
**Natural gas — Calculation of
compression factor —**

**Part 3:
Calculation using physical properties**

*Gaz naturel — Calcul du facteur de compression —
Partie 3: Calcul à partir des caractéristiques physiques*
(standards.iteh.ai)

[ISO 12213-3:2006](https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006)

<https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006>



PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.

iTeh STANDARD PREVIEW
(standards.iteh.ai)

ISO 12213-3:2006

<https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006>

© ISO 2006

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Published in Switzerland

Contents

Page

| | |
|---|-----------|
| Foreword..... | iv |
| 1 Scope | 1 |
| 2 Normative references | 1 |
| 3 Terms and definitions..... | 1 |
| 4 Method of calculation | 2 |
| 4.1 Principle | 2 |
| 4.2 The SGERG-88 equation | 2 |
| 4.3 Input variables..... | 3 |
| 4.4 Ranges of application..... | 3 |
| 4.5 Uncertainty | 5 |
| 5 Computer program | 6 |
| Annex A (normative) Symbols and units | 7 |
| Annex B (normative) Description of the SGERG-88 method | 10 |
| Annex C (normative) Example calculations | 21 |
| Annex D (normative) Conversion factors | 22 |
| Annex E (informative) Specification for pipeline quality natural gas | 25 |
| Annex F (informative) Performance over wider ranges of application | 28 |
| Annex G (informative) Subroutine SGERG.FOR in Fortran | 33 |
| Bibliography | 38 |

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.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. 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.

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.

ISO 12213-3 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 12213-3:1997), which has been technically revised. The revision includes changes to Subclause 4.4.1 and the addition of a new annex, Annex E.

ISO 12213 consists of the following parts, under the general title *Natural gas — Calculation of compression factor*:

- *Part 1: Introduction and guidelines*
- *Part 2: Calculation using molar-composition analysis*
- *Part 3: Calculation using physical properties*

<https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006>

Natural gas — Calculation of compression factor —

Part 3: Calculation using physical properties

1 Scope

ISO 12213 specifies methods for the calculation of compression factors of natural gases, natural gases containing a synthetic admixture and similar mixtures at conditions under which the mixture can exist only as a gas.

This part of ISO 12213 specifies a method for the calculation of compression factors when the superior calorific value, relative density and carbon dioxide content are known, together with the relevant pressures and temperatures. If hydrogen is present, as is often the case for gases with a synthetic admixture, the hydrogen content also needs to be known.

NOTE In principle, it is possible to calculate the compression factor when any three of the parameters superior calorific value, relative density, carbon dioxide content (the usual three) and nitrogen content are known, but subsets including nitrogen content are not recommended.

The method is primarily applicable to pipeline quality gases within the ranges of pressure p and temperature T at which transmission and distribution operations normally take place, with an uncertainty of about $\pm 0,1\%$. For wider-ranging applications the uncertainty of the results increases (see Annex F).

More detail concerning the scope and field of application of the method is given in ISO 12213-1.

2 Normative references

The following referenced documents are indispensable for the application 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 6976:1995, *Natural gas — Calculation of calorific values, density, relative density and Wobbe index from composition*

ISO 12213-1, *Natural gas — Calculation of compression factor — Part 1: Introduction and guidelines*

ISO 80000-4, *Quantities and units — Part 4: Mechanics*

ISO 80000-5, *Quantities and units — Part 5: Thermodynamics*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 12213-1 apply.

4 Method of calculation

4.1 Principle

The method recommended uses equations which are based on the concept that pipeline quality natural gas may be uniquely characterized for calculation of its volumetric properties by an appropriate and distinctive set of measurable physical properties. These characteristics, together with the pressure and temperature, are used as input data for the method.

The method uses the following physical properties: superior calorific value, relative density and carbon dioxide content. The method is particularly useful in the common situation where a complete molar composition is not available, but may also be preferred for its relative simplicity. For gases with a synthetic admixture, the hydrogen content needs to be known.

