
Notranji zrak - 6. del: Določevanje hlapnih organskih spojin (VVOC, VOC, SVOC) v notranjem zraku in zraku v preskusnih komorah z aktivnim vzorčenjem v cevkah z adsorpcijskim polnilom, termično desorpcijo in plinsko kromatografijo z MS ali MS-FID

Indoor air - Part 6: Determination of organic compounds (VVOC, VOC, SVOC) in indoor and test chamber air by active sampling on sorbent tubes, thermal desorption and gas chromatography using MS or MS FID

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Air intérieur - Partie 6: Dosage des composés organiques (COTV, COV, COSV) dans l'air intérieur et l'air de chambre d'essai par prélèvement actif sur tubes à sorbant, désorption thermique et chromatographie en phase gazeuse avec détection MS ou MS-FID

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Indoor air —

Part 6:

Determination of organic compounds (VOC, SVOC) in indoor and test chamber air by active sampling on sorbent tubes, thermal desorption and gas chromatography using MS or MS FID

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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 on 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 the following URL: www.iso.org/iso/foreword.html. (standards.iteh.ai)

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This second edition cancels and replaces the first edition (ISO 16000-6:2012), which has been technically revised.

The main changes compared to the previous edition are as follows:

- other sorbents than Tenax TA[®] are allowed to be used;
- descriptions of VVOC and SVOC measurements are included in the mandatory part of the document.

A list of all parts in the ISO 16000 series can be found on the ISO website.

Introduction

ISO 16000-1 establishes general requirements relating to the measurement of indoor air pollutants and the important conditions to be observed before or during the sampling of individual pollutants or groups of pollutants. Aspects of the determination (sampling and analysis) and the sampling strategy of specific pollutants or groups of pollutants are specified in the subsequent parts of ISO 16000 (see Foreword).

ISO 16000-5 (dealing with VOC sampling strategy) is a link between ISO 16000-1 (a generic standard establishing the principles) and this part of ISO 16000, which deals with sampling and analytical measurements.

ISO 16017 (see [Clause 2](#) and Reference [8]) and ISO 12219 [3]-[7] also focus on measuring vapour-phase organic chemicals in air.

This standard can be applied to measure vapour phase organic compounds in indoor environments that include buildings with varying designs and purposes and cabins for different modes of transport, as well as measurement in product emission test chambers. These measurements can be for a range of purposes as described in ISO 16000-1 and -5, therefore the requirement for the measurement may be well defined by the task descriptor or may be quite open. For example, the task may be to determine a specific list of target chemicals with a defined sampling time and sensitivity of measurement or it may be to investigate the cause of a reported and poorly understood indoor air quality problem. Depending upon the task of measurement the user of this standard must select the most appropriate sampling and analytical instrumentation and conditions. This standard provides that information in the normative part combined with informative guidance. [Figure 1](#) refers to the most critical parts of the standard with regard to selection of the most appropriate methodology for the task to be undertaken. Tenax TA® only or multisorbents can be used to capture ranges of vapour phase organic compounds. Multisorbents are used for wider ranges and may improve recovery of compounds.

