------|----------|------------------------|------------------------|---|
| | Total Cu | Cu (II) | Total Cu | Cu (II) | Total Cu + Cu (II) | Total Cu + Cu (II) |
| Α | N.A. | N.A. | N.A. | N.A. | N.A. | N.A. |
| В | N.A. | N.A. | N.A. | N.A. | N.A. | N.A. |
| С | N.A. | N.A. | N.A. | N.A. | N.A. | N.A. |
| 0 | N.A. | N.A. | N.A. | N.A. | N.A. | N.A. |
| 1 | 55 mL | 55 mL | 20 mL | 10 mL | 140 mL | 190 L |
| 3 | 55 mL | 55 mL | 20 mL | 10 mL | 140 mL | 190 L |

Remark

The indicated volumes are an estimation of the consumption for rinse and dilution water, based on a standard operating procedure, as defined in the specifications of the EZ analyser. Please be aware that, depending on the sample matrix, the rinse water volumes might increase.

4.3 Storage and quality of chemicals

Quality of chemicals

All chemicals should be of Reagent grade, ACS grade or better (*). The use of pro analysis chemicals is recommended. Poor quality of the reagents can affect the analyser performance.

(*) Analytical Reagent (AR), Guaranteed Reagent (GR), UNIVAR, AnalaR, Premium Reagent (PR), ReagentCertified ACS reagent, ACS Plus reagent, puriss p.a. ACS reagent, ReagentPlus®, TraceCERT®, Suprapur®, Ultrapur®, or better are also possible.

Quality of DI-water

All EZ analysers are tested with standard solutions, reagents and dilution water prepared using type I water or better as defined by ASTM D1193-91.

To achieve the specifications as stated on the data sheet, method and reagents sheet and acceptance test reports, the same water quality (or better) must be used for the preparation of the standard solutions, reagents and dilution water.

Additionally the water used for the preparation of the standard solutions for an EZ analyser must be free of the parameter or any of the interferences for the method of that EZ analyser.

Storage of Reagents

While operating the instrument, keep in mind the reagent requirements as stated in the reagent overview, the chapters below and/or in the data sheet of the instrument.

A CAUTION



For longer-term storage: Store the reagents cold; Store the reagents in the dark;

If applicable: Store the reagents in a fridge during operation

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Refresh the reagents after one month (unless stated differently in the chapters below).

Do not mix old reagents with freshly prepared reagents. Remove old reagents from the container before adding freshly prepared reagents.

4.4 Acid solution (1 M)

| Products | Formula | MW (g/mol) | CAS No. | 1 litre solution |
|-------------------|------------------|------------|-----------|------------------|
| Nitric acid (65%) | HNO ₃ | 63.01 | 7697-37-2 | 69 mL |

Preparation

Take 69 mL nitric acid (HNO₃ 65%) and dilute to 1 litre with de-ionized water.

4.5 Acid solution (0.25 M)

| Products | Formula | MW (g/mol) | CAS No. | 1 litre solution |
|-------------------|------------------|------------|-----------|------------------|
| Nitric acid (65%) | HNO ₃ | 63.01 | 7697-37-2 | 17 mL |

Preparation

Take 17 mL nitric acid (HNO₃ 65%) and dilute to 1 litre with de-ionized water.

4.6 Colour solution

| Products | Formula | MW (g/mol) | CAS No. | 1 litre solution |
|--|--|------------|-------------|------------------|
| 2,2-Biquinoline-4,4'- dicarbolxylic acid dipotassium salt trihydrate | C ₂₀ H ₁₀ K ₂ N ₂ O ₄ * 3H ₂ O | 474.55 | 207124-63-8 | 3 g |
| Ammonium hydroxide solution (25%)* | NH ₄ OH | 35.05 | 1336-21-6 | 50 mL |
| Hydroxylamine hydrochloride | H ₄ CINO | 69.49 | 5470-11-1 | 1 g |

^{*} Density: 0.91 g/ml (20°C)

Preparation

Dissolve 3.0 g 2,2'-biquinoline-4,4'-dicarboxylic acid dipotassium salt trihydrate ($C_{20}H_{10}K_2N_2O_4*3H_2O$) in 100 ml de-ionized water. Add 50 mL ammonium hydroxide solution (NH₄OH, 25%) and dissolve completely.

Dissolve 1.0 g hydroxylamine hydrochloride in 100 mL de-ionized water.

Mix both solutions together in a volumetric flask of 1 litre and fill up to the mark grade with de-ionized water.

4.7 Calibration solution

| Products | Formula | MW (g/mol) | CAS No. | 1 litre solution |
|----------------------------------|--|------------|-----------|------------------|
| Copper (II) sulfate pentahydrate | CuO ₄ S * 5H ₂ O | 249.69 | 7758-99-8 | 3.9293 g |
| Nitric acid (65%) | HNO ₃ | 63.01 | 7697-37-2 | 1 mL |

Preparation

1000 mg/L Cu stock solution

Prepare a stock solution of 1000 mg/L Cu: Dissolve accurately 3.9293 g copper(II)sulfate pentahydrate (CuO_4S*5H_2O) in 500 mL de-ionized water using a volumetric flask of 1000 mL. Add 1 mL of concentrated nitric acid (HNO_3 65%). This addition is done to keep the solution stable. Fill up to 1 litre with de-ionized water.

Cu standard solution - REF2

Prepare a standard solution for calibration according to the following table: take accurately x mL of the 1000 mg/L Cu stock solution and transfer into a volumetric flask of 1000 mL. Add de-ionized water up to the mark grade.

| | Measuring range | Concentration REF2 | Amount of stock solution to add to 1 litre |
|---|-----------------|--------------------|--|
| Α | 0.3 mg/L Cu | 0.3 mg/L Cu | 0.30 mL |
| В | 0.75 mg/L Cu | 0.75 mg/L Cu | 0.75 mL |
| С | 1.5 mg/L Cu | 1.5 mg/L Cu | 1.50 mL |
| 0 | 3.0 mg/L Cu | 3.0 mg/L Cu | 3.0 mL |
| 1 | 12 mg/L Cu | 12 mg/L Cu | 12 mL |
| 3 | 30 mg/L Cu | 30 mg/L Cu | 30 mL |

Cu standard solution - REF1

Prepare a standard solution of 0 mg/L Cu. Use de-ionized water.

4.8 Cleaning solution (facultative)

The cleaning procedure should prevent any build-up of chemicals in the analyser. To obtain an effective cleaning procedure one has to test the cleaning solution and the cleaning interval for each application. Perform the selected cleaning solution and interval for a trial period, check then the effectiveness of the procedure and change if necessary.

| | Change Information | | | | |
|--|--|--|--|--|--|
| Date: 25/05/2022 | Previous version: Edition 1.01 to Edition 1.02 | | | | |
| | | | | | |
| | Reason for Change | | | | |
| - Correction of | - Correction of CAS number of Nitric Acid | | | | |
| Description of Change | | | | | |
| - Correction of CAS number of Nitric Acid from 7697-32-2 to 7697-37-2 (chapter 4.4, 4.5 and 4.7) | | | | | |