



SLOVENSKI STANDARD

SIST EN ISO 6975:2005

01-julij-2005

Zemeljski plin – Širše analize – Plinsko-kromatografska metoda (ISO 6975:1997)

Natural gas - Extended analysis - Gas-chromatographic method (ISO 6975:1997)

Erdgas - Erweiterte Analyse - Gaschromatographisches Verfahren (ISO 6975:1997)

Gaz naturel - Analyse étendue - Méthode par chromatographie en phase gazeuse (ISO 6975:1997)

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ICS:

75.060

Zemeljski plin

Natural gas

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en

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EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

EN ISO 6975

May 2005

ICS 75.060

English version

**Natural gas - Extended analysis - Gas-chromatographic method
(ISO 6975:1997)**

Gaz naturel - Analyse étendue - Méthode par
chromatographie en phase gazeuse (ISO 6975:1997)

Erdgas - Erweiterte Analyse - Gaschromatographisches
Verfahren (ISO 6975:1997)

This European Standard was approved by CEN on 17 April 2005.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the Central Secretariat or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the Central Secretariat has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

Management Centre: rue de Stassart, 36 B-1050 Brussels

EN ISO 6975:2005 (E)**Foreword**

The text of ISO 6975:1997 has been prepared by Technical Committee ISO/TC 193 "Natural gas" of the International Organization for Standardization (ISO) and has been taken over as EN ISO 6975:2005 by CMC.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by November 2005, and conflicting national standards shall be withdrawn at the latest by November 2005.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

Endorsement notice

The text of ISO 6975:1997 has been approved by CEN as EN-ISO 6975:2005 without any modifications.

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INTERNATIONAL STANDARD

ISO
6975

Second edition
1997-04-01

Natural gas — Extended analysis — Gas-chromatographic method

iTeh STANDARD PREVIEW

*Gaz naturel — Analyse étendue — Méthode par chromatographie
en phase gazeuse*

SIST EN ISO 6975:2005

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Reference number
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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.

Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

International Standard ISO 6975 was prepared by Technical Committee ISO/TC 193, *Natural gas*, Subcommittee SC 1, *Analysis of natural gas*.

This second edition cancels and replaces the first edition (ISO 6975:1986), which has been technically revised.

Annexes A to D of this International Standard are for information only.

Introduction

This International Standard describes the specifications that an accurate method for the complete and extended analysis of natural gas shall fulfil.

The analytical methods involve injection of natural gas on to packed or open tubular (capillary) columns in one or more gas chromatographs. The components coming off the columns are detected by thermal-conductivity detector (TCD) or flame ionization detector (FID).

Compared to other analytical methods, the extended gas-chromatographic analysis supplies considerable knowledge of the individual components in the natural-gas mixture. This allows quantitative measurement of particular components in the C_6+ fraction and calculation of physical properties. Although the higher hydrocarbons influence physical properties such as calorific values and density by less than 0,3 % and 0,004 %, respectively, knowledge of them is required for evaluation of the pseudo-values for the C_6+ fraction.

In addition, vapour phase equilibrium calculations often require detailed composition data, especially of hydrocarbons with carbon numbers higher than 6.

The simultaneous determination of benzene and other aromatics is needed for gas consumers using natural gas as a chemical feedstock since these components can interfere in some processes.

Natural gas — Extended analysis — Gas-chromatographic method

1 Scope

This International Standard describes the specifications for the quantitative analysis of the following components of natural gas:

- helium
- hydrogen
- argon
- oxygen
- nitrogen
- carbon dioxide
- saturated hydrocarbons from C₁ to C₅
- hydrocarbon fractions from C₆ upwards
- aromatic compounds as benzene and toluene

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The gas-chromatographic methods determine the components in the following ranges:

oxygen:	0,001 % (n/n)	to	0,5 % (n/n)
helium:	0,001 % (n/n)	to	0,5 % (n/n)
hydrogen:	0,001 % (n/n)	to	0,5 % (n/n)
argon:	0,001 % (n/n)	to	0,5 % (n/n)
nitrogen:	0,001 % (n/n)	to	40 % (n/n)
carbon dioxide:	0,001 % (n/n)	to	40 % (n/n)
methane:	50 % (n/n)	to	100 % (n/n)
ethane:	0,02 % (n/n)	to	15 % (n/n)
propane:	0,001 % (n/n)	to	5 % (n/n)

higher hydrocarbons: The method can measure hydrocarbon components from 10⁻⁶ (n/n) up to their maximum concentration, which is compatible with the requirement that the gas is free from hydrocarbon condensate at any pressure in the range 1 × 10² kPa to 7 × 10³ kPa.