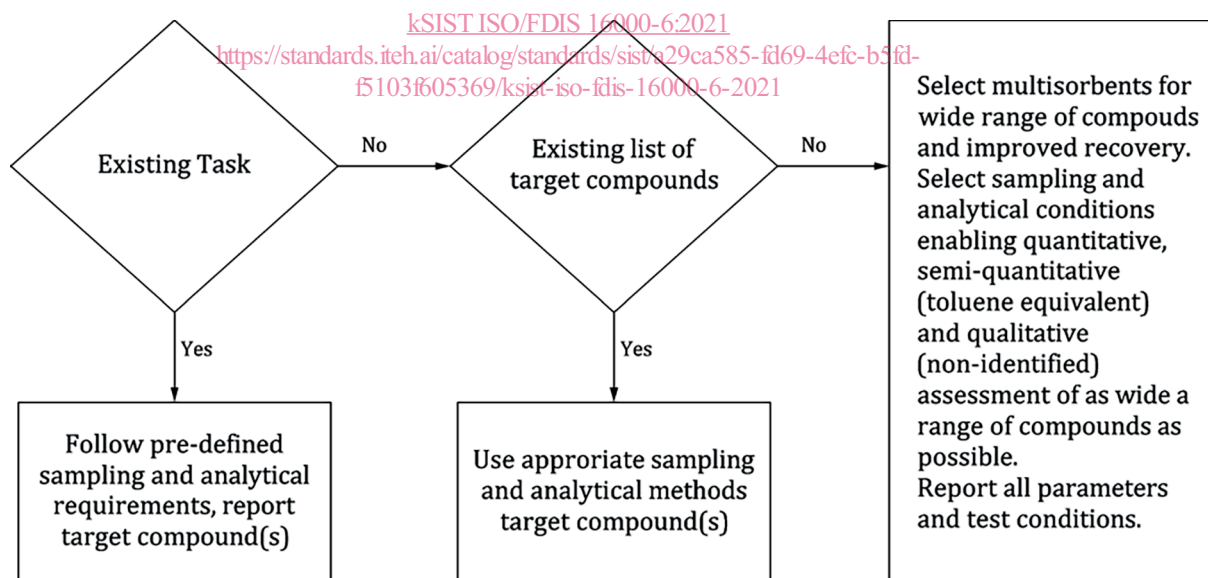


Figure 1 — Measurement scheme showing different ways of analysing air samples depending on the respective task including target compounds

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Indoor air —

Part 6:

Determination of organic compounds (VVOC, VOC, SVOC) in indoor and test chamber air by active sampling on sorbent tubes, thermal desorption and gas chromatography using MS or MS FID

1 Scope

This part of ISO 16000 specifies a method for determination of volatile organic compounds (VOCs) in indoor air and in air sampled for the determination of the emission from products or materials used in indoor environments (according to ISO 16000-1) using test chambers and test cells. The method uses sorbent sampling tubes with subsequent thermal desorption (TD) and gas chromatographic (GC) analysis^[13] employing a capillary column and a mass spectrometric (MS) detector with or without additional Flame Ionisation detector (FID).

The method is applicable to the measurement of most GC-compatible vapor-phase organic compounds at concentrations ranging from micrograms per cubic metre to several milligrams per cubic metre. Many very volatile organic compounds (VVOC) and semi-volatile organic compounds (SVOC) can be analysed depending on sorbents used.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content 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.

ISO 16000-1, *Indoor air — Part 1: General aspects of sampling strategy*

ISO 16017-1:2000, *Indoor, ambient and workplace air — Sampling and analysis of volatile organic compounds by sorbent tube/thermal desorption/capillary gas chromatography — Part 1: Pumped sampling*

EN 16516:2017, *Construction products: Assessment of release of dangerous substances. Determination of emissions into indoor air*

3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- IEC Electropedia: available at <http://www.electropedia.org/>
- ISO Online browsing platform: available at <https://www.iso.org/obp>

ISO/DIS 16000-6:2020(E)**3.1
semi-volatile organic compound
SVOC**

Organic compound eluting after n-hexadecane and up to and including n-C₃₀ on a gas chromatographic column specified as a 5% phenyl 95% methyl polysiloxane capillary column.

Note 1 to entry: The vapour-fraction of SVOC ranging in volatility from n-C₃₀ to n-C₄₄ can also be analysed by thermal desorption GC-MS but requires specific sampling and analytical conditions for optimum performance [22, 25]

**3.2
volatile organic compound
VOC**

Organic compound eluting between and including n-hexane and n-hexadecane on a gas chromatographic column specified as a 5% phenyl 95% methyl polysiloxane capillary column.

**3.3
very volatile organic compound
VVOC**

Organic compound eluting before n-hexane on a gas chromatographic column specified as 5% phenyl 95% methyl polysiloxane capillary column.

**3.4
total volatile organic compounds
TVOC**

Sum of the concentration of the identified and unidentified volatile organic compounds, as defined in [3.2](#) and calculated as detailed in [Annex A](#)

**3.5
total semi-volatile organic compounds
TSVOC**

Sum of the concentrations of identified and unidentified semi-volatile organic compounds as defined in [3.1](#) and calculated as detailed in [Annex A](#).

**3.6
target compound**

Individual vapour phase compound in indoor air with a concentration determined quantitatively and reported as a result of this method

**3.7
task list**

Specific list of requirements for sampling and analysis defined prior to testing and reflected in the reporting of the results

Note 1 to entry: The requirements may include a specific target list with or without associated limit criteria, and/or require investigations of unknowns. They may also include particular control of aspects such as the location, duration and frequency of sampling.