4.2 The SGERG-88 equation

The calculation method using physical properties is based on the standard GERG 88 (SGERG-88) virial equation for natural gases [1], [2], [3]. The standard GERG 88 virial equation is derived from the master GERG 88 (MGERG-88) virial equation, which is a method of calculation based on a molar-composition analysis [4].

The SGERG-88 virial equation from which the compression factor Z is calculated may be written as

$$Z = 1 + B\rho_m + C\rho_m^2 \tag{1}$$

where

iTeh STANDARD PREVIEW
(standards.iteh.ai)

B and C are functions of the input data comprising the superior calorific value H_S , the relative density d , the contents of both inert and combustible non-hydrocarbon components of the gas mixture (CO_2 and H_2) and the temperature T ; [ISO 12213-3:2006](https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006)

ρ_m is the molar density given by [674e9ec46872/iso-12213-3-2006](https://standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006)

$$\rho_m = p/(ZRT) \tag{2}$$

where

$$Z = f_1(p, T, H_S, d, x_{CO_2}, x_{H_2}) \tag{3}$$

However, the SGERG-88 method treats the natural-gas mixture internally as a five-component mixture consisting of an equivalent hydrocarbon gas (with the same thermodynamic properties as the sum of the hydrocarbons present), nitrogen, carbon dioxide, hydrogen and carbon monoxide. To characterize the thermodynamic properties of the hydrocarbon gas adequately, the hydrocarbon heating value H_{CH} is also needed. Therefore, the calculation of Z uses

$$Z = f_2(p, T, H_{CH}, x_{CH}, x_{N_2}, x_{CO_2}, x_{H_2}, x_{CO}) \tag{4}$$

In order to be able to model coke oven gas mixtures, the mole fraction of carbon monoxide is taken to have a fixed relation to the hydrogen content. If hydrogen is not present ($x_{H_2} < 0,001$), then set $x_{H_2} = 0$. The natural-gas mixture is then treated in the calculation method as a three-component mixture (see Annex B).

The calculation is performed in three steps:

First, the five-component composition from which both the known superior calorific value and the known relative density can be calculated satisfactorily may be found from the input data by an iterative procedure described in detail in Annex B.

Secondly, once this composition is known, B and C may be found using relationships also given in Annex B.

In the third step, Equations (1) and (2) are solved simultaneously for ρ_m and Z by a suitable numerical method.

A flow diagram of the procedure for calculating Z from the input data is shown in Figure B.1.

4.3 Input variables

4.3.1 Preferred input data set

The input variables required for use with the SGERG-88 equation are the absolute pressure, temperature and superior calorific value (volumetric basis), the relative density, the carbon dioxide content and the hydrogen content. Thus the physical properties used in the input data set (set A) are

$$H_S, d, x_{CO_2} \text{ and } x_{H_2}$$

Relative density is referred to normal conditions (101,325 kPa and 0 °C) and superior calorific value is referred to normal conditions (101,325 kPa and 0 °C) and a combustion temperature of 25 °C.

4.3.2 Alternative input data sets

Three alternatives to the preferred input data set (see 4.3.1) may be used with the standard GERG 88 virial equation:

$$x_{N_2}, H_S, d \text{ and } x_{H_2} \text{ (set B)}$$

$$x_{N_2}, x_{CO_2}, d \text{ and } x_{H_2} \text{ (set C)}$$

$$x_{N_2}, x_{CO_2}, H_S \text{ and } x_{H_2} \text{ (set D)}$$

The alternative input data sets are considered fully in GERG Technical Monograph TM5^[3]. Use of the alternative input data sets gives results which may differ at the fourth decimal place. This part of ISO 12213 recommends the use of input data set A.

