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ISO TC 35/SC 16/WG 1

Secretariat: DIN•

Date: 2024-12-18

Paints and varnishes — Determination of preservatives in water-dilutable coating materials — Part 1: Determination of in-can free formaldehyde

<u>Peintures et vernis — Dosage des agents de préservation dans les produits de peinture diluables à</u> <u>l'eau — Partie 1: Dosage du formaldéhyde libre en pot</u>

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2	Normative references	L		
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4	Principle	3		
5	Apparatus	5		
6	Reagents and materials	,		
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8	Precision1	5		
9	Test report1	,		
Anne	x A (normative) Titration methods for the determination of formaldehyde content	3		
<u>A.1</u>	- Determination of formaldehyde concentration: Iodometric method	3		
<u>A.2</u>	Determination of formaldehyde concentration: pH-value method1	3		
Anne	x B (informative) Example of HPLC conditions for method B	L		
Anne	x C (informative) - Example of HPLC conditions for method C22	2		
Anne	x D (informative) Results of the interlaboratory test	3		
Bibli	ography	5		
<u>Forev</u>	vord	,		
<u>1 ht</u>	<u>- Scope</u>	La		
2	Normative references	L		
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Annex B (informative) Example of HPLC conditions for method B	<u></u> 21
Annex C (informative) Example of HPLC conditions for method C	<u></u> 22
Annex D (informative) Results of the interlaboratory test	<u></u> 23
Bibliography	25

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This document was prepared by Technical Committee ISO/TC 35, Paints and varnishes, Subcommittee SC 16, Chemical analysis, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 139, Paints and varnishes, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

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Paints and varnishes — Determination of preservatives in water-	lieadei
Bart 1. Determination of in-can free formaldehyde	
Tart 1. Determination of m-can nee for maldenyue	
1 Scope	
This document specifies the apparatus and the analytical methodmethods for determining th concentration of in-can free formaldehyde in water-dilutable coating materials.	e
This document can also be applied <u>forto</u> polymer dispersions.	Formatted: Body Text, Tab stops: Not at 0.7 cm + 1.4
The determination method A for in-can free formaldehyde described in this document <u>can beis</u> only o limited suitability for pigmented systems, as the inherent coloration of the material can have an influence on the detection.	f 5.6 cm + 2.1 cm + 2.8 cm + 3.5 cm + 4.2 cm + 4.9 cm + 5.6 cm + 6.3 cm + 7 cm
2 Normative references iTeh Standards	
The following documents are referred to in the text in such a way that some or all of their conten 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.	^t ai)
ISO 1513, Paints and varnishes — Examination and preparation of test samples	
ISO 3696, Water for analytical laboratory use — Specification and test methods	
ISO 15528, Paints, varnishes and raw materials for paints and varnishes — Sampling —	
3 Terms and definitions	
For the purposes of this document, the following terms and definitions apply.	
ISO and IEC maintain terminology databases for use in standardization at the following addresses:	
 ISO Online browsing platform: available at <u>https://www.iso.org/obp</u>https://www.iso.org/obp 	
 IEC Electropedia: available at <u>https://www.electropedia.org/</u>https://www.electropedia.org/ 	
3.1 ready for use state of a product when it is mixed in accordance with the manufacturer's instructions in the correct proportions and thinned if required using the correct thinners so that it is ready for application by the approved method	t Formatted: Don't adjust space between Latin and Asian text, Don't adjust space between Asian text and numbers
[SOURCE: ISO 11890-2:2020, 3.7]	Formatted: Normal, Centered, Space After: 24 pt, Tab stops: 17.2 cm, Right

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3.2	Formatted: Don't keep with next
in-can free formaldehyde content	
concentration of formaldehyde which is available in the coating as an <i>in-can preservative</i> (3.9) as	Formatted: cite_sec
determined by- <u>i</u>	
<u>—</u> method A of this document, i.e. photometric detection after derivatisation<u>derivatization</u> with acetylacetone;	
<u>—</u> method B of this document, i.e. liquid chromatography detection after <u>derivatisationderivatization</u> with DNPH (2,4-dinitrophenyl-hydrazine); or	
method C of this document i.e. liquid chromatography detection with post-column derivatisationderivatization with acetylacetone; where water is used as an extraction solvent for all three methods	Formatted: List Continue 1, Bulleted + Level: 1 + Aligned at: 0.63 cm + Indent at: 1.27 cm, Adjust space between Latin and Asian text, Adjust space between Asian text and numbers
Note 1 to entry. In-can free formal dehyde content corresponds to the amount of formal dehyde in milligrams, based	
on 1-kg of <i>coating material</i> (3.4) or polymer dispersion, which is available unbound in the sample at the time of	Formatted: cite sec
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4 Principle

4.1 General

This document describes three methods for the determination of in-can free formaldehyde in coating matricesmaterials which have been validated to show comparable results for selected samples with respect to their comparability limits statistically for methods A and B and with a single comparison for method C.

4.2 Equilibria of in-can free formaldehyde in the coating matrix

When a formaldehyde depot substance is added to a coating material, equilibrium is established between the formaldehyde bound to the formaldehyde depot substance and the in-can free formaldehyde. This equilibrium depends on the pH value, the coating matrix, the formaldehyde depot substance and the temperature. Furthermore, in water-dilutable coating materials, there is also equilibrium between the incan free formaldehyde and the matrix itself. The in-can free formaldehyde can undergo a variety of equilibrium reactions, e.g. with water to form a hydrate or with other formaldehyde molecules to form trimers.

