
Capability of detection —
Part 7:
Methodology based on stochastic
properties of instrumental noise

Capacité de détection —

Partie 7: Méthodologie basée sur les propriétés stochastiques du bruit instrumental

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Contents

	Page
Foreword.....	iv
Introduction.....	v
1 Scope	1
2 Normative references	1
3 Terms and definitions	2
4 Quantitative analysis and background noise	3
4.1 Error sources of analysis.....	3
4.2 Random processes in background.....	3
5 Theories for precision	4
5.1 Theory based on auto-covariance function.....	4
5.2 Theory based on power spectrum.....	6
6 Practical use of FUMI theory	9
6.1 Estimation of noise parameters by Fourier transform.....	9
6.2 Estimation of noise parameters by autocovariance function.....	11
6.3 Procedures for estimation of SD.....	11
Annex A (informative) Symbols and abbreviated terms used in this document	14
Annex B (informative) Derivation of Formula (7)	15
Annex C (informative) Derivation of Formulae (14) to (16)	16
Bibliography	18

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 of 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 www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 69, *Applications of statistical methods*, Subcommittee SC 6, *Measurement methods and results*.

This second edition cancels and replaces the first edition (ISO 11843-7:2012), which has been technically revised.

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

- created a new [6.2](#);
- [6.2](#) of the first edition is renumbered [6.3](#).

A list of all parts in the ISO 11843 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

The series of ISO 11843 is based on the probability distributions of the net state variable (measurand) for both the linear and nonlinear calibration situations. The focus is implicitly, though sometimes explicitly, on the uncertainty associated with an estimate of the measured response predominantly coming from the baseline noise in instrumental analysis. In many, if not most, analytical instruments, the baseline noise is considered the prime cause of uncertainty when the sample amount is as low as the minimum detectable value. Within its domain of applicability, the method given in this document can dispense with the repetition of real samples, thus helping to improve global environments by saving time and energy that would be required by repetition.

The basic concept of ISO 11843-7 is the mathematical description of the probability distribution of the response variable in terms of mathematically well-defined random processes. This description straightforwardly leads to the minimum detectable value. As for the relation of the response and measurand, linear and nonlinear calibration functions can be applied. In this manner, compatibility with ISO 11843-2 and ISO 11843-5 is ensured.

The definition and applicability of the minimum detectable value are described in ISO 11843-1 and ISO 11843-2; the definition and applicability of the precision profile are described in ISO 11843-5. The precision profile expresses how the precision changes depending on the net state variable. ISO 11843-7 specifies the practical use of the fundamental concepts in ISO 11843 in case of the background noise predominance in instrumental analysis.

The minimum detectable value, x_d , is generally expressed in the unit of the net state variable. If the calibration function is linear, the SD or CV of the response variable estimated in this document can linearly be transformed to the SD or CV of the net state variable, which in turn can be used for the estimation of the minimum detectable value, x_d .

If the calibration function is nonlinear, the precision profile of the response variable in this document needs to be transformed to the precision profile of the net state variable as shown in ISO 11843-5. In this situation, the contents of ISO 11843-5 can be used for this purpose without modification.

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Capability of detection —

Part 7:

Methodology based on stochastic properties of instrumental noise

1 Scope

Background noise exists ubiquitously in analytical instruments, whether or not a sample is applied to the instrument. This document is concerned with mathematical methodologies for estimating the minimum detectable value in case that the most predominant source of measurement uncertainty is background noise. The minimum detectable value can directly and mathematically be derived from the stochastic characteristics of the background noise.

This document specifies basic methods to

- extract the stochastic properties of the background noise,
- use the stochastic properties to estimate the standard deviation (SD) or coefficient of variation (CV) of the response variable, and
- calculate the minimum detectable value based on the SD or CV obtained above.

The methods described in this document are useful for checking the detection of a certain substance by various types of measurement equipment in which the background noise of the instrumental output predominates over the other sources of measurement uncertainty. Feasible choices are visible and ultraviolet absorption spectrometry, atomic absorption spectrometry, atomic fluorescence spectrometry, luminescence spectrometry, liquid chromatography and gas chromatography.

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 3534-1, *Statistics — Vocabulary and symbols — Part 1: General statistical terms and terms used in probability*

ISO 3534-2, *Statistics — Vocabulary and symbols — Part 2: Applied statistics*

ISO 3534-3, *Statistics — Vocabulary and symbols — Part 3: Design of experiments*

ISO 5725-1, *Accuracy (trueness and precision) of measurement methods and results — Part 1: General principles and definitions*

ISO 11843-1, *Capability of detection — Part 1: Terms and definitions*

ISO 11843-2, *Capability of detection — Part 2: Methodology in the linear calibration case*

ISO 11843-5, *Capability of detection — Part 5: Methodology in the linear and non-linear calibration cases*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 3534-1, ISO 3534-2, ISO 3534-3, ISO 5725-1, ISO 11843-1, ISO 11843-2, ISO 11843-5 and the following apply. A list of symbols and abbreviated terms used in this document is provided in Annex A.

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/>

3.1
precision profile
<detection capability> mathematical description of the standard deviation (SD) of the response variable $[\sigma_Y(X)]$ or net state variable $[\sigma_X(X)]$ as a function of the net state variable

Note 1 to entry: The coefficient of variation (CV) of the response variable or net state variable as a function of the net state variable is also referred to as a precision profile.

