

SLOVENSKI STANDARD oSIST ISO/FDIS 15819:2008

01-junij-2008

?cnaYhj_U'!'5 bU']hg_Y'aYhcXY'!'B]hfcnUa]b].'XYhY_V]/U']b'Xc`cYjUb/Y'B! b]hfcncX]YhUbc`Ua]bU'fB89@5'L'j '_cnaYh] b]\ 'gfYXghj]\ 'g'hY_c]bg_c _fcaUhc[fU2]/c'j]gc_Y``c`'fjcgh]'n'aUgbc'gdY_hfcaYhf]/g_c'XYhY_V]/c

Cosmetics - Analytical methods - Nitrosamines: Detection and determination of Nnitrosodiethanolamine (NDELA) in cosmetics by HPLC-MS

Cosmétiques - Méthodes analytiques - Nitrosamines: Recherche et dosage des Nnitrosodiéthanolamines (NDELA) dans les produits cosmétiques par CLHP-SM

Ta slovenski standard je istoveten z:

ICS:

71.100.70 S[:{^cāǎædeŽ√[æ‡^c}ã]¦ā][{[∖ã

Cosmetics. Toiletries

oSIST ISO/FDIS 15819:2008

en

FINAL DRAFT

INTERNATIONAL STANDARD

ISO/FDIS 15819

ISO/TC 217

Secretariat: ISIRI

Voting begins on: 2008-02-25

Voting terminates on: 2008-04-25

Cosmetics — Analytical methods — Nitrosamines: Detection and determination of N-nitrosodiethanolamine (NDELA) in cosmetics by HPLC-MS

Cosmétiques — Méthodes analytiques — Nitrosamines: Recherche et dosage des N-nitrosodiéthanolamines (NDELA) dans les produits cosmétiques par CLHP-SM

RECIPIENTS OF THIS DRAFT ARE INVITED TO SUBMIT, WITH THEIR COMMENTS, NOTIFICATION OF ANY RELEVANT PATENT RIGHTS OF WHICH THEY ARE AWARE AND TO PROVIDE SUPPORT-ING DOCUMENTATION.

IN ADDITION TO THEIR EVALUATION AS BEING ACCEPTABLE FOR INDUSTRIAL, TECHNO-LOGICAL, COMMERCIAL AND USER PURPOSES, DRAFT INTERNATIONAL STANDARDS MAY ON OCCASION HAVE TO BE CONSIDERED IN THE LIGHT OF THEIR POTENTIAL TO BECOME STAN-DARDS TO WHICH REFERENCE MAY BE MADE IN NATIONAL REGULATIONS.



Reference number ISO/FDIS 15819:2008(E)

PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.

Copyright notice

This ISO document is a Draft International Standard and is copyright-protected by ISO. Except as permitted under the applicable laws of the user's country, neither this ISO draft nor any extract from it may be reproduced, stored in a retrieval system or transmitted in any form or by any means, electronic, photocopying, recording or otherwise, without prior written permission being secured.

Requests for permission to reproduce should be addressed to either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office Case postale 56 • CH-1211 Geneva 20 Tel. + 41 22 749 01 11 Fax + 41 22 749 09 47 E-mail copyright@iso.org Web www.iso.org

Reproduction may be subject to royalty payments or a licensing agreement.

Violators may be prosecuted.

Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 15819 was prepared by Technical Committee ISO/TC 217, Cosmetics.

Introduction

Human exposure to N-nitrosamines can occur through diverse sources such as environment, food or personal care products. As a result of their perceived carcinogenic potential on several animal species, minimization of exposure to N-nitrosamines is recognised as important to the preservation of human health. Among N-nitrosamines, N-nitrosodiethanolamine (NDELA) has been recognised as a potential contaminant of cosmetics.

In this context, several analytical methods have been developed to detect and determine its presence in cosmetics – such as gas chromatography/thermal energy analysis, high performance liquid chromatography (HPLC) coupled either with photolysis and colorimetric quantification or with mass spectrometry (MS) determination. This latter method uses advanced technology to ensure the maximum specificity towards NDELA, to minimize the risk of artifactual formation of the analyte of interest and to allow precise quantification.