4.4 Ranges of application

4.4.1 Pipeline quality gas

The ranges of application for pipeline quality gas are as defined below:

| | | | |
|---------------------------------|-----------------------|-----------------|------------------------------|
| absolute pressure | 0 MPa | $\leq p$ | ≤ 12 MPa |
| temperature | 263 K | $\leq T$ | ≤ 338 K |
| mole fraction of carbon dioxide | 0 | $\leq x_{CO_2}$ | $\leq 0,20$ |
| mole fraction of hydrogen | 0 | $\leq x_{H_2}$ | $\leq 0,10$ |
| superior calorific value | 30 MJ·m ⁻³ | $\leq H_S$ | ≤ 45 MJ·m ⁻³ |
| relative density | 0,55 | $\leq d$ | $\leq 0,80$ |

The mole fractions of other natural-gas components are not required as input. These mole fractions shall, however, lie within the following ranges (the ratio of successive mole fractions in the alkane homologous series is typically 3:1 — see Annex E):

| | | | |
|----------|-----|-----------------|-------------|
| methane | 0,7 | $\leq x_{CH_4}$ | $\leq 1,0$ |
| nitrogen | 0 | $\leq x_{N_2}$ | $\leq 0,20$ |

| | |
|----------------------------------|-----------------------------------|
| ethane | $0 \leq x_{C_2H_6} \leq 0,10$ |
| propane | $0 \leq x_{C_3H_8} \leq 0,035$ |
| butanes | $0 \leq x_{C_4H_{10}} \leq 0,015$ |
| pentanes | $0 \leq x_{C_5H_{12}} \leq 0,005$ |
| hexanes | $0 \leq x_{C_6} \leq 0,001$ |
| heptanes | $0 \leq x_{C_7} \leq 0,000\ 5$ |
| octanes plus higher hydrocarbons | $0 \leq x_{C_{8+}} \leq 0,000\ 5$ |
| carbon monoxide | $0 \leq x_{CO} \leq 0,03$ |
| helium | $0 \leq x_{He} \leq 0,005$ |
| water | $0 \leq x_{H_2O} \leq 0,000\ 15$ |

The method applies only to mixtures in the single-phase gaseous state (above the dew point) at the conditions of temperature and pressure of interest. For pipeline quality gas, the method is applicable over wider ranges of temperature and pressure but with increased uncertainty (see Figure 1). In the computer implementation, the lower temperature limit is set at 250 K.

4.4.2 Wider ranges of application

iTech STANDARD PREVIEW
(standards.iteh.ai)

The ranges of application tested beyond the limits given in 4.4.1 are:

| | |
|---------------------------------|---|
| absolute pressure | $0\text{ MPa} \leq p \leq 12\text{ MPa}$ |
| temperature | $263\text{ K} \leq T \leq 338\text{ K}$ |
| mole fraction of carbon dioxide | $0 \leq x_{CO_2} \leq 0,30$ |
| mole fraction of hydrogen | $0 \leq x_{H_2} \leq 0,10$ |
| superior calorific value | $20\text{ MJ}\cdot\text{m}^{-3} \leq H_S \leq 48\text{ MJ}\cdot\text{m}^{-3}$ |
| relative density | $0,55 \leq d \leq 0,90$ |

The allowable mole fractions of other major natural-gas components are extended to:

| | |
|----------|-------------------------------|
| methane | $0,5 \leq x_{CH_4} \leq 1,0$ |
| nitrogen | $0 \leq x_{N_2} \leq 0,50$ |
| ethane | $0 \leq x_{C_2H_6} \leq 0,20$ |
| propane | $0 \leq x_{C_3H_8} \leq 0,05$ |

The limits for other minor natural-gas components remain as given in 4.4.1 for pipeline quality gas.