A distinction shall be made in a coating material between the <u>in-can</u> total formaldehyde content (free and bound formaldehyde) and the in-can free formaldehyde content. In this document, the in-can free formaldehyde content is understood to be the formaldehyde concentration determined by either-<u>:</u>

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- <u>method A</u>, i.e. <u>derivatisationderivatization</u> with acetylacetone and subsequent photometric detection and quantification;
- <u>— method B, i.e. derivatisationderivatization</u> with DNPH and subsequent detection and quantification with liquid chromatography-ultraviolet/visible light detector_(LC-UV/VIS) analysis; or
- ___method C, i.e. LC(liquid chromatography_(LC) separation followed by post-column derivatisationderivatization with acetylacetone and subsequent quantification with UV/VIS(ultraviolet/visible light detector_(UV/VIS) detection.

4.3 Equilibria of in-can free formaldehyde during extraction

In addition to the equilibria in the coating matrix mentioned in 4.2, during extraction, the equilibrium of in-can free formaldehyde is also affected by the extraction solvent.

Depending on the nature of the extraction solvent, the sample/solvent ratio, extraction time and pH value, the equilibrium of bound and free in-can formaldehyde can be shifted to either side. Therefore, it is crucial to select the appropriate extraction solvent and extraction conditions.

NOTE If a solvent other than water is used for extraction, the value for the in-can free formaldehyde obtained is no longer defined by clause 3.2. The definition in 3.2 applies exclusively to equilibrium in water.

4.4 Equilibria of in-can free formaldehyde during derivatisation derivatization

The in-can free formaldehyde in the extract (or after chromatographic separation) is derivatized by chemical reaction with a derivatizing agent. The reaction is also an equilibrium reaction. To ensure complete derivatisationderivatization, an excess of derivatisationderivatization agent is added.

4.5 Principle of Method A: DerivatisationDerivatization with Acetylacetoneacetylacetone (ACAC) combined with photometric detection

4.5.1 Derivatisation Derivatization with ACAC

In order to enable photometric detection of formaldehyde, <u>derivatisationderivatization</u> shall be carried out. <u>DerivatisationDerivatization</u> introduces a chromophore after reaction with formaldehyde, which has the property to absorb ultraviolet (UV) or visible (VIS) light.

The <u>derivatisationderivatization</u> is performed prior to the photometric analysis (pre-column <u>derivatisationderivatization</u>).

In-can free formaldehyde is derivatised to 3,5-diacetyl-1,4-dihydrolutidine by using ACAC and ammonium ions (Hantzsch reaction). The absorption maximum of the compound is at a wavelength of 412-nm.

4.5.2 Detection and quantification with a spectral photometer

A spectral photometer is applied to quantify the formaldehyde-ACAC reaction product, which, in turn, can be used to calculate the concentration of in-can free formaldehyde in the coating matrix.

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4.6 Principle of Method B: DerivatisationDerivatization with Dinitrophenylhydrazinedinitrophenylhydrazine (DNPH) combined with Liquidliquid chromatography (LC) separation and UV/VIS detection

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4.6.1 Derivatisation Derivatization with DNPH

The in-can free formaldehyde can be derivatized with DNPH leading to the formation of a compound that can be separated by liquid chromatography and detected with a UV/VIS-detector at 360-nm.

4.6.2 Separation, detection and quantification with LC-UV/VIS

Liquid chromatography is applied to separate the different aldehyde-DNPH reaction products. Subsequent UV/VIS detection is used to identify and quantify the concentration of the formaldehyde-DNPH reaction product, which enables the calculation of the in-can free formaldehyde content in the coating sample.

If <u>derivatisation</u><u>derivatization</u> with DNPH is used, the determination of in-can free formaldehyde content shall be performed by LC-UV/VIS analysis, since the reaction with DNPH is also sensitive to other aldehydes, which can interfere when using a determination with a spectral photometer.

4.7 Principle of Method C: Liquid chromatography (LC) separation followed by postcolumn derivatisation<u>derivatization</u> with ACAC and subsequent quantification with UV/VIS detection

4.7.1 Separation of free formaldehyde by liquid chromatography

Liquid chromatography is performed prior to derivatisation.derivatization. It is applied to separate free formaldehyde from all other components present in the water extract to avoid interferences in the detection.

4.7.2 Post-column derivatisationderivatization with acetylacetone (ACAC) and quantification with UV/VIS detection

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To enable detection of the separated formaldehyde, <u>derivatisationderivatization</u> is performed as described in 4.5.1, in this case by mixing ACAC to the column outlet. Quantification is done in-line with UV/VIS detection at the maximum absorption of the formaldehyde ACAC reaction product, i.e. 412-_nm.

5 Apparatus

5.1 General

The usual laboratory apparatus and, in particular, the following apparatus stated in 5.2 to 5.6 shall be used.

5.2 General apparatus

5.2.1 <u>5.2.1</u> Precision scale, capable of weighing to an accuracy of 0,000-<u>1 g</u>.

5.2.2 Suitable volumetric flasks for calibration, with ground-glass stoppers.

5.2.3 5.2.3 Suitable pipettes.

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