

SLOVENSKI STANDARD oSIST prEN 14211:2022

01-december-2022

Zunanji zrak - Standardna metoda za določanje koncentracije dušikovega dioksida in dušikovega monoksida s kemiluminiscenco

Ambient air - Standard method for the measurement of the concentration of nitrogen dioxide and nitrogen monoxide by chemiluminescence

Außenluft - Messverfahren zur Bestimmung der Konzentration von Stickstoffdioxid und Stickstoffmonoxid mit Chemilumineszenz

Air ambiant - Méthode normalisée pour le mesurage de la concentration en dioxyde d'azote et monoxyde d'azote par chimiluminescence

Ta slovenski standard je istoveten z: prEN 14211

ICS:

13.040.20 Kakovost okoljskega zraka Ambient atmospheres

oSIST prEN 14211:2022 en,fr,de

oSIST prEN 14211:2022

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oSIST prEN 14211:2022 https://standards.iteh.ai/catalog/standards/sist/f8960ae4-6c7a-4ac2-913b-a438306ba0d4/osist-pren-14211-2022

EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

DRAFT prEN 14211

October 2022

ICS 13.040.20

Will supersede EN 14211:2012

English Version

Ambient air - Standard method for the measurement of the concentration of nitrogen dioxide and nitrogen monoxide by chemiluminescence

Air ambiant - Méthode normalisée pour le mesurage de la concentration en dioxyde d'azote et monoxyde d'azote par chimiluminescence Außenluft - Messverfahren zur Bestimmung der Konzentration von Stickstoffdioxid und Stickstoffmonoxid mit Chemilumineszenz

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

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Contents Pag European foreword		
2	Normative references	6
3	Terms and definitions	6
4	Abbreviated terms	
5	Principle	
5 5.1	General	
5.2	Measuring principle	
5.3	Type testing	
5.4	Field operation and quality control	
6	Sampling	14
6.1	General	14
6.2	Sampling location	14
6.3	Sampling system	
6.4	Control and regulation of sample flow rate	16
6.5	Sampling pump for the manifold	16
7	Analyser equipmentGeneral	16
7.1		
7.2	Converter	
7.3	Ozone generator	
7.4	Reaction chamberSIST DEN 142112022	
7.5	Optical filter s. //standards.iteh.ai/catalog/standards/sist/f8960ac4-6c7a-4ac2-913h-	
7.6 7.7	Detector	
7.7 7.8	Ozone removal deviceSampling pump for the analyser	
7.8 7.9	Particle filter	
	Type testing of nitrogen dioxide and nitrogen monoxide analysers	
8 8.1	GeneralGeneral	
8.2	Relevant performance characteristics and performance criteria	
8.3	Design change	
8.4	Procedures for determination of the performance characteristics during the	
	laboratory test	22
8.5	Determination of the performance characteristics during the field tests	
8.6	Type testing and uncertainty calculation	
9	Field operation and ongoing quality control	42
9.1	General	
9.2	Suitability evaluation	42
9.3	Initial installation	44
9.4	Ongoing quality assurance/quality control	45
9.5	Calibration of the analyser	
9.6	Checks	
9.7	Maintenance	
9.8	Data handling and data reports	
9.9	Measurement uncertainty	56

10	Expression of results	56
11 11.1 11.2	Test reports and documentation	57
	A (normative) Test of lack of fit	
Annex A.1	Establishment of the regression line	
A.1 A.2	_	
	Calculation of the residuals of the averages	
	B (informative) Sampling equipment	
	C (informative) Types of chemiluminescence analysers	
	D (informative) Manifold testing	
Annex	E (normative) Type testing	
E.1	Type testing and uncertainty calculation	69
E.2	Type testing Requirement a)	69
E.3	Type testing Requirement b)	71
E.4	Type testing Requirement c)	83
E.5	Type testing Requirement d)	83
Annex	F (informative) Calculation of uncertainty in field operation at the hourly limit value	90
F.1	General	90
F.2	Combined standard uncertainty	90
F.3	Standard uncertainties	91
F.4	Example calculation	98
Annex	G (informative) Calculation of uncertainty in field operation at the annual limit value	100
G.1	General	100
Annex	H (informative) Test stand for the test point "sensitivity coefficient of sample gas pressure"	101
H.1	General	101
Annex	I (informative) Significant technical changes	103
I.1	Details of significant technical changes between this document and the previous edition	
Biblio	graphy	
	5^ ~ T ~ V	

European foreword

This document (prEN 14211:2022) has been prepared by Technical Committee CEN/TC 264 "Air quality", the secretariat of which is held by DIN.