**3.8
laboratory blank**

Conditioned sampling tube from the batch selected for each sampling exercise, retained in the laboratory, sealed with long term storage caps throughout the sampling exercise to be used as a blank tube. These tubes are analysed with the sampled tubes

**3.9
field blank**

A conditioned sample tube from the batch used for the sampling exercise, subjected to the same handling procedure in the field as the sample tubes, including removal and replacement of storage caps, but not used for sample collection

3.10

internal standard

Compound of known concentration added to a sample to facilitate the qualitative identification and/or quantitative determination of the sample components

4 Symbols and abbreviated terms

For the purpose of this document, the following abbreviated terms apply:

VOC	volatile organic compounds
VVOC	very volatile organic compound
SVOC	semi-volatile organic compounds
TVOC	total volatile organic compounds
TSVOC	total semi-volatile organic compounds
TIC	total ion chromatogram
MS	mass spectrometric detector
FID	flame ionisation detector
GC	gas chromatograph
TD	thermal desorption

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5 Principle

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A measured volume of sample air is actively collected from indoor air, vehicle interior air and an emission test chamber (see ISO 16000-9, ISO 12219-4, ISO 12219-6) or an emission test cell (see ISO 16000-10) by drawing through one (or more) sorbent tube. VOC, VVOC and SVOC are retained by the sorbent tube, and the compounds are subsequently analysed in the laboratory to determine the identity, retained mass and associated air concentration of as many individual compounds as required by the specific test. Depending upon the range of target compounds the most appropriate sorbent tube(s), sampling and analytical conditions are applied. The collected compounds are desorbed by heat and transferred under inert carrier gas via a focussing trap into a gas chromatograph equipped with a capillary column and a mass spectrometric detector, with or without an additional flame ionization detector (FID).

6 Reagents and materials

6.1 Organic compounds for calibration of chromatographic quality

6.2 Dilution solvent for preparing calibration blend solution for liquid spiking, of chromatographic quality, free from compounds co-eluting with the compound(s) of interest ([6.1](#))

6.3 Solid sorbents

6.3.1 Introduction

Multiple sorbents, suitable for thermal desorption, are commercially available. They range in strength from very retentive sorbents required to retain and release VVOC to very weak sorbents suitable for quantitative sampling and release of SVOC. For particulate sorbents, the relevant particle size is 0,18 mm to 0,60 mm (30 mesh to 80 mesh). For a detailed list of sorbents see [Annex D](#).

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6.3.2 Quartz wool or glas/quartz beads, clean (i.e. do not produce analytically significant artefacts) and not prone to particle formation

6.3.3 Porous Polymers, i.e. Tenax TA^{®1)} particle size ~0,25 mm to ~0,6 mm (30 mesh to 60 mesh). Tenax TA[®] is a porous polymer based on 2,6-diphenyleneoxide. Manufactured Tenax TA[®] contains quantities of impurities, which shall be removed before using it for air sampling

6.3.4 “Carbon black” sorbents, such as Carbopack X^{®2)} or Carbograph 5 TD^{®3)}, particle size 0,25 mm to 0,5 mm (40 mesh to 60 mesh). Hydrophobic carbon sorbents suitable for VOC and VVOC with vapour pressures below those typical for C₄ hydrocarbons

6.3.5 Carbon melcular sieve (very strong) sorbents can also be used at the non-sampling end of the tube for trapping VVOC with vapour pressures above those typical for C₄ hydrocarbons. However, note that these sorbents are not completely hydrophobic. Therefore, if such sorbents are included, the tube needs to be dry purged in the sampling direction before analysis

6.4 Preparing calibration standards on sorbent tubes

As many identified substances as possible, or as required, should be calibrated using original reference compounds. Standards should be introduced to the sampling end of conditioned sorbent tubes using either liquid or gas phase standards.

6.4.1 Gas-phase standards

Standard atmospheres of known concentrations of the compound(s) of interest, shall be prepared by a recognized procedure such as ISO 16141⁴⁾ or ISO 6145⁴⁾. Typical concentrations are around 100 µg/m³ but levels will vary depending on test requirements. Alternatively, gas standards of appropriate quality and concentration shall be sourced commercially.

If the concentrations in any prepared standard atmosphere are not traceable to primary standards and/or if the inertness and stability of the atmospheres generated cannot be guaranteed, the concentration shall be confirmed using an independent procedure.

NOTE Producing gas phase standards of reactive and/or high boiling compounds can be particularly difficult. Frequent monitoring of the standard will be needed.

