



SLOVENSKI STANDARD
oSIST prEN ISO 2613-2:2023
01-april-2023

Analiza zemeljskega plina - Biometan - 2. del: Določevanje siloksana s plinsko kromatografijo s spektrometrijo na osnovi mobilnosti ionov (GC-IMS) (ISO/DIS 2613-2:2023)

Analysis of natural gas - Biomethane - Part 2: Determination of siloxane content by gas chromatography ion mobility spectrometry (ISO/DIS 2613-2:2023)

Analyse von Erdgas - Siliziumanteil von Biomethan - Teil 2: Bestimmung des Siloxananteils durch gaschromatographische Ionenmobilitätsspektroskopie (ISO/DIS 2613-2:2023)

Analyse du gaz naturel - Biométhane - Partie 2: Détermination de la teneur en siloxanes par chromatographie en phase gazeuse spectrométrie de mobilité ionique (ISO/DIS 2613-2:2023)

Ta slovenski standard je istoveten z: prEN ISO 2613-2

ICS:

75.060 Zemeljski plin Natural gas

oSIST prEN ISO 2613-2:2023 **en,fr,de**

DRAFT INTERNATIONAL STANDARD

ISO/DIS 2613-2

ISO/TC 193/SC 1

Secretariat: NEN

Voting begins on:
2023-02-13Voting terminates on:
2023-05-08

Analysis of natural gas — Biomethane —

Part 2:

Determination of siloxane content by gas chromatography ion mobility spectrometry

ICS: 75.060

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Reference number
ISO/DIS 2613-2:2023(E)

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Published in Switzerland

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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.

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This document was prepared by Technical Committee ISO/TC 193, Natural gas, Subcommittee SC 1, Analysis of natural gas.

A list of all parts in the ISO 2613 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 describes a method for the determination of the siloxane content of biomethane that is applicable to both laboratory and online analysis.

Siloxanes are a common contaminant in biogas, caused by their extensive use in common household products such as cosmetics and detergents. These products eventually become entrained in waste streams that are used to generate biogas. Additionally, siloxanes are sometimes added to digesters as anti-foaming agents to improve generation efficiency.

If not effectively removed, siloxanes can cause damage to gas processing and vehicle injection equipment through formation of silica deposits at high temperatures. Many gas distributors require measurement reports to demonstrate that the resultant biomethane conforms with their requirements, which typically includes siloxane content due to its potential to cause damage. Additionally, producers may wish to monitor siloxane content for process optimisation purposes.

The technique described in this standard is commercially available and tailored specifically to measuring siloxanes within biomethane at low concentrations (for instance, as specified in EN 16723-1^[1] and EN 16723-2^[2]). It is suitable for use both in the laboratory and field.

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Analysis of natural gas — Biomethane —

Part 2:

Determination of siloxane content by gas chromatography ion mobility spectrometry

1 Scope

This document describes a Gas Chromatography – Ion Mobility Spectroscopy (GC-IMS) method for the determination of the concentration of siloxanes in biomethane. The method is applicable to the following siloxanes:

- hexamethyldisiloxane (L2)
- octamethyltrisiloxane (L3)
- decamethyltetrasiloxane (L4)
- dodecamethylpentasiloxane (L5)
- hexamethylcyclotrisiloxane (D3)
- octamethylcyclotetrasiloxane (D4)
- octamethylcyclotetrasiloxane (D5)
- dodecamethylcyclohexasiloxane (D6)

This document describes suitable calibration and measurement strategies to quantify siloxanes in (bio) methane around and above the $0,3 \text{ mg m}^{-3}$ ($14 \text{ } \mu\text{mol mol}^{-1}$) level and applies to analysis within absolute pressure ranges of 1 bar – 2 bar, temperatures of $0 \text{ } ^\circ\text{C}$ – $40 \text{ } ^\circ\text{C}$ and relative humidity $< 90 \%$.

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 6142-1, *Gas analysis — Preparation of calibration gas mixtures — Part 1: Gravimetric method for Class I mixtures*

ISO 6143, *Gas analysis — Comparison methods for determining and checking the composition of calibration gas mixtures*

ISO 6144, *Gas analysis — Preparation of calibration gas mixtures — Static volumetric method*

ISO 6145-1, *Gas analysis — Preparation of calibration gas mixtures using dynamic methods — Part 1: General aspects*

ISO 6974-1, *Natural gas — Determination of composition and associated uncertainty by gas chromatography — Part 1: General guidelines and calculation of composition*

ISO 6974-2, *Natural gas — Determination of composition and associated uncertainty by gas chromatography — Part 2: Uncertainty calculations*

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ISO 7504, *Gas analysis — Vocabulary*

ISO 10715, *Natural gas — Gas sampling*

ISO 10723, *Natural gas — Performance evaluation for analytical systems*

ISO 14532, *Natural gas — Vocabulary*

ISO 14912, *Gas analysis — Conversion of gas mixture composition data*

ISO/IEC guide 98-3:2008, *Uncertainty of measurement — Part 3: Guide to the expression of uncertainty in measurement (GUM:1995)*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 7504, ISO/IEC guide 98-3:2008, ISO 14532 and the following apply.

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 <https://www.electropedia.org/>

**3.1 ion mobility spectrometry
IMS**
analytical method used to separate and identify ionised molecules in the gas phase based on their mobility in a carrier gas under an electric field

4 Principle

A gaseous biomethane sample is drawn into a sample loop via a vacuum pump. The sample is injected onto a GC column, which separates the individual components in the gas mixture. As they exit the column, the components are ionised and passed into a drift tube. The drift tube applies an electric field and a flow of inert nitrogen drift gas that causes the ions to drift at different rates based on their mass and geometric structure onto a detector.

The instrument is calibrated by sampling an appropriate range of siloxane concentrations and using the detector output to generate a calibration function. The function can then be applied to unknown samples to calculate the concentration of siloxanes. This can then be summed to calculate total siloxanes and total silicon.

The method is applicable to siloxanes within the concentration ranges provided in [Table 1](#).

Table 1 — Application ranges

Component	Abbreviation	Formula	Lower limit (mg m ⁻³)	Upper limit (mg m ⁻³)
hexamethyldisiloxane	L2	C ₆ H ₁₈ Si ₂ O	0,1	10,0
octamethyltrisiloxane	L3	C ₈ H ₂₄ Si ₃ O ₂	0,1	10,0
decamethyltetrasiloxane	L4	C ₁₀ H ₃₀ Si ₄ O ₃	0,1	10,0
dodecamethylpentasiloxane	L5	C ₁₂ H ₃₆ Si ₅ O ₄	0,1	10,0
hexamethylcyclotrisiloxane	D3	C ₁₂ H ₁₈ O ₃ Si ₃	0,1	10,0
octamethylcyclotetrasiloxane	D4	C ₈ H ₂₄ O ₄ Si ₄	0,1	10,0
octamethylcyclotetrasiloxane	D5	C ₁₀ H ₃₀ O ₅ Si ₅	0,1	10,0
dodecamethylcyclohexasiloxane	D6	C ₁₂ H ₃₆ O ₆ Si ₆	0,3	10,0