This document is currently submitted to the CEN Enquiry.

This document will supersede EN 14211:2012.

The technical modifications in comparison with the previous edition are listed in Annex I of this document.

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1 Scope

This document specifies a continuous measurement method for the determination of the concentrations of nitrogen dioxide and nitrogen monoxide present in ambient air based on the chemiluminescence measuring principle. This document describes the performance characteristics and sets the relevant minimum criteria required to select an appropriate chemiluminescence analyser by means of type testing. It also includes the evaluation of the suitability of an analyser for use in a specific fixed site so as to meet the data quality requirements as specified in Annex I of Directive 2008/50/EC [1] and requirements during sampling, calibration and quality assurance for use.

The method is applicable to the determination of the concentration of nitrogen dioxide present in ambient air up to $500~\mu g/m^3$. This concentration range represents the certification range for nitrogen dioxide for type testing.

The method is applicable to the determination of the concentration of nitrogen monoxide present in ambient air up to $1\,200\,\mu\text{g/m}^3$. This concentration range represents the certification range for nitrogen monoxide for the type testing.

NOTE 1 It is possible to use other ranges depending on the levels present in ambient air.

NOTE 2 When this document is used for purposes other than for measurements required by Directive 2008/50/EC, the ranges and uncertainty requirements possibly do not apply.

The method covers the determination of ambient air concentrations of nitrogen dioxide and nitrogen monoxide in zones classified as rural areas, urban-background areas, traffic-orientated locations and locations influenced by industrial sources.

The results are expressed in µg/m³ (at 20 °C and 101,3 kPa).

NOTE 3 500 μ g/m³ of nitrogen dioxide corresponds to 261 nmol/mol of nitrogen dioxide at 20 °C and 101,3 kPa. 1 200 μ g/m³ of nitrogen monoxide corresponds to 962 nmol/mol of nitrogen monoxide at 20 °C and 101,3 kPa.

This document contains information for different groups of users. 4-6c7a-4ac2-913b-

Clause 5 to Clause 7 and Annex B and Annex C contain general information about the principles of NO_x measurement by chemiluminescence analyser and sampling equipment.

Clause 8 and Annex E are specifically directed towards test houses and laboratories that perform type testing of NO_x analysers. These sections contain information about:

- type testing conditions, test procedures and test requirements;
- analyser performance requirements;
- evaluation of the type testing results;
- evaluation of the uncertainty of the measurement results of the NO_x analyser based on the type testing results.

Clause 9 to Clause 11 and Annex F and Annex G are directed towards monitoring networks performing the practical measurements of NO_x in ambient air. These sections contain information about:

- initial installation of the analyser in the monitoring network and acceptance testing;
- ongoing quality assurance/quality control;
- calculation and reporting of measurement results;
- evaluation of the uncertainty of measurement results under practical monitoring conditions.

This document represents an evolution of earlier editions (EN 14211:2005 and EN 14211:2012). It is advisable that when equipment is procured it complies fully with this document.

NOTE 4 Type testing performed prior to the publication of this document for the purpose of demonstrating equivalence are still valid.

NOTE 5 Analysers type tested prior to the publication of this document remain valid for use for regulated monitoring purposes.

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.

EN 15267-1, Air quality - Certification of automated measuring systems - Part 1: General principles

EN 15267-2, Air quality - Certification of automated measuring systems - Part 2: Initial assessment of the AMS manufacturer's quality management system and post certification surveillance for the manufacturing process

EN ISO 6142, Gas analysis — Preparation of calibration gas mixtures — Gravimetric method (ISO 6142)

EN ISO 6143, Gas analysis - Comparison methods for determining and checking the composition of calibration gas mixtures (ISO 6143)

EN ISO 6144, Gas analysis - Preparation of calibration gas mixtures - Static volumetric method (ISO 6144)

EN ISO 6145-6, Gas analysis - Preparation of calibration gas mixtures using dynamic methods - Part 6: Critical flow orifices (ISO 6145-6)

EN ISO 6145-7, Gas analysis - Preparation of calibration gas mixtures using dynamic methods - Part 7: Thermal mass-flow controllers (ISO 6145-7)

EN ISO 6145-10, Gas analysis - Preparation of calibration gas mixtures using dynamic volumetric methods - Part 10: Permeation method (ISO 6145-10)

EN ISO 14956, Air quality - Evaluation of the suitability of a measurement procedure by comparison with a required measurement uncertainty (ISO 14956)

ISO 19229, Gas analysis - Purity analysis and the treatment of purity data

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:

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

3.1

adjustment

adjustment of a measuring system

set of operations carried out on a measuring system so that it provides prescribed indications corresponding to given values of a quantity to be measured

Note 1 to entry: Types of adjustment of a measuring system include zero adjustment of a measuring system, offset adjustment, and span adjustment (sometimes called gain adjustment).

