
**Cigarettes — Determination
of benzo[a]pyrene in cigarette
mainstream smoke using GC/MS —**

**Part 2:
Method using cyclohexane as
extraction solvent**

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*Cigarettes — Dosage du benzo[a]pyrène dans le courant principal de
la fumée de cigarette par GC/SM —*

*Partie 2: Méthode utilisant du cyclohexane comme solvant
d'extraction*

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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)

This document was prepared by Technical Committee ISO/TC 126, *Tobacco and tobacco products*.

This second edition cancels and replaces the first edition (ISO 22634-2:2017), which has been technically revised.

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

- reagents have modified by adding nitrogen;
- storage of standard solutions has been modified;
- sample clean-up has been modified;
- vortex mixer has been added;
- Bibliography has been extended.

A list of all parts in the ISO 22634 series can be found on the ISO website.

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, produced through collaborative experiments involving many laboratories in many countries, provides a procedure for the determination of B[a]P in cigarette mainstream smoke. The repeatability and reproducibility of this method have been assessed according to ISO recommendations and are included.

No machine smoking regime can represent all human smoking behaviours.

- It is recommended that cigarettes also be tested under conditions of a different intensity of machine smoking than those specified in this document.
- Machine smoking testing is useful to characterize cigarette emissions for design and regulatory purposes, but communication of machine measurements to smokers can result in misunderstandings about differences in exposure and risk across brands.
- Smoke emission data from machine measurements may be used as inputs for product hazard assessment, but they are not intended to be nor are they valid measures of human exposure or risks. Communicating differences between products in machine measurements as differences in exposure or risk is a misuse of testing using ISO standards.

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Cigarettes — Determination of benzo[a]pyrene in cigarette mainstream smoke using GC/MS —

Part 2: Method using cyclohexane as extraction solvent

WARNING — The use of this document can involve hazardous materials, operations and equipment. This document does not purport to address all the safety problems associated with its use. It is the responsibility of the user of this document to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.

1 Scope

This document specifies a method for the determination of benzo[a]pyrene (B[a]P) in the total particulate matter (TPM) of cigarette mainstream smoke using gas chromatography/mass spectrometry (GC/MS) with cyclohexane as extraction solvent.

This method is specified using ISO 3308 smoking parameters. This document provides an alternative method to that specified in ISO 22634-1, with a different clean-up, and a shorter total analytical run allowing a potential increase of sample throughput in comparison with ISO 22634-1.

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 3308, *Routine analytical cigarette-smoking machine — Definitions and standard conditions*

ISO 3402, *Tobacco and tobacco products — Atmosphere for conditioning and testing*

ISO 4387, *Cigarettes — Determination of total and nicotine-free dry particulate matter using a routine analytical smoking machine*

ISO 8243, *Cigarettes — Sampling*

3 Terms and definitions

No terms and definitions are listed in this document.

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

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

4 Principle

- Sampling of the test cigarettes according to the sampling procedure specified in ISO 8243.
- Conditioning of the test cigarettes according to the conditioning procedure specified in ISO 3402.
- Smoking of the test cigarettes according to the smoking procedure specified in ISO 4387.

- Extraction of the TPM, collected on the appropriate glass-fibre filter pad, with cyclohexane.
- Sample clean-up using solid phase extraction (SPE).
- Analytical determination of B[a]P by gas chromatography/mass spectrometry.

5 Apparatus

The usual laboratory apparatus and equipment and, in particular, the following.

5.1 Routine analytical cigarette-smoking machine, complying with the requirements of ISO 3308 and equipped for smoking in accordance with ISO 4387.

5.2 Gas chromatograph with a mass selective detector, equipped with its computerized control and data acquisition and processing system. This system shall be able to pilot the mass spectrometer in order to obtain chromatographic data under single ion monitoring (SIM) detection mode. The gas chromatograph shall be configured to perform splitless injections on a capillary column. It is recommended to equip the gas chromatograph with an autosampler for sample injection.

