

INTERNATIONAL
STANDARD

ISO
19070

IULTCS/IUC 34

First edition
2016-03-01

**Leather — Chemical determination
of *N*-methyl-2-pyrrolidone (NMP)
in leather**

*Cuir — Détermination de la teneur en N-méthyl-2-
pyrrolidone (NMP) dans le cuir*

iTeh STANDARD PREVIEW
(standards.iteh.ai)

[ISO 19070:2016](https://standards.iteh.ai/catalog/standards/sist/a0ff8-a9e1-48cf-b0c0-3f9c6d717107/iso-19070-2016)

<https://standards.iteh.ai/catalog/standards/sist/a0ff8-a9e1-48cf-b0c0-3f9c6d717107/iso-19070-2016>



Reference numbers
ISO 19070:2016(E)
IULTCS/IUC 34:2016(E)

© ISO 2016

iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 19070:2016

<https://standards.iteh.ai/catalog/standards/sist/a1a0ff8-a9e1-48cf-b0c0-39c6d717107/iso-19070-2016>



COPYRIGHT PROTECTED DOCUMENT

© ISO 2016, Published in Switzerland

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Ch. de Blandonnet 8 • CP 401
CH-1214 Vernier, Geneva, Switzerland
Tel. +41 22 749 01 11
Fax +41 22 749 09 47
copyright@iso.org
www.iso.org

Contents

	Page
Foreword.....	iv
Introduction.....	v
1 Scope.....	1
2 Normative references.....	1
3 Principle.....	1
4 Reagents.....	1
5 Apparatus.....	1
6 Procedure.....	2
6.1 Preparation of standard solutions.....	2
6.1.1 NMP stock solution (1 000 µg/ml).....	2
6.1.2 NMP Calibration solution.....	2
6.2 Sample preparation.....	2
6.3 Extraction procedure.....	2
7 Chromatographic determination.....	2
8 Quantification.....	3
9 Performance of the method.....	3
10 Test report.....	3
Annex A (informative) Chromatographic analysis.....	4

ISO 19070:2016
<https://standards.iteh.ai/catalog/standards/sist/a1a0ff8-a9e1-48cf-b0c0-3f9c6d717107/iso-19070-2016>

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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT) see the following URL: [Foreword - Supplementary information](#)

ISO 19070 was prepared by the Chemical Testing Commission of the International Union of Leather Technologists and Chemists Societies (IUC Commission, IULTCS) in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 289, *Leather*, the secretariat of which is held by UNI, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

IULTCS, originally formed in 1897, is a world-wide organization of professional leather societies to further the advancement of leather science and technology. IULTCS has three Commissions, which are responsible for establishing international methods for the sampling and testing of leather. ISO recognizes IULTCS as an international standardizing body for the preparation of test methods for leather.

Introduction

Under the REACH regulation in the European Union (EU Regulation 1907/2006), *N*-methyl-2-pyrrolidone has been included in the candidate list of substances of very high concern (SVHC) and its use is restricted.

Accordingly, a new method is given for the determination of *N*-methyl-2-pyrrolidone in leather.

iTeh STANDARD PREVIEW (standards.iteh.ai)

ISO 19070:2016

<https://standards.iteh.ai/catalog/standards/sist/a1a0ff8-a9e1-48cf-b0c0-39c6d717107/iso-19070-2016>

iTeh STANDARD PREVIEW
(standards.iteh.ai)

ISO 19070:2016

<https://standards.iteh.ai/catalog/standards/sist/a1a0ff8-a9e1-48cf-b0c0-39c6d717107/iso-19070-2016>

Leather — Chemical determination of *N*-methyl-2-pyrrolidone (NMP) in leather

1 Scope

This International Standard specifies a method to determine the amount of *N*-methyl-2-pyrrolidone (NMP) in leather and leather components.

This method may also be used for the determination of *N*-ethyl-2-pyrrolidone (NEP) in leather.

2 Normative references

The following documents, in whole or in part, are normatively referenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 2418, *Leather — Chemical, physical and mechanical and fastness tests — Sampling location*

ISO 4044, *Leather — Chemical tests — Preparation of chemical test samples*

iTeh STANDARD PREVIEW
(standards.iteh.ai)

3 Principle

The test sample is extracted using acetone at 50 °C in an ultrasonic bath for 1 h. An aliquot is then analysed using a gas chromatograph with a mass selective detector (GC-MS).