Note 2 to entry: Adjustment of a measuring system should not be confused with calibration, which is a prerequisite for adjustment.

[SOURCE: JCGM 200:2012, 3.11] [2]

Note 3 to entry: In the context of this document, adjustment is generally performed on measurement data rather than on the analyser.

3.2

ambient air

outdoor air in the troposphere, excluding workplaces as defined by Directive 89/654/EEC, where provisions concerning health and safety at work apply and to which members of the public do not have regular access

[SOURCE: Council Directive 2008/50/EC] [1]

3.3

analyser

measuring system that provides an output signal which is a function of the concentration, partial pressure, flow or temperature of one or more components of a gas mixture

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availability of the analyser

fraction of the time period for which valid measuring data of the ambient air concentration is available from an analyser

3.5

calibration

operation that, under specified conditions, in a first step, establishes a relation between the quantity values with measurement uncertainties provided by measurement standards and corresponding indications with associated measurement uncertainties and, in a second step, uses this information to establish a relation for obtaining a measurement result from an indication

Note 1 to entry: A calibration may be expressed by a statement, calibration function, calibration diagram, calibration curve, or calibration table. In some cases, it may consist of an additive or multiplicative correction of the indication with associated measurement uncertainty.

Note 2 to entry: Calibration should not be confused with adjustment of a measuring system, often mistakenly called "self-calibration", nor with verification of a calibration.

Note 3 to entry: Often, the first step alone in the above definition is perceived as being calibration.

[SOURCE: JCGM 200:2012, 2.39] [2]

Note 4 to entry: In the context of this document, calibration is a comparison of the analyser response to a known gas concentration with a known uncertainty when the information obtained from the comparison is used for the successive adjustment (if needed) of the analyser.

3.6

certification range

concentration range for which the analyser is type tested

3.7

check

verification that the analyser is still operating within specified performance limits

3.8

combined standard uncertainty

standard uncertainty of the result of a measurement when that result is obtained from the values of a number of other quantities, equal to the positive square root of a sum of terms, the terms being the variances or co-variances of these other quantities weighted according to how the measurement result varies with changes in these quantities

[SOURCE: JCGM 100:2008, 2.3.4] [3]

3.9

converter efficiency

degree of conversion of nitrogen dioxide present in the sample gas into NO, given as a percentage

3.10

coverage factor

numerical factor used as a multiplier of the combined standard uncertainty in order to obtain an expanded uncertainty $oSIST\ prEN\ 142\ 11:2022$

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[SOURCE: JCGM 100:2008, 2.3.6] [3] 438306ba0d4/osist-pren-14211-2022

3.11

designated body

body which has been designated for a specific task (type testing and/or QA/QC activities in the field) by the competent authority in the Member States

3.12

detection limit

smallest concentration of a measurand that can be reliably detected by a specific measurement process

Note 1 to entry: The detection limit is calculated as $3.3 \times (s_z/B)$ where s_z is the standard deviation of analyser response at zero measurand concentration (see 8.4.5) and B is the slope of the calibration function [4].

3.13

expanded uncertainty

quantity defining an interval about the result of a measurement that may be expected to encompass a large fraction of the distribution of values that could reasonably be attributed to the measurand

Note 1 to entry: The fraction may be viewed as the coverage probability or level of confidence of the interval.

Note 2 to entry: To associate a specific level of confidence with the interval defined by the expanded uncertainty requires explicit or implicit assumptions regarding the probability distribution characterized by the measurement result and its combined standard uncertainty. The level of confidence that may be attributed to this interval can be known only to the extent to which such assumptions may be justified.

[SOURCE: JCGM 100:2008, 2.3.6] [3]

Note 3 to entry: For the purpose of this document, the expanded uncertainty is the combined standard uncertainty multiplied by a coverage factor k = 2 resulting in an interval with a level of confidence of 95 %.

3.14

fall time

difference between the response time (fall) and the lag time (fall)

3.15

independent measurement

individual measurement that is not influenced by a previous individual measurement by separating two individual measurements by at least four response times

Note 1 to entry: The largest value of response time (rise) and response time (fall) are intended.

