



SLOVENSKI STANDARD

SIST EN 14666:2005

01-november-2005

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Surface active Agents - Determination of tetraacetyl ethylene diamine (TAED) content in TAED granules - Gas chromatographic method

Grenzflächenaktive Stoffe - Bestimmung des Tetraacetylenethylen-diamin (TAED)-Gehalts in TAED-Granulaten - Gaschromatographisches Verfahren

Agents de surface - Détermination de la teneur en tétra-acétyl-éthylène diamine (TAED) dans les granulés de TAED - Méthode par chromatographie en phase gazeuse

Ta slovenski standard je istoveten z: EN 14666:2005

ICS:

71.100.40 Površinsko aktivna sredstva Surface active agents

SIST EN 14666:2005

en

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EUROPEAN STANDARD

EN 14666

NORME EUROPÉENNE

EUROPÄISCHE NORM

June 2005

ICS 71.100.40

English version

Surface active Agents - Determination of tetraacetyl ethylene diamine (TAED) content in TAED granules - Gas chromatographic method

Agents de surface - Détermination de la teneur en tétra-acétyl-éthylène diamine (TAED) dans les granulés de TAED - Méthode par chromatographie en phase gazeuse

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This European Standard was approved by CEN on 19 May 2005.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the Central Secretariat or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the Central Secretariat has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

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

Management Centre: rue de Stassart, 36 B-1050 Brussels

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Foreword

This document (EN 14666:2005) has been prepared by Technical Committee CEN/TC 276 “Surface active agents”, the secretariat of which is held by AFNOR.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by December 2005, and conflicting national standards shall be withdrawn at the latest by December 2005.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

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5.3 Syringe, capacity 5 µl.

5.4 Filter, porosity 0,45 µm.

6 Sampling and preparation of the sample

The laboratory sample shall be prepared and stored in accordance with ISO 607. Granules shall be ground for 1 min in an analytic mill or in a mortar to reach homogeneity.

7 Procedure

7.1 Calibration

Weigh into sample vials 25 mg, 50 mg, 75 mg and 100 mg of TAED (4.3) to the nearest 0,1 mg.

NOTE It is not required to perform the calibration before each series of samples. However a standard of 50 mg of TAED in 20 ml of the internal standard solution (4.4) should be analysed before each measurement. The value of response factor should not differ by more than 0,5 % from the average value of the calibration.

Add 20 ml of the internal standard solution (4.4) and dissolve the samples.

Inject each calibration solution three times into the gas chromatograph (5.1). An example of a calibration chromatogram is given in Figure A.1.

Record the peak areas of TAED and internal standard for each chromatogram and carry out a linear regression analysis of (A_{IS}/A_I) versus (m_{IS}/m_I). The response factor of TAED referring to the internal standard is the slope of the calculated calibration curve.

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If a correlation factor of less than 0,98 is obtained prepare new calibration solution and re-inject.

If a data acquisition system is not available, calculate the response factor RF_i for each chromatogram according to the following formula:

$$RF_i = \frac{A_{IS} \times m_I}{A_I \times m_{IS}} \quad (1)$$

where

A_I the peak area of TAED in the chromatogram i of calibration solution;

A_{IS} the peak area of the stearic acid methyl ester (4.2) in the chromatogram i of calibration solution;

m_I the mass of TAED in the calibration solution of the chromatogram i , in milligrams;

m_{IS} the mass of the stearic acid methyl ester in the calibration solution of the chromatogram i , in milligrams.

The response factor RF of TAED referring to the internal standard is the mean value of all RF_i . If the relative standard deviation of RF is higher than 3 % repeat the calibration procedure and re inject.

7.2 Determination

Immediately after grinding or crushing the sample (see Clause 6), weigh approximately 500 mg of the sample (m) to the nearest 0,1 mg into a 100 ml volumetric flask. Add about 250 mg stearic acid methyl ester (4.2) to the nearest 0,1 mg and make up to the mark with acetonitrile (4.1).

Insert a stirring rod into the flask and stir for 20 min on a magnetic stirrer at room temperature.

Inject the clear solution directly into the gas chromatograph (5.1). If the solution is cloudy, filter it through the filter (5.4) and only then inject it into the gas chromatograph.

Set up the gas chromatograph to give results similar to the example chromatogram in Figure A.1.

8 Calculation and expression of results

The content of TAED, $w(\text{C}_{10}\text{H}_{16}\text{O}_4\text{N}_2)$, in the sample, expressed in grams per 100 g, is calculated by the equation (2):

$$w(\text{C}_{10}\text{H}_{16}\text{O}_4\text{N}_2) = \frac{RF \times m_{\text{IS}} \times A_{\text{S}} \times 100}{m \times A_{\text{IS}}} \quad (2)$$

where

RF the response factor of TAED relating to the stearic acid methyl ester (4.2), (see 7.1);

m_{IS} the mass of the stearic acid methyl ester, in milligrams (see 7.2);

m the mass of the sample, in milligrams (see 7.2);

A_{IS} the peak area of the stearic acid methyl ester in the chromatogram of the sample solution;

A_{S} the peak area of TAED in the chromatogram of the sample solution.

9 Precision

9.1 Repeatability limit

The absolute difference between two independent single test results, obtained using the same method on identical test material in the same laboratory by the same operator using the same equipment within a short interval of time, will not exceed the repeatability limit, r , in more than 5 % of cases.

Precision data are given in Annex B.

9.2 Reproducibility limit

The absolute difference between two independent single test results, obtained using the same method on identical test material in different laboratories by different operators using different equipment, will not exceed the reproducibility limit, R , in more than 5 % of the cases.

Precision data are given in Annex B.

10 Test report

The test report shall include the following information:

- a) all information necessary for the complete identification of the sample;
- b) method used (a reference to this document);
- c) test results;
- d) details of any operation not specified in this document or in the European Standards to which reference is made, and any operations regarded as optional, as well as any incidents like to have affected the results.

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