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Organo-mineral fertilizers - Identification of chelating agents - Part 1: Determination of EDTA, HEEDTA and DTPA by ion chromatography

Organisch-mineralische Düngemittel - Identifizierung von Chelatbildnern - Teil 1: Bestimmung von EDTA, HEEDTA und DTPA mittels Ionenchromatographie

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English Version

**Organo-mineral fertilizers - Identification of chelating
agents - Part 1: Determination of EDTA, HEEDTA and
DTPA by ion chromatography**

Organisch-mineralische Düngemittel - Identifizierung
von Chelatbildnern - Teil 1: Bestimmung von EDTA,
HEEDTA und DTPA mittels Ionenchromatographie

This draft Technical Specification is submitted to CEN members for Vote. It has been drawn up by the Technical Committee CEN/TC 260.

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

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European foreword

This document (FprCEN/TS 17789-1:2021) has been prepared by Technical Committee CEN/TC 260 "Fertilizers and liming materials", the secretariat of which is held by DIN.

This document is currently submitted to the Vote on TS.

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Introduction

Micronutrients are considered to be, in plant nutrition, a number of elements known to be needed in small amounts for proper plant growth and development. The most common are Iron (Fe), Manganese (Mn), Molybdenum (Mo), Copper (Cu), Zinc (Zn) and Boron (B).

If an organo-mineral fertilizer contains a substance, or one of the substances in the mixture, which is intended to enhance the long term availability to plants of micronutrients in the EU fertilizing product, that substance is either a chelating agent or a complexing agent.

The chelating agents are divided into two groups¹:

- Group 1: EDTA, DTPA, HEEDTA, IDHA and [S,S]-EDDS;
- Group 2: Chelating agents present in UVCB (unknown or variable composition, complex reaction products or biological materials) chelates including [o,o] EDDHA, [o,p] EDDHA, [o,o] EDDHMA, HBED and EDDHSA.

This document specifies the test method to be used in order to determine the content of certain chelating agents (EDTA, DTPA and HEEDTA) in organo-mineral fertilizers (product function category (PFC) 1 (B) according to Regulation (EU) 2019/1009 [4]).

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¹ Abbreviated terms are described in Annex A.

1 Scope

This document specifies a method for the determination by ion chromatography of the total amount of each of the individual chelating agents EDTA, HEEDTA, and DTPA in organo-mineral fertilizers, having an organic matrix based on vegetal residues (cocoa shells, grape residue, soybean residue, ...), algae extract, and animal meal (feather, bones, blood, ...) and containing one or more of these substances.

The method allows the identification and the determination of the total water-soluble fraction of each of these chelating agents. It does not allow to distinguish between the free form and the metal bound form of the chelating agents.

This method applies to organo-mineral fertilizers containing chelates of one or more of the following micronutrients: cobalt, copper, iron, manganese, zinc and with a mass fraction of at least 0,1 %.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

EN 12944-1, *Fertilizers and liming materials and soil improvers - Vocabulary - Part 1: General terms*

EN 12944-2, *Fertilizers and liming materials and soil improvers - Vocabulary - Part 2: Terms relating to fertilizers*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in EN 12944-1 and EN 12944-2 apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at <https://www.iso.org/obp>
- IEC Electropedia: available at <https://www.electropedia.org/>

4 Principle

The micronutrients associated with the chelating agents present in an aqueous extract of the sample are replaced by iron(III). The iron chelates are separated and determined by ion chromatography. The separation is based on anion exchange, by elution with a nitrate acetate solution. The detection is based on UV photometry at 330 nm, after post-column reaction with diluted perchloric acid.

5 Interferences

Several substances can interfere, to a degree largely dependent on the type of column used. With the column described in 7.2, the following phenomena have been observed.

- a) Injection of solutions having high concentrations of salts can cause shifts in the retention times, mostly decreasing the retention when compared to the standard solutions. In these cases, the identity of the peaks can be confirmed by standard addition.
- b) Solutions having high concentrations of salts can also create a large signal at the void volume, poorly resolved from the HEEDTA peak.