3.16

individual measurement

measurement averaged over a time period equal to the response time of the analyser

Note 1 to entry: The largest value of response time (rise) and response time (fall) are intended.

Note 2 to entry: This definition differs from the meaning of the concept "individual measurement" in Directive 2008/50/EC [1].

3.17

influence quantity

quantity that is not the measurand but that affects the result of the measurement

[SOURCE: JCGM 100:2008, B.2.10] [3]

3.18

interferent

component of the air sample, excluding the measured constituent, that affects the output signal

3.19

lack of fit

maximum deviation from the linear regression line of the average of a series of measurement results at the same concentration

3.20

lag time

time interval from the moment at which a step change of sample concentration occurs at the inlet of the analyser to the moment at which the output reading reaches a level corresponding to a predefined change of the stable output reading

3.21

limit value

level fixed on the basis of scientific knowledge, with the aim of avoiding, preventing or reducing harmful effects on human health and/or the environment as a whole, to be attained within a given period and not to be exceeded once attained

[SOURCE: Council Directive 2008/50/EC] [1]

3.22

long term drift

difference between zero or span readings over a determined period of time (e.g. period of unattended operation)

3.23

monitoring station

enclosure located in the field in which an analyser has been installed to monitor concentrations of one or more ambient air pollutants

3.24

parallel measurement

measurement from different analysers, sampling from one and the same sampling manifold, starting at the same time and ending at the same time

3.25

performance characteristic

one of the parameters assigned to equipment in order to define its performance

3.26

performance criterion

limiting quantitative numerical value assigned to a performance characteristic, to which conformance is tested

3.27

period of unattended operation (standards.iteh.ai)

time period over which the drift complies with the performance criterion for long term drift

3.28

repeatability (of results of measurement) talog/standards/sist/f8960ae4-6c7a-4ac2-913b-

closeness of the agreement between the results of successive individual measurements of nitrogen monoxide and nitrogen dioxide carried out under the same conditions of measurement

Note 1 to entry: These conditions include:

- the same measurement procedure;
- the same observer;
- the same analyser, used under the same conditions;
- at the same location;
- repetition over a short period of time.

3.29

reproducibility under field conditions

closeness of the agreement between the results of simultaneous measurements with two analysers in ambient air carried out under the same conditions of measurement

Note 1 to entry: These conditions are called field reproducibility conditions and include:

- the same measurement procedure;
- two identical analysers, used under the same conditions;

- at the same monitoring station;
- the period of unattended operation.

3.30

residence time inside the analyser

time period for the sampled air to be transported from the inlet of the analyser to the reaction chamber for the NO-channel

3.31

residence time in the sampling system

time period for the sampled air to be transferred to the inlet of the analyser

3.32

response time

time interval from the instant at which a step change of sample concentration occurs at the inlet of the analyser to the instant at which the output reading reaches a level corresponding to a predefined change of the output reading

3.33

sampled air

part of ambient air that is transferred through the sampling inlet and sampling system for subsequent measurement

3.34

sample gas temperature (standards.iteh.ai)

temperature of the sampled gas at the sample inlet

Note 1 to entry: The term "gas" can refer to a test gas used in type testing or to ambient air transferred to the analyser. $\frac{1}{3} \frac{1}{3} \frac{1}{3}$

3.35

sampling system

the assembly of components needed to transfer ambient air to the analyser

3.36

short-term drift

difference between zero or span readings at the beginning and end of a 12 h period

3.37

standard uncertainty

uncertainty of the result of a measurement expressed as a standard deviation

[SOURCE: JCGM 100:2008, 2.3.1] [3]

3.38

surrounding temperature

temperature of the air directly surrounding the analyser

3.39

total residence time

sum of the residence time in the sampling system and the residence time inside the analyser

3.40

type testing

examination of two or more analysers of the same design which are submitted by a manufacturer to a specified body including the tests necessary for approval of the design (determination of fitness for purpose of a specific device model / of an unambiguous designated analyser)

3.41

uncertainty

uncertainty of measurement parameter associated with the result of a measurement that characterises the dispersion of the values that could reasonably be attributed to the measurand

4 Abbreviated terms

AMS automated measuring system (sometimes referred to as continuous automated measuring

system (CAM))