6.4.2 Loading sorbent tubes with gas-phase standards

Pass a known volume of standard atmosphere or gas standard through a conditioned sorbent tube from the sampling end, e.g. by means of a pump operating at 50 ml/min.

The volume of gas-phase standard sampled shall not exceed the breakthrough volume of sorbent tube for any of the compounds of interest.

After loading disconnect and seal the tube. Prepare fresh standards with each batch of samples. For indoor air and test chamber air load sorbent tubes with e.g. 100 ml, 200 ml, 400 ml, 1l, 2l, 4l or 10l of the 100 µg/m³ standard atmosphere selected.

1) Tenax TA[®] is a trade name of a product supplied by Buchem. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of the product named. Equivalent products may be used, if they can be shown to lead to the same results.

2) Carbopack X[®] is a trade name of Supelco. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of the product named. Equivalent products may be used if they can be shown to lead to the same results

3) Carbograph 5 TD[®] is a trade name of Lara. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of the product named. Equivalent products may be used, if they can be shown to lead the the same results.

6.4.3 Calibration blend solution for liquid spiking

Standard solution concentrations will vary depending on test requirements. The selected compound(s) shall be prepared or obtained as a liquid standard in chromatographic-grade solvent (e.g. in methanol) at an appropriate level – typically between 10 ng/μl and 1000ng/μl – depending on system sensitivity and the analytical conditions selected, for example split ratios. A suitably precise micro-syringe shall be used to introduce 1 μl aliquots of the check material onto the sampling end of sorbent tubes in a stream of inert gas as described. 1 μl is the suggested volume unless solvent can be selectively and efficiently purged without jeopardising breakthrough of the most volatile compounds(s) of interest.

The stability and safe storage times of calibration blend solutions shall be determined. Fresh standard solutions shall be prepared accordingly or if there is evidence of deterioration, e.g. reactions between alcohols and ketones.

Other techniques such as direct liquid spiking onto the sorbent bed without gas stream applied are also possible. In this case it is important to use tubes where the syringe needle can directly reach the sorbent bed.

6.4.4 Loading sorbent tubes with liquid standards

The sampling end of a sorbent tube is fitted to the unheated injection unit of the gas chromatograph (GC) (7.6) through which inert purge gas is passed at 100 ml/min, and a maximum 1μl aliquot of an appropriate standard solution is injected through the septum. After 5 min, the tube is disconnected and sealed. Prepare fresh standard tubes with each batch of samples.

NOTE 1 Multi-sorbent tubes, in particular with stronger sorbents are more difficult to solvent and standard solution. Smaller injection volumes are recommended for stronger sorbents and multi-sorbent tubes.

Introducing liquid standards on to sorbent tubes via a GC injector is considered the optimum approach to liquid standard introduction, as components reach the sorbent bed in the vapour phase.

Calibration mixtures should be prepared in controlled ambient temperature conditions. Before use, temper the solutions accordingly.

NOTE 2 When preparing standard tubes from liquid standards containing SVOC analytes, efficient transfer is enhanced if the configuration of the injector allows the tip of the syringe to make gentle contact with the sorbent retaining mechanism (e.g. gauze or frit) at the sampling end of the tube. It is important to keep liquid standard injection volumes to 1 μl or less unless the solvent can be selectively purged from the tube prior to analysis. Using small injection volumes minimises the difference between standards and samples during analysis thus minimising uncertainty.

NOTE 3 Standard tubes containing VVOC are more typically prepared either from standard atmospheres (see 6.4 and 6.5) or from concentrated gas standards sourced commercially. It is appropriate for concentrated gas standards to be introduced to the sampling end of sorbent tubes in a stream of carrier gas via an unheated GC injector or similar device.

An internal standard can be added by mixing with the calibration solution or by spiking separately.

NOTE 4 If standard tubes are being prepared by introducing aliquots from more than one standard solution or gas, it is appropriate to first introduce the standard containing higher boiling components and to introduce the most volatile organic compounds last. This minimizes risk of analyte breakthrough during the standard tube loading process.

NOTE 5 The purity of the inert carrier gas used to purge sorbent tubes during standard introduction (e.g. He, Ar, N₂) should permit the detection of an injection of 0,5 ng toluene. The quality of the carrier gas is of great importance, as any contaminants contained in the gas are enriched on the sorbent together with the substances to be analysed.

6.4.5 Commercial, pre-loaded standard tubes

Certified pre-loaded standard tubes are available and can be used for establishing analytical quality control and for routine calibration.