5.3 Fused silica capillary column, for example a 50 % phenyl-, 50 % methyl-polysiloxane stationary phase and a 30 m length, 0,25 mm internal diameter column with a 0,25 µm film thickness are suitable for this analysis.

NOTE Other columns can be used, provided that appropriate peak separation is obtained.

5.4 TurboVap®¹⁾ evaporator or equivalent equipment

5.5 Vacuum sample preparation unit or equivalent equipment

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5.6 Solid phase extraction cartridges, NH₂ bonded silica phase volume of 3 ml and packed with 500 mg is suitable.

NOTE Other cartridges with the same phase but different dimensions can be used as long as it is proved that results are equivalent.

5.7 Positive displacement pipettes, suitable for a volume range of 10 µl to 1 000 µl.

5.8 General laboratory equipment, for the preparation of samples, standards and reagents, e.g. sample vials (vial inserts may be required). All glassware shall be cleaned before use to avoid any contamination. Amber glassware may be required.

5.9 Ultrasonic bath.

5.10 Shaker, set to 200 r/min.

5.11 Vortex mixer.

6 Reagents

All reagents shall be of analytical grade quality.

6.1 Hexane, of known purity, not less than 99 %, CAS 110-54-3.

1) TurboVap® is an example of a suitable product available commercially. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of this product.

- 6.2 Cyclohexane**, of known purity, not less than 99 %, CAS 110-82-7.
- 6.3 Benzo[a]pyrene**, of known purity, not less than 98 %, CAS 50-32-8.
- 6.4 Benzo[a]pyrene-d12**, of known purity, not less than 98 %, CAS 63466-71-7.
- 6.5 Helium**, carrier gas of known purity, not less than 99,999 %, CAS 7440-59-7.
- 6.6 Nitrogen**, of known purity, not less than 99,999 %, CAS 7727-37-9.

WARNING — Benzo[a]pyrene and benzo[a]pyrene-d12 are carcinogens. Appropriate safety precautions shall be taken when manipulating these compounds or any solution containing these compounds.

7 Standards

7.1 General

Certified B[a]P or B[a]P-d12 solutions can be used as reference material.

7.2 Primary B[a]P-d12 stock solution: 100 µg/ml

Dissolve 10 mg B[a]P-d12, weighed to the nearest 0,01 mg, into a 100 ml volumetric flask and fill to the mark with cyclohexane. Sonicate to ensure dissolution.

7.3 Secondary B[a]P-d12 spiking solution: 40 ng/ml

Transfer 800 µl of the primary B[a]P-d12 stock solution (7.2) into a 2 000 ml volumetric flask and fill to the mark with cyclohexane.

7.4 Primary B[a]P stock solution: 100 µg/ml

Dissolve 10 mg B[a]P, weighed to the nearest 0,01 mg, into a 100 ml volumetric flask and fill to the mark with secondary B[a]P-d12 spiking solution (7.3).

7.5 Secondary B[a]P stock solution: 1 000 ng/ml

Dilute 1 ml of the primary B[a]P stock solution (7.4) into a 100 ml volumetric flask and fill to the mark with secondary B[a]P-d12 spiking solution (7.3).

7.6 Working standard solutions

Prepare six working standard solutions that cover the concentration range of interest. For example, transfer 100 µl of the secondary B[a]P stock solution (7.5) into a 20 ml volumetric flask and then fill to the mark with secondary B[a]P-d12 spiking solution (7.3). These solutions have a mass concentration of approximately 40 ng/ml of B[a]P-d12 and mass concentrations from 5,0 ng/ml to 250 ng/ml of B[a]P.

7.7 Storage of standard solutions

The standard solutions (7.2 to 7.6) are stable for up to four months if stored in the refrigerator at maximum 4 °C. Storage in amber glassware and away from the light is recommended.