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- c) High concentrations of nitrate, carbonate, sulfate, and phosphate do not interfere. However, large amounts of chloride create a negative fronting peak poorly resolved from the DTPA peak and altering its peak shape.
- d) Compounds, related to the group of polyamino polycarboxylic acids can interfere. While signals for [o,o] EDDHA, [o,o] EDDHMA, and EDDHSA are not detected, relatively weak signals are observed for NTA and CDTA. Under some conditions NTA can coelute with HEEDTA or EDTA.
- e) No signals have been detected for the following complexing agents: citrate, oxalate, tartrate, phthalate, and 20 naturally occurring amino acids.

6 Reagents

All reagents shall be of recognized analytical grade.

6.1 Water

All water used should conform to EN ISO 3696 and be degassed.

6.2 Nitric acid, substance concentration $c(\text{HNO}_3) = 7,2 \text{ mol/l}$.

Nitric acid, diluted 1 + 1 with water.

6.3 Sodium hydroxide solution, $c(\text{NaOH}) = 0,5 \text{ mol/l}$.

Dissolve 20 g of NaOH in pellet form in a 1 000 ml volumetric flask with water. Dilute to the mark and homogenize.

6.4 EDTA stock solution, $c(\text{EDTA}) = 2 \text{ mmol/l}$.

Dissolve 372 mg of the sodium dihydrogen salt of ethylene diamine tetraacetic acid dihydrate in 400 ml of water in a 500 ml volumetric flask. After dissolution, dilute to the mark with water and homogenize. Store the solution in a plastic bottle.

6.5 HEEDTA stock solution, $c(\text{HEEDTA}) = 2 \text{ mmol/l}$.

Dissolve 380 mg of the trisodium salt of hydroxyethyl ethylene diamine triacetic acid dihydrate in 400 ml of water in a 500 ml volumetric flask. After dissolution, dilute to the mark with water and homogenize. Store the solution in a plastic bottle.

6.6 DTPA stock solution, $c(\text{DTPA}) = 2 \text{ mmol/l}$.

Dissolve 393 mg of diethylene triamine pentaacetic acid in 10 ml of NaOH (6.3) in a 50 ml beaker. After dissolution, transfer quantitatively into a 500 ml volumetric flask, dilute to the mark with water and homogenize. Store the solution in a plastic bottle.

6.7 EDTA/HEEDTA/DTPA standard mixtures.

Into a set of three volumetric flasks of 100 ml, pipette respectively 2,5 ml, 5 ml and 10 ml of each stock solution of 2 mmol/l (6.4, 6.5 and 6.6). Dilute to the mark with water and homogenize. These solutions contain a mixture of EDTA, HEEDTA and DTPA, in concentrations of respectively 0,05 mmol/l, 0,1 mmol/l and 0,2 mmol/l. They should be used within two days.

6.8 Eluent nitrate (50 mmol/l) and acetate (50 mmol/l).

Dissolve 4,10 g of anhydrous sodium acetate (NaCH_3COO) in a mixture of 800 ml of water and 6,95 ml of nitric acid (6.2) in a 1 l volumetric flask. Dilute to the mark with water and homogenize. Adjust the pH of the eluent to $2,75 \pm 0,20$. Before use, filter the solution through a $0,45 \mu\text{m}$ membrane filter (7.4).

NOTE The concentration of the eluent influences the speed and the efficiency of the separation, which can be carried out with eluent concentrations varying between 25 mmol/l and 75 mmol/l of nitrate and acetate. At low concentrations, an improved separation between the void signal, the HEEDTA and the EDTA peaks can be obtained, while at high concentrations a better DTPA peak shape is observed.

6.9 Iron(III) nitrate solution.

Dissolve 5 g of ferric nitrate nonahydrate ($\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$) in a mixture of 800 ml of water and 21 ml of nitric acid (6.2) in a 1 000 ml volumetric flask. Dilute to the mark with water and homogenize. Store the solution in a plastic bottle.

NOTE Nitric acid is added in order to stabilize the solution and to allow the complete replacement of other chelated micronutrients by iron(III).

6.10 Perchloric acid solution, 2 %.

Dilute 29 ml of perchloric acid (70 % HClO_4 , $\rho = 1,67 \text{ g/ml}$) to 1 l with water.

7 Apparatus

Usual laboratory equipment, glassware, and the following:

7.1 **Rotary shaker**, capable of operating at a rotational speed of about 35 min^{-1} to 45 min^{-1} .