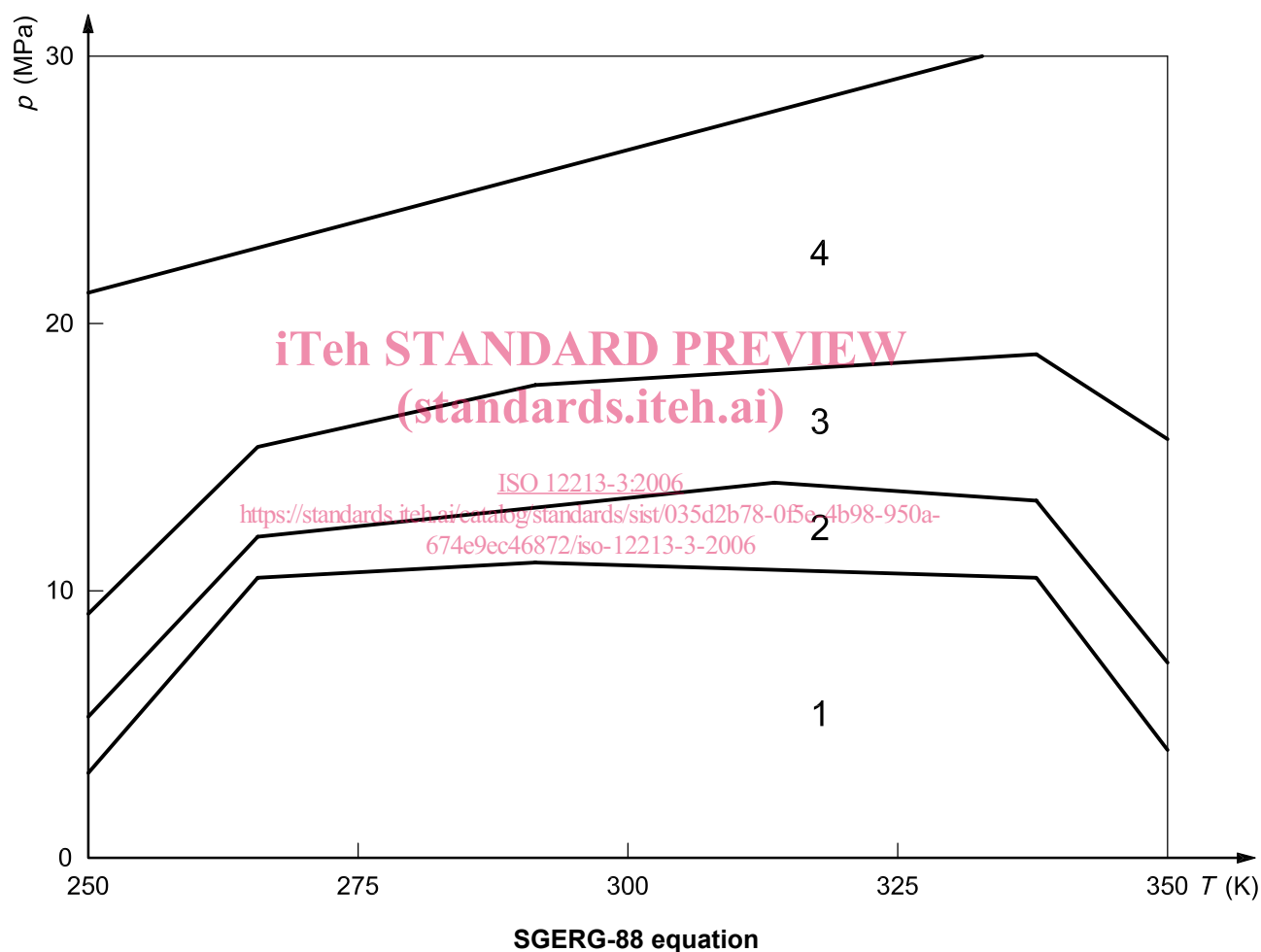
The method is not applicable outside these ranges; the computer implementation described in Annex B will not allow violation of the limits of composition quoted here.

