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Ambient air — Determination of numerical concentration of inorganic fibrous particles — Scanning electron microscopy method

Air ambient — Détermination de la concentration en nombre des particules inorganiques fibreuses — Méthode par microscopie électronique à balayage



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Contents

	Page
Foreword	v
Introduction	vi
1 Scope	1
2 Normative references	1
3 Terms and definitions	1
4 Abbreviated terms	4
5 Principle	4
6 Apparatus and materials	4
6.1 Air sampling.....	4
6.1.1 Sampling head.....	4
6.1.2 Sampling train.....	5
6.1.3 Sampling pump.....	5
6.1.4 Needle valve.....	6
6.1.5 Volumetric flowmeter (rotameter).....	6
6.1.6 Timer.....	6
6.1.7 Dry type gas meter (optional).....	6
6.1.8 Meteorological instruments (optional).....	6
6.1.9 Instruments for unattended sampling (optional).....	7
6.2 Preparation of filters.....	7
6.2.1 Vacuum evaporator.....	7
6.2.2 Plasma asher.....	8
6.3 Sample analysis.....	8
6.3.1 Scanning electron microscope (SEM).....	8
6.3.2 Energy-dispersive X-ray system.....	8
6.3.3 Stereo-microscope.....	9
6.3.4 Gold-coated capillary-pore polycarbonate filters.....	9
6.3.5 Backing filters.....	9
6.3.6 Disposable plastic field monitors (optional).....	9
6.3.7 Technically pure oxygen.....	9
6.3.8 Rubber connecting hoses.....	9
6.3.9 Filter containers.....	9
6.3.10 Routine electron microscopy tools and supplies.....	9
6.3.11 Sample for resolution adjustment.....	9
6.3.12 Sample for magnification calibration.....	10
7 Air sample collection and analysis	10
7.1 Measurement planning.....	10
7.2 Collection of air samples.....	10
7.3 SEM specimen preparation.....	13
7.4 Analysis in the scanning electron microscope.....	13
7.4.1 General instructions.....	13
7.4.2 Fibre-counting criteria.....	14
7.4.3 Fibre classification.....	19
7.4.4 Analysis using reference spectra and peak height ratios.....	26
7.4.5 Measurement of fibre dimensions.....	28
7.4.6 Recording of data on the fibre counting form.....	28
8 Calculation of results	28
8.1 Calculation of the mean fibre concentration.....	28
8.2 Calculation of the 95 % confidence interval.....	30
9 Performance characteristics	30
9.1 General.....	30
9.2 Measurement uncertainty.....	30

This is a preview of "ISO 14966:2019". [Click here to purchase the full version from the ANSI store.](#)

9.2.1	Systematic errors.....	30
9.2.2	Random errors.....	30
9.2.3	Errors due to sampling.....	31
9.2.4	Errors associated with the SEM examination.....	31
9.2.5	Total error of the measurement.....	31
9.2.6	Random errors due to fibre counting.....	32
9.3	Limit of detection.....	34
10	Test report.....	35
Annex A	(normative) Preparation of filters for air sampling.....	37
Annex B	(normative) Procedures for calibration and adjustment of the SEM.....	38
Annex C	(informative) Characteristics and chemical composition of inorganic fibres.....	40
Annex D	(informative) Poisson variability as a function of fibre density on sampling filter and area of filter analysed.....	45
Annex E	(informative) Combination of the results from multiple samples.....	47
Bibliography	48

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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 of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 146, *Air quality*, Subcommittee SC 3, *Ambient atmospheres*.

This second edition cancels and replaces the first edition (ISO 14966:2002), which has been technically revised. It also incorporates the corrected version ISO 14699:2002/Cor 1:2007. The main changes compared to the previous edition are as follows:

- Counting rules, changed to the recommended method (membrane filter method) of the WHO (World Health Organization);
- Analytical procedure (classification), using normalized peak height ratios in addition to the method of the previous edition;
- Rule for early termination of filter evaluation (counting and analysis). A formula is given to terminate the filter evaluation, if the calculated (asbestos) fibre concentration is above a set limit value for this fibre concentration.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

Introduction

This document describes a method for measurement of the numerical concentration of inorganic fibrous particles in ambient air using the scanning electron microscope. This document is based on VDI 3492^[1].

The method is also suitable for determining the numerical concentrations of inorganic fibres in the interior atmospheres of buildings, for example measurement of residual airborne fibre concentrations after the removal of asbestos-containing building materials.

Biological research has shown that the fibrogenic or carcinogenic effect of a fibre is related to its length, diameter and its resistance to dissolution in a biological environment. The point at which fibres are too short, too thick or of insufficient durability to produce a fibrogenic or carcinogenic effect is uncertain. Fibres with lengths greater than 10 µm and diameters of a few tenths of 1 µm, which also have durabilities such that they remain unchanged for many years in the body, are regarded as particularly carcinogenic. Based on current knowledge, fibres shorter than 5 µm are thought to have a lower carcinogenic potential^{[2]-[5]}.

For the purposes of this document, a fibre is defined as a particle which has a minimum length to width (aspect) ratio of 3:1. Fibres with lengths greater than 5 µm and widths extending from the lower limit of visibility up to 3 µm are counted. Fibres with diameters less than 3 µm are considered to be respirable. Since the method requires recording the lengths and widths of all fibres, the data can be re-evaluated if it is required to derive concentrations for fibres with a higher minimum aspect ratio^[6].

The range of concentration to be measured extends from that found in clean air environments, in which the mean value of a large number of individual measurements of asbestos fibre concentrations has been found to be generally lower than 100 fibres/m³ (fibres longer than 5 µm), up to higher exposure scenarios in which concentrations as much as two orders of magnitude higher have been found^{[4][6]}.

This method is used to measure the numerical concentration of inorganic fibres with widths smaller than 3 µm and lengths exceeding 5 µm up to a maximum of 100 µm. Using energy-dispersive X-ray analysis (EDXA), fibres are classified as fibres with compositions consistent with those of asbestos fibres, calcium sulfate fibres and other inorganic fibres.

Calcium sulfate fibres are separated from other inorganic fibres and are not included in the final result, because on the basis of current knowledge, they do not represent any health hazard. Nevertheless, the numerical concentration of calcium sulfate fibres should be determined, since a high concentration of these fibres can negatively bias the results for probable asbestos fibres, and in some circumstances the sample may have to be rejected^[7]. In addition, knowledge of the numerical concentration of calcium sulfate fibres is of importance in the interpretation of fibre concentrations in ambient atmospheres.

Detection and identification of fibres becomes progressively more uncertain as the fibre width is reduced below 0,2 µm. Identification of a fibre as a specific species is more confident if the source of emission is known or suspected, such as in a building for which bulk materials are available for analysis.

In order to facilitate the scanning electron microscope examination, organic particles collected on the filter are almost completely removed by a plasma ashing treatment.

Except in situations where fibre identification is difficult, there should be only minor differences between fibre counting results obtained by this method and those obtained using the procedures for determination of PCM-equivalent fibres in [Annex E](#) of the transmission electron microscopy method ISO 10312^[8].