<https://standards.iteh.ai/catalog/standards/sist/a1a0ff8-a9e1-48cf-b0c0-3f9c6d717107/iso-19070-2016>

4 Reagents

All reagents shall be analytical grade.

4.1 Acetone, CAS¹⁾ No. 67-64-1.

4.2 *N*-methyl-2-pyrrolidone, at least 99,5 %, CAS No. 872-50-4.

5 Apparatus

Usual laboratory equipment and, in particular, the following.

5.1 Analytical balance, capable of reading to 0,000 1 g.

5.2 Sealable jar, approximately 20 ml, with lid (suitable for carrying out an acetone extraction at 50 °C).

5.3 Ultrasonic bath (temperature controlled).

5.4 Micro-pipettes, 50 µl and 100 µl.

5.5 Pipettes, 0,5 ml to 5 ml capacity.

5.6 Volumetric flasks, 100 ml.

1) CAS = Chemical Abstracts Service

5.7 **Gas chromatograph**, with mass selective detector (GC-MS).

5.8 **PTFE-membrane filter**, pore size 0,45 µm.

5.9 **Sample vials**, PTFE seal.

6 Procedure

6.1 Preparation of standard solutions

6.1.1 NMP stock solution (1 000 µg/ml)

Weigh accurately 0,1 g of *N*-methyl-2-pyrrolidone using the analytical balance (5.1), into a 100 ml volumetric flask (5.6) and fill up to the mark with acetone (4.1).

6.1.2 NMP Calibration solution

Four calibration solutions are used to establish the calibration curve. They are all prepared in 100 ml volumetric flasks, as specified in Table 1.

Table 1 — Preparation of calibration solutions

Calibration solution	Volume of the NMP stock solution (6.1.1)	Concentration of NMP in the calibration solution
1	0,1 ml	1 µg/ml
2	0,5 ml	5 µg/ml
3	1 ml	10 µg/ml
4	5 ml	50 µg/ml
Filled up to the mark with acetone		

6.2 Sample preparation

Take a representative sample from the leather or leather component. When possible sample according to ISO 2418.

The leather specimen shall be cut into pieces in accordance with ISO 4044. The dimensions of the pieces shall not be larger than 2 mm to 3 mm. Do not condition the sample.

6.3 Extraction procedure

Weigh accurately approximately 1 g of the leather pieces with the analytical balance (5.1) and place them into the sealable jar (5.2). Add 10 ml of acetone and extract the leather at (50 ± 2) °C in an ultrasonic bath for (1,0 ± 0,1) h.

After cooling to room temperature, filter the solution through a PTFE membrane filter (5.8) and transfer to a vial (5.9), and seal with a PTFE cap. This solution is now ready for instrumental analysis, which shall be performed within 24 h.

If there is to be any delay in analysing the prepared sample then it may be stored at (4 ± 2) °C for up to 24 h.

7 Chromatographic determination

The detection of the *N*-methyl-2-pyrrolidone may be performed using the GC-MS chromatographic technique given in Annex A.

If the amount of NMP in the sample is above the highest calibration point, a dilution step is necessary, which has to be considered in the calculation in [Clause 8](#).

8 Quantification

The content of *N*-methyl-2-pyrrolidone is calculated as a mass fraction, w , in mg/kg, according to Formula (1):

$$w = \frac{A_{\text{NMP-S}} \cdot C_{\text{NMP-Std}} \cdot V}{A_{\text{NMP-Std}} \cdot m_{\text{S}}} \quad (1)$$

where

- $A_{\text{NMP-S}}$ is the peak area of NMP in the sample;
- $A_{\text{NMP-Std}}$ is the peak area of NMP in the calibration standard;
- $C_{\text{NMP-Std}}$ is the concentration of NMP in the calibration standard, in $\mu\text{g/ml}$;
- V is the final volume of sample, in ml, ($V = 10$ ml according to [6.3](#));
- m_{S} is the mass of the sample as received, in g.

9 Performance of the method

The quantification limit shall be sufficient to ensure the analysis complies with any specification or legislative framework requirements, for example less than 1 000 mg/kg for *N*-methyl-2-pyrrolidone in the leather. The typical limit of quantification for this method is 5 mg/kg.

10 Test report

The test report shall include at least the following:

- a) a reference to this International Standard, i.e. ISO 19070;
- b) the date of the test;
- c) all details necessary for complete identification of the sample tested;
- d) the amount of *N*-methyl-2-pyrrolidone determined in the leather sample, in mg/kg;
- e) the limit of detection for this method;
- f) any deviation from the analytical procedure specified in this International Standard.