This method is not intended for the determination of oxygen compounds (water vapour, methanol, glycols) or sulfur compounds.

It is not possible to make unambiguous identifications of hydrocarbons above C_6 . Even where "spiking" a gas mixture with known components shows where they elute, it cannot be stated with certainty that such a component is the only one with that retention time. Unidentified components are classified according to the carbon number which the analysis indicates to be appropriate. While this is a necessary simplification, it does allow a reasonable quantitative value to be obtained.

This method is intended for use in situations where the hexanes plus compositional breakdown and/or the complete analysis is desired.

The method is not intended for dense phase gases whose pressure exceeds the cricondebar (critical condensation pressure), or for gas samples containing any measurable hydrocarbon condensate, liquid water or process fluid such as methanol or glycols (see ISO 6570-1 and ISO 10715).

Gases which have been treated for transmission are unlikely to contain detectable levels of hydrocarbons above C_{12} . Samples taken from nearer the well head, before the gas has reached gas treatment plants, may contain hydrocarbons to C_{16} or above.

2 Normative references

The following standards contain provisions which, through reference in this text, constitute provisions of this International Standard. At the time of publication, the editions indicated were valid. All standards are subject to revision, and parties to agreements based on this International Standard are encouraged to investigate the possibility of applying the most recent editions of the standards indicated below. Members of IEC and ISO maintain registers of currently valid International Standards.

ISO 5725-1:1994, *Accuracy (trueness and precision) of measurement methods and results — Part 1: General principles and definitions*. <https://standards.iteh.ai/catalog/standards/sist/8a973b69-1e17-49fe-b65b-f84b38369166/sist-en-iso-6975-2005>

ISO 6142:1981, *Gas analysis — Preparation of calibration gas mixtures — Weighing methods*.

ISO 6143:1981, *Gas analysis — Determination of composition of calibration gas mixtures — Comparison methods*.

ISO 6326-1:1989, *Natural gas — Determination of sulfur compounds — Part 1: General introduction*.

ISO 6326-2:1981, *Gas analysis — Determination of sulfur compounds in natural gas — Part 2: Gas chromatographic method using an electrochemical detector for the determination of odoriferous sulfur compounds*.

ISO 6326-3:1989, *Natural gas — Determination of sulfur compounds — Part 3: Determination of hydrogen sulfide, mercaptan sulfur and carbonyl sulfide sulfur by potentiometry*.

ISO 6326-4:1994, *Natural gas — Determination of sulfur compounds — Part 4: Gas chromatographic method using a flame photometric detector for the determination of hydrogen sulfide, carbonyl sulfide and sulfur-containing odorants*.

ISO 6326-5:1989, *Natural gas — Determination of sulfur compounds — Part 5: Lingener combustion method*.

ISO 6570-1:1983, *Natural gas — Determination of potential hydrocarbon liquid content — Part 1: Principles and general requirements*.

ISO 6974-1:—¹⁾, *Natural gas — Determination of composition with defined uncertainty by gas chromatography — Part 1: Guidelines for tailored analysis*.

1) To be published. (Revision, in parts, of ISO 6974:1984)

ISO 6974-2:—²⁾, *Natural gas — Determination of composition with defined uncertainty by gas chromatography — Part 2: Measuring system characteristics and statistics for data treatment.*

ISO 6974-3:—²⁾, *Natural gas — Determination of composition with defined uncertainty by gas chromatography — Part 3: Determination of hydrogen, helium, inert gases and hydrocarbons up to C₈.*

ISO 6974-4:—²⁾, *Natural gas — Determination of composition with defined uncertainty by gas chromatography — Part 4: Determination of nitrogen, carbon dioxide and hydrocarbons (C₁ up to C₅ and C₆₊) for a laboratory and on-line measuring system.*

ISO 6974-5:—²⁾, *Natural gas — Determination of composition with defined uncertainty by gas chromatography — Part 5: Determination of nitrogen, carbon dioxide and hydrocarbons (C₁ up to C₅ and C₆₊) for a laboratory and on-line process application.*

ISO 10715:1997, *Natural gas — Sampling guidelines.*

ISO 14111:1997, *Natural gas — Guidelines for traceability in analysis.*

3 Definitions

For the purposes of this International Standard, the following definitions apply.