7.2 **Chromatograph**, equipped with:
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a) an isocratic pump delivering the eluent (6.8) at a flow rate of $0,5 \text{ ml/min}$;

b) an injection valve with an injection loop of about $50 \mu\text{l}$;

c) an anion exchange separator column $10 \mu\text{m}$ particles – 2 % substrate x – linking 100 mmol_c capacity (per column) – alkyl quaternary ammonium – medium high (hydrophobic)²;

NOTE 1 The column mentioned in c) is not exclusive. Any anion exchange column with comparable capacity, hydrophobicity, and selectivity can probably be used.

NOTE 2 Columns can, in their native state, exhibit some sensitivity towards various anions (e.g. nitrate, sulfate, phosphate) which can disappear after treatment with excess iron. It can be necessary to precondition the column prior to its use by repeated injections, at least 10, of the iron(III) solution (6.9), followed by equilibration under eluent flow for at least 6 h.

After many injections, especially of products having complex matrices, the column can lose some of the capacity and the separating efficiency. Substances like phenolic compounds (e.g. [o,o] EDDHA, humic acids) can be strongly adsorbed on the column particles. Suspended and colloidal matter can block the column entrance and disturb the eluent flow. The column manufacturer cleaning instructions shall be consulted for the suitable cleaning procedure.

² A combination of IonPac™ AS 7 separator and Ion Pac™ AG 7 guard column from Dionex Co, Sunnyvale, CA USA, or equivalent are examples of suitable products available commercially. This information is given for the convenience of users of this document and does not constitute an endorsement by CEN of these products.

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The injection of un-dissolved matter severely decreases the lifetime of the column. Solutions should always be allowed to equilibrate, and then filter through a 0,2 µm membrane filter before injection.

- d) a post-column reagent delivery module, delivering the reagent (6.10) at a flow rate between 0,5 ml/min and 0,6 ml/min;

NOTE 3 The post-column reagent stabilizes the eluting ferric chelates and suppresses the signal of some possibly interfering substances. For the analysis of samples with simple matrices, the post column reaction can be considered as superfluous.

- e) an UV/VIS detector with a 330 nm filter, output range set at an absorbance of 0,1;

NOTE 4 Ferric chelates of EDTA, HEEDTA, and DTPA have a useful spectral absorbance between 250 nm and 350 nm. The absorbance at 254 nm offers a better sensitivity, but also produces a noisier background, more sensitive to interferences.

- f) an integrator.

7.3 Balance, capable of weighing to the nearest 0,1 mg.

7.4 Membrane filters, micro membrane filters, resistant to aqueous solutions, with porosity of respectively 0,45 µm and 0,2 µm.

8 Sampling and sample preparation

Sampling and sample preparation are not part of the method specified in this document. Recommended sampling methods are given in EN 1482-1 and, for sample preparation, in EN 1482-2.

For the size reduction of samples with a high amount of chelating agents, it is not recommended to use a high speed laboratory mill. It is more convenient to grind the sample to a particle size less than 1 mm with a mortar.

9 Procedure

9.1 Preparation of the test solution

Weigh 5 g of the sample (m) to the nearest 1 mg, into a 250 ml volumetric flask (V_0). Add 200 ml of water and put the flask on the rotary shaker (7.1) for 1 h. Make up to the mark with water, homogenize, and filter through a paper filter. If necessary, dilute the filtrate with water, in order to obtain a concentration of chelating agent between 0,02 mmol/l and 0,2 mmol/l. Let D be the dilution factor. Pipette 20 ml of the solution into a 100 ml beaker. Add 5 ml of the iron(III) solution (6.9), homogenize, and allow to stand for 15 min.

The addition of iron can cause precipitation, especially if phosphate is present in the sample solution. For this reason, the precipitate should be allowed to settle and the solution to equilibrate for 15 min.

9.2 Preparation of the standard solutions

Pipette 20 ml of each of the EDTA/HEEDTA/DTPA standard mixtures of respectively 0,05 mmol/l, 0,1 mmol/l and 0,2 mmol/l (6.7) into a set of 100 ml beakers. Add 5 ml of the iron(III) solution (6.9), homogenize, and allow to stand for 15 min.