FEP perfluoro-ethylene-propylene

MFC mass flow controller

PFA perfluoro-alkoxy

PTFE polytetrafluoroethylene

5 Principle iTeh STANDARD PREVIEW

5.1 General

This document describes the method for measurement of the concentrations of nitrogen dioxide and nitrogen monoxide in ambient air by means of chemiluminescence. The requirements, the specific components of the chemiluminescence analyser and its sampling system are described. A number of performance characteristics with associated minimum performance criteria are given for the analyser. The actual values of these performance characteristics for a specific type of analyser shall be determined in a so-called type test for which procedures have been described. The type testing comprises a laboratory test and a field test. The selection of a type-tested analyser for a specific measuring task in the field is based on the calculation of the expanded uncertainty of the measurement method. In this expanded uncertainty calculation, the actual values of various performance characteristics of a type tested analyser and the site-specific conditions at the monitoring station are taken into account (see 9.6). The expanded uncertainty of the method shall not exceed the limits (this is also specified in Annex I of

Directive 2008/50/EC [1]). Requirements and recommendations for quality assurance and quality

5.2 Measuring principle

control are given for the measurements in the field (see 9.4).

Chemiluminescence is based on the reaction of nitrogen monoxide with ozone. In a chemiluminescence analyser, air is sampled through a filter (to prevent contamination of the gas conveying system, especially the optical components of the analyser) and fed at a constant flow rate into the reaction chamber of the analyser, where it is mixed with an excess of ozone for the determination of nitrogen monoxide only. The emitted radiation (chemiluminescence) is proportional to the number of nitrogen monoxide molecules in the detection volume and thus proportional to the concentration of nitrogen monoxide. The emitted radiation is filtered by a selective optical filter and converted into an electric signal by a photomultiplier tube or a photodiode.

For the determination of nitrogen dioxide, the sampled air is fed through a converter where the nitrogen dioxide is reduced to nitrogen monoxide and analysed in the same way as described previously. The electrical signal obtained from the photomultiplier tube or photodiode is proportional to the sum of

concentrations of nitrogen dioxide and nitrogen monoxide. The amount of nitrogen dioxide is calculated from the difference between this concentration and that obtained for nitrogen monoxide only (when the sampled air has not passed through the converter).

Chemiluminescence is the emission of light during a chemical reaction. During the gas-phase reaction of nitrogen monoxide and ozone light with an intensity proportional to the concentration of nitrogen monoxide is produced when electrons of the excited nitrogen dioxide molecules decay to lower energy states.

This chemiluminescence method is based on the reaction:

$$NO + O_3 \rightarrow NO_2^* + O_2$$
 (1)

$$NO_2^* \to NO_2 + hv \tag{2}$$

Excited nitrogen dioxide (NO_2^*) emits radiation in the near infrared region (600 nm to 3 000 nm) with a maximum centred around 1 200 nm. For the determination of nitrogen dioxide, the nitrogen dioxide present in sampled air is converted to nitrogen monoxide in a converter as a result of the reaction:

$$\begin{array}{c} converter \\ NO_2 \rightarrow NO \end{array} \tag{3}$$

The nitrogen monoxide is then analysed according to the reactions (1) and (2).

The concentrations of nitrogen dioxide and nitrogen monoxide are expressed in the units provided by the calibration standard. The final results for reporting are expressed in $\mu g/m^3$ using standard conversion factors (see Clause 10).

5.3 Type testing

The type test is based on the evaluation of performance characteristics determined under a prescribed series of tests. In this document, test procedures are described for the determination of the actual values of the performance characteristics for at least two analysers in a laboratory and the same analysers in the field, operated in parallel in both cases. The laboratory tests shall not include the sampling inlet, sampling system and external data acquisition system, but shall include analyser sampling line and filter. The field tests may include a sampling inlet and a sampling system. However, the influence of these components on the test results shall be minimized by proper maintenance.

The performance test shall be performed by a competent body. The evaluation for type testing of an analyser is based on the calculation of the expanded uncertainty in the measuring result based on the numerical values of the tested performance characteristics and compared with a prescribed maximum uncertainty.

The type testing of an analyser and subsequent QA and QC procedures provide evidence that the defined requirements concerning data quality laid out in Annex I of Directive 2008/50/EC [1] can be satisfied.

Appropriate experimental evidence shall be provided by:

- type tests performed under conditions of intended use of the specified method of measurement, and
- calculation of expanded uncertainty of results of measurement by reference to JCGM 100 [3].

5.4 Field operation and quality control

Prior to the installation and operation of a type-tested analyser at a monitoring station, an expanded uncertainty calculation shall be performed with the actual values of the performance, obtained during the type tests, and the site-specific conditions at that monitoring station. This calculation shall be used