3.1 resolution: Gas-chromatographic resolution is a characteristic of the separation of two adjacent peaks and is measured as twice the distance between the maxima of the named peaks divided by the sum of the intercepts on the baseline made by tangents drawn to the peaks at half their height (see figure 1). The resolution R_{AB} may be expressed by the following equation:

$$R_{AB} = 2 \times \frac{d_R(B) - d_R(A)}{w(B) + w(A)}$$

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where

$d_R(A)$ and $d_R(B)$ are the retention distances of the eluted components A and B;

$w(A)$ and $w(B)$ are the widths of the respective peaks at their base.

3.2 main components: The nitrogen, carbon dioxide and saturated hydrocarbons from methane to *n*-pentane present in a natural-gas sample.

3.3 associated components: The helium, hydrogen, argon and oxygen present in a natural-gas sample.

3.4 trace components: The hydrocarbons and/or groups of hydrocarbons from *n*-pentane upwards present in a natural-gas sample.

3.5 other components: Those components for which this method is not intended, such as oxygen compounds (water vapour, methanol, glycol) and sulfur compounds.

3.6 response: The response is the output signal for a component that is measured as peak area or peak height (more often as peak area).

²⁾ To be published. (Revision, in parts, of ISO 6974:1984)

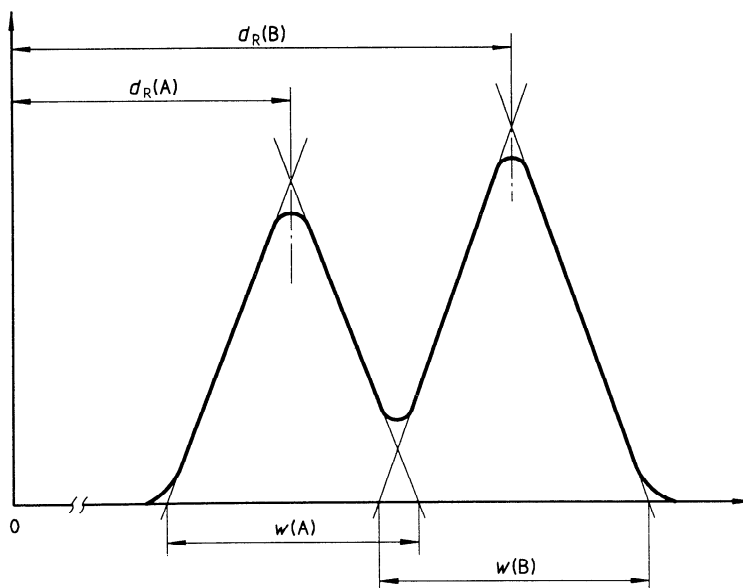


Figure 1 — Resolution of two adjacent peaks

3.7 reference component: The component present in the working-reference gas mixture against which the relative response factors of sample components not present in the working-reference gas mixture are defined.

3.8 relative response factor (for an FID): The relative response factor K_i is calculated as the ratio of the carbon number of the reference component to the carbon number of a particular sample component. Thus if the reference component is *n*-butane, then the relative response factor for C_7 components in the sample (heptane isomers, methylcyclohexane, toluene, etc.) is calculated as

$$K_i = \frac{4}{7} = 0,571$$

3.9 concentration of a group of components: The concentration of a group of components is the sum of the concentrations of the normal hydrocarbon components and their isomers. The response of the group is the sum of the responses of these components. The relative response of the group is equal to the relative response of the normal alkane of the group. The group is named according to the normal alkane of the group.

4 Principle

The components to be determined in a gaseous sample are physically separated by gas chromatography and compared with calibration data obtained under the same set of conditions.

The main components are separated using a packed or open tubular column in a gas chromatograph and detected by TCD. An FID may be used additionally.

The analytical system used separates oxygen from nitrogen to check air contamination in the sample.

The hydrocarbons from propane upwards are separated using an open tubular column in a gas chromatograph and detected by FID. The method assumes that the response of an FID to hydrocarbons is proportional to the carbon number. Hence, components not present in the calibration gas are quantified by means of knowledge of their carbon number.

Associated components can be determined by following special gas-chromatographic techniques (see e.g. the various parts of ISO 6974).