This analytical method uses high performance liquid chromatography coupled with mass spectrometry to separate and detect trace levels of NDELA from a cosmetic ingredient or product matrix with maximum specificity for NDELA.

Cosmetics — Analytical methods — Nitrosamines: Detection and determination of N-nitrosodiethanolamine (NDELA) in cosmetics by HPLC-MS

1 Scope

This International Standard describes a method for the detection and quantification of NDELA in cosmetics and raw materials used in cosmetics.

This method is not applicable to the detection and/or quantification of nitrosamines other than NDELA nor to the detection and/or quantification of NDELA in products other than cosmetics or raw materials used in cosmetics.

If a product has a possibility of either NDELA contamination from ingredients or NDELA formation by the composition of ingredients, the method shall be applied for quantitative determination of NDELA. Accordingly the method would not be applied in routine testing of cosmetic products. Because of the large variety of cosmetic products within this field of application, this method might need to be adapted for certain matrices.

Therefore, International Standards dedicated to alternative methods for testing nitrosamines in cosmetic products are being developed separately. Other methods can be employed provided that they are verified as to their detection of NDELA and validated in terms of recovery and quantification of the analyte.

2 Normative references

The following referenced documents are indispensable for the application 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.

ISO 3696:1987, Water for analytical laboratory use — Specification and test methods

3 Principle

Extraction of the nitrosamine NDELA in cosmetic samples is carried out with water in the presence of deuterated d8-NDELA used as internal standard. Clean-up is performed either using solid phase extraction (SPE clean-up, see 6.3.1) with a C18 cartridge or liquid–liquid extraction using dichloromethane (DCM clean-up, see 6.3.2) when the samples are not dispersible in water. The extracts are analysed by HPLC-MS-MS (high performance liquid chromatography coupled with tandem mass spectrometric detection).

NDELA quantification is done by comparing the ratio of the major fragment ions of NDELA and d8-NDELA with the calibration curve.

Confirmation of the presence of NDELA is carried out by using the molecular ion and two diagnostic ions.

4 Reagents

During the analysis, unless otherwise stated, use only reagents of recognized analytical grade and only distilled water or water of grade 1 in accordance with ISO 3696:1987. Solvent shall be of quality for HPLC analysis.

- 4.1 Methanol (MeOH), HPLC grade.
- 4.2 Ethanol (EtOH), HPLC grade.
- **4.3 Dichloromethane**, HPLC grade.
- 4.4 N-nitrosodiethanolamine, with known purity greater than 95 %.
- 4.5 d8-N-nitrosodiethanolamine, with known purity greater than 95 %.
- **4.6** Ammonium acetate (NH₄Ac), analytical grade.
- 4.7 1 mol/l ammonium acetate solution, formed by dissolving 77,08 g of NH₄Ac in 1,0 l water.

4.8 Eluent A: 2 mmol NH₄Ac in water, formed by taking 2 ml of 1 mol/l NH₄Ac (4.7) and making up to 1 l with water.

4.9 Eluent B: 2 mmol NH₄Ac in 90 % MeOH/water, formed by taking 2 ml of 1 mol/l NH₄Ac (4.7) and adding 900 ml MeOH and 98 ml water.

5 Apparatus

Use standard laboratory glassware and equipment, with the addition of:

5.1 Vortex mixer.

5.2 Sample processing station, in SPE application (such as Vacmaster^{®1)} sample processing station, IST).

5.3 Centrifuge, capable of reaching not less than 20 000 G.

5.4 Solid phase extraction columns, e.g. Bakerbond^{®1)} C18 – 6 ml, 500 mg reversed phase octadecylsilane bonded to silicagel, 40 APD, 60 Å.

5.5 HPLC-MS-MS equipment consisting of:

5.5.1 High performance liquid chromatography apparatus, consisting of an eluent reservoir, a pump, an injection system, a data processor, e.g. an integrator with plotter, coupled with tandem mass spectrometry using electrospray ionization.

¹⁾ Vacmaster[®], Bakerbond[®] and Spherisorb[®] are examples of suitable products available commercially. This information is given for the convenience of users of this International Standard and does not constitute an endorsement by ISO of these products.