4.5 Uncertainty

4.5.1 Uncertainty for pipeline quality gas

The uncertainty in the prediction of the compression factor ΔZ (for the temperature range 263 K to 338 K) is $\pm 0,1\%$ at pressures up to 10 MPa and $\pm 0,2\%$ between 10 MPa and 12 MPa for natural gases with $x_{N_2} \leq 0,20$, $x_{CO_2} \leq 0,09$, $x_{C_2H_6} \leq 0,10$ and $x_{H_2} \leq 0,10$, and for $30 \text{ MJ}\cdot\text{m}^{-3} \leq H_S \leq 45 \text{ MJ}\cdot\text{m}^{-3}$ and $0,55 \leq d \leq 0,80$ (see Figure 1).

For gases with a CO_2 content exceeding a mole fraction of 0,09, the uncertainty of $\pm 0,1\%$ is maintained for pressures up to 6 MPa and for temperatures between 263 K and 338 K. This uncertainty level is determined by comparison with the GERG databank on measurements of the compression factor for natural gases [5], [6] and with the Gas Research Institute data [9].



Key

p pressure

T temperature

1 $\Delta Z \leq \pm 0,1\%$

2 $\Delta Z \pm 0,1\%$ to $\pm 0,2\%$

3 $\Delta Z \pm 0,2\%$ to $\pm 0,5\%$

4 $\Delta Z \pm 0,5\%$ to $\pm 3,0\%$

Figure 1 — Uncertainty limits for the calculation of compression factors

(The uncertainty limits given are expected to be valid for natural gases and similar gases with $x_{N_2} \leq 0,20$; $x_{CO_2} \leq 0,09$; $x_{C_2H_6} \leq 0,10$ and $x_{H_2} \leq 0,10$, and for $30 \text{ MJ}\cdot\text{m}^{-3} \leq H_S \leq 45 \text{ MJ}\cdot\text{m}^{-3}$ and $0,55 \leq d \leq 0,80$)

4.5.2 Uncertainty for wider ranges of application

The estimated uncertainties involved in calculations of compression factors beyond the limits of quality given in 4.5.1 are discussed in Annex F.

4.5.3 Impact of uncertainties of input variables

Listed in Table 1 are typical values for the uncertainties of the relevant input variables. These values may be achieved under optimum operating conditions.

As a general guideline only, an error propagation analysis using the above uncertainties in the input variables produces an additional uncertainty of about $\pm 0,1\%$ in the result at 6 MPa and within the temperature range 263 K to 338 K. Above 6 MPa, the additional uncertainties are greater and increase roughly in direct proportion to the pressure (see Reference [3]).

4.5.4 Reporting of results

Results for the compression factor shall be reported to four places of decimals, together with the pressure and temperature values and the calculation method used (ISO 12213-3, SGERG-88 equation). For verification of calculation procedures, it is useful to carry extra digits.

Table 1 — Uncertainties of input variables

| Input variable | Absolute uncertainty |
|---------------------------------|-------------------------------|
| Absolute pressure | $\pm 0,02$ MPa |
| Temperature | $\pm 0,15$ K |
| Mole fraction of carbon dioxide | $\pm 0,002$ |
| Mole fraction of hydrogen | $\pm 0,005$ |
| Relative density | $\pm 0,0013$ |
| Superior calorific value | $\pm 0,06$ MJ·m ⁻³ |

5 Computer program

Software which implements this International Standard has been prepared. Users of this part of ISO 12213 are invited to contact ISO/TC 193/SC 1, either directly or through their ISO member body, to enquire about the availability of this software.

Annex A (normative)

Symbols and units

The symbols specified in this annex are those which are used in both the main text and in Annex B. The units specified here are those which give consistency with the values of the coefficients given in Annex B.

| Symbol | Meaning | Units |
|-------------|--|---|
| b_{H0} | Zero-order (constant) term in the molar heating value (H_{CH}) expansion of B_{11} [Equation (B.20)] | $\text{m}^3 \cdot \text{kmol}^{-1}$ |
| b_{H1} | First-order (linear) term in the molar heating value (H_{CH}) expansion of B_{11} [Equation (B.20)] | $\text{m}^3 \cdot \text{MJ}^{-1}$ |
| b_{H2} | Second-order (quadratic) term in the molar heating value (H_{CH}) expansion of B_{11} [Equation (B.20)] | $\text{m}^3 \cdot \text{kmol} \cdot \text{MJ}^{-2}$ |
| $b_{H0(0)}$ | Terms in the temperature expansion of b_{H0} [Equation (B.21)] | $\text{m}^3 \cdot \text{kmol}^{-1}$ |
| $b_{H0(1)}$ | | $\text{m}^3 \cdot \text{kmol}^{-1} \cdot \text{K}^{-1}$ |
| $b_{H0(2)}$ | | $\text{m}^3 \cdot \text{kmol}^{-1} \cdot \text{K}^{-2}$ |
| $b_{H1(0)}$ | Terms in the temperature expansion of b_{H1} [Equation (B.21)] | $\text{m}^3 \cdot \text{MJ}^{-1}$ |
| $b_{H1(1)}$ | | $\text{m}^3 \cdot \text{MJ}^{-1} \cdot \text{K}^{-1}$ |
| $b_{H1(2)}$ | | $\text{m}^3 \cdot \text{MJ}^{-1} \cdot \text{K}^{-2}$ |
| $b_{H2(0)}$ | Terms in the temperature expansion of b_{H2} [Equation (B.21)] | $\text{m}^3 \cdot \text{kmol} \cdot \text{MJ}^{-2}$ |
| $b_{H2(1)}$ | | $\text{m}^3 \cdot \text{kmol} \cdot \text{MJ}^{-2} \cdot \text{K}^{-1}$ |
| $b_{H2(2)}$ | | $\text{m}^3 \cdot \text{kmol} \cdot \text{MJ}^{-2} \cdot \text{K}^{-2}$ |
| $b_{ij(0)}$ | Terms in the temperature expansion of b_{ij} [Equation (B.22)] | $\text{m}^3 \cdot \text{kmol}^{-1}$ |
| $b_{ij(1)}$ | | $\text{m}^3 \cdot \text{kmol}^{-1} \cdot \text{K}^{-1}$ |
| $b_{ij(2)}$ | | $\text{m}^3 \cdot \text{kmol}^{-1} \cdot \text{K}^{-2}$ |
| B | Second virial coefficient [Equation (1)] | $\text{m}^3 \cdot \text{kmol}^{-1}$ |
| B_{ij} | Second virial coefficient for binary interaction between component i and component j [Equation (B.22)] | $\text{m}^3 \cdot \text{kmol}^{-1}$ |
| c_{H0} | Zero-order (constant) term in the molar heating value (H_{CH}) expansion of C_{111} [Equation (B.29)] | $\text{m}^6 \cdot \text{kmol}^{-2}$ |
| c_{H1} | First-order (linear) term in the molar heating value (H_{CH}) expansion of C_{111} [Equation (B.29)] | $\text{m}^6 \cdot \text{kmol}^{-1} \cdot \text{MJ}^{-1}$ |
| c_{H2} | Second-order (quadratic) term in the molar heating value (H_{CH}) expansion of C_{111} [Equation (B.29)] | $\text{m}^6 \cdot \text{MJ}^{-2}$ |

| Symbol | Meaning | Units |
|--|---|--|
| $c_{H0}(0)$ $c_{H0}(1)$ $c_{H0}(2)$ | Terms in the temperature expansion of c_{H0} [Equation (B.30)] | $m^6 \cdot kmol^{-2}$ |
| | | $m^6 \cdot kmol^{-2} \cdot K^{-1}$ |
| | | $m^6 \cdot kmol^{-2} \cdot K^{-2}$ |
| $c_{H1}(0)$ $c_{H1}(1)$ $c_{H1}(2)$ | Terms in the temperature expansion of c_{H1} [Equation (B.30)] | $m^6 \cdot kmol^{-1} \cdot MJ^{-1}$ |
| | | $m^6 \cdot kmol^{-1} \cdot MJ^{-1} \cdot K^{-1}$ |
| | | $m^6 \cdot kmol^{-1} \cdot MJ^{-1} \cdot K^{-2}$ |
| $c_{H2}(0)$ $c_{H2}(1)$ $c_{H2}(2)$ | Terms in the temperature expansion of c_{H2} [Equation (B.30)] | $m^6 \cdot MJ^{-2}$ |
| | | $m^6 \cdot MJ^{-2} \cdot K^{-1}$ |
| | | $m^6 \cdot MJ^{-2} \cdot K^{-2}$ |
| $c_{ijk}(0)$ $c_{ijk}(1)$ $c_{ijk}(2)$ | Terms in the temperature expansion of c_{ijk} [Equation (B.31)] | $m^6 \cdot kmol^{-2}$ |
| | | $m^6 \cdot kmol^{-2} \cdot K^{-1}$ |
| | | $m^6 \cdot kmol^{-2} \cdot K^{-2}$ |
| C | Third virial coefficient [Equation (1)] | $m^6 \cdot kmol^{-2}$ |
| C_{ijk} | Third virial coefficient for ternary interaction between components i, j and k [Equation (B.31)] | $m^6 \cdot kmol^{-2}$ |
| d | Relative density [$d(\text{air}) = 1$; Equation (B.1)] | — |
| DH_{CH} | Change in the molar heating value H_{CH} during iteration [Equations (B.10) and (B.11)] | $MJ \cdot kmol^{-1}$ |
| H_S | Superior calorific value [gas at normal conditions (0 °C, 1,013 25 bar), combustion temperature 25 °C] | $MJ \cdot m^{-3}$ |
| H | Molar heating value (combustion temperature 25 °C) | $MJ \cdot kmol^{-1}$ |
| M | Molar mass [Equations (B.5) and (B.8)] | $kg \cdot kmol^{-1}$ |
| p | Absolute pressure | bar |
| R | (Universal) gas constant | $m^3 \cdot bar \cdot kmol^{-1} \cdot K^{-1}$ |
| T | Absolute temperature | K |
| t | Celsius temperature [= $T - 273,15$; Equation (B.27)] | °C |
| V_m | Molar volume (= $1/\rho_m$) | $m^3 \cdot kmol^{-1}$ |
| x | Mole fraction of a component | — |
| y | Combination rule parameters for the binary unlike-interaction virial coefficients B_{12} and B_{13} (Table B.2) and the ternary unlike-interaction virial coefficient C_{ijk} [Equation (B.32)] | — |
| Z | Compression factor | — |
| ρ | Mass density [Equations (B.8) and (B.42)] | $kg \cdot m^{-3}$ |
| ρ_m | Molar density (= V_m^{-1}) | $kmol \cdot m^{-3}$ |

Additional subscripts

| | |
|-----------------|--|
| n | Value at normal conditions ($T_n = 273,15 \text{ K}$, $p_n = 1,013 25 \text{ bar}$) |
| CH | For the equivalent hydrocarbon |
| CO | For carbon monoxide |
| CO ₂ | For carbon dioxide |
| H ₂ | For hydrogen |
| N ₂ | For nitrogen |

Additional qualifiers

| | |
|---------|---|
| (air) | For dry air of standard composition [Equation (B.1)] |
| (D) | For special value of ρ used in Equation (B.11) |
| 1 | For the equivalent hydrocarbon [Equations (B.12) and (B.15)] |
| 2 | For nitrogen [Equations (B.12) and (B.16)] |
| 3 | For carbon dioxide [Equations (B.12) and (B.17)] |
| 4 | For hydrogen [Equations (B.12) and (B.18)] |
| 5 | For carbon monoxide [Equations (B.12) and (B.19)] |
| (id) | Ideal gas state standards.iteh.ai/catalog/standards/sist/035d2b78-0f5e-4b98-950a-674e9ec46872/iso-12213-3-2006 |
| (u) | Iteration counter (B.2.1) |
| (v) | Iteration counter (B.2.2) |
| (w) | Iteration counter (B.4) |