Note 2 to entry: Precision means the SD or CV of the observed response variable or SD or CV of the net state variable when estimated by the calibration function (see ISO 11843-5).

[SOURCE: ISO 11843-5:2008, 3.4, modified — “coefficient of variation” has been removed and Note 1 to entry has been added instead. Note 2 to entry has also been added.]

3.2
minimum detectable value of the net state variable

x_d
value of the net state variable in the actual state that will lead, with probability $1 - \beta$, to the conclusion that the system is not in the basic state

Note 1 to entry: Under the assumption that the SD, $\sigma_X(X)$, of the net state variable is constant $[(\sigma_X(X) = \sigma_X)]$, the minimum detectable value, x_d , is defined as

$$x_d = (k_c + k_d) \sigma_X \quad (1)$$

where

k_c denotes a coefficient to specify the probability of an error of the first kind;

k_d is a coefficient to specify the probability of an error of the second kind.

If the SD, σ_Y , of the response variable is assumed to be constant $[\sigma_Y(X) = \sigma_Y]$, then the minimum detectable value can be calculated by the following Formula (2):

$$x_d = (k_c + k_d) (\sigma_Y / |dY / dX|) \quad (2)$$

where $|dY/dX|$ denotes the absolute value of the slope of the linear calibration function and is constant.

Note 2 to entry: If the net state variable is normally distributed, the coefficients $k_c = k_d = 1,65$ specify the probabilities of an error of the first and second kinds (= 5 %) and Formula (1) can simply be written as $x_d = 3,30 \sigma_X$.

Note 3 to entry: If $k_c = k_d = 1,65$, Formula (1) takes the form that $\sigma_X / x_d = 1/3,30 = 30$ %. Therefore, x_d can be found in the *precision profile* (3.1). x_d is located at X , the CV of which is 30 %.

Note 4 to entry: Different types of *precision profiles* (3.1) are defined, but they can be transformed to each other.

For example, the SD, $\sigma_Y(X)$, of the response variable can be transformed to the SD, $\sigma_X(X)$, of the net state variable by means of the absolute value of the derivative, $|dY/dX|$, of the calibration function $[Y = f(X)]$: $\sigma_X(X) = [\sigma_Y(X) / |dY/dX|]$ (see ISO 11843-5). This treatment is an approximation, the extent of which depends on local curvature, involving d^2Y/dX^2 .

[SOURCE: ISO 11843-5:2008, 3.2, modified — Notes to entry 1 to 4 have been added.]

4 Quantitative analysis and background noise

4.1 Error sources of analysis

The quantitative analysis to obtain a measurand from a sample is generally considered to consist of preparation, instrumental analysis, data handling and calibration. These steps of analysis are mechanically independent of each other and so are probabilistically independent as well.

This document applies only to instrumental analysis. However, the errors from the other steps affect the error of the final value of the measurand, as well. That is, the combined uncertainty associated with an estimate of the measurand depends on the propagation of all uncertainties relating to the relevant steps. The following conditions are necessary for the use of ISO 11843-7.

At concentrations near the minimum detectable value in chromatography, the error from the sample injection into a chromatograph is even less important (e.g. CV = 0,3 % in a recent apparatus) than the background noise (CV = 30 % by definition). If the importance of a factor other than noise is comparable to that of the noise, the methodologies of this document are not applicable.

Data handling is usually a process to extract a signal component from noisy instrumental output such as peak height or area, which is a relative height of a summit of a peak-shaped signal or integration of intensities over a signal region, respectively. The statistical influences of this process are the major concern of this document. The use of a digital or analogue filter can also be taken into account, if the noise after the filtration is analysed for this purpose.

4.2 Random processes in background

Typical examples of the response variable are area and height measured in chromatography. In this document, intensity difference [Formula (6)] and area [Formulae (10) and (11)] are taken as the difference and summation of intensities Y_i of instrumental output. The response variables are usually independent of each other even if they are obtained from consecutive measurement by the same instrument. On the other hand, the consecutive intensities Y_i are formulated as a time-dependent random process, and in many cases, can be considered $1/f$ noise [1].

The power spectrum, $P(f)$, of $1/f$ noise has a slope inversely proportional to frequency, f :

$$P(f) \propto \frac{1}{f} \quad (3)$$

when f is near zero.

The simplest model of random processes is the white noise. Let w_i denote the random variable of the white noise at point i . By definition, the mean of the white noise is zero and the SD, $\tilde{w}$, of the white noise is constant at every point i . A prominent feature of the white noise is that the noise intensities, w_i and w_j , are independent of each other, if $i \neq j$.

The autoregressive process [AR(1)-process] of first order is a mathematical model in which the intensities, M_i and M_j , are not independent of each other ($i \neq j$). The AR(1)-process is treated as a major component of time-dependent changes of instrumental output [see Formula (9)]. The AR(1)-process at point i is defined to take the form

$$M_i = \rho M_{i-1} + m_i \quad (4)$$

where

m_i denotes the random variable of the white noise at point i ;

ρ is a constant parameter ($-1 < \rho < 1$).

5 Theories for precision

5.1 Theory based on auto-covariance function

A theory has been proposed [2][3][4] based on an auto-covariance function:

$$\psi(\tau_s) \equiv E[Y_{t_0+\tau_s} Y_{t_0}] \quad (5)$$

where $E[\cdot]$ denotes the mean of a random variable inside the square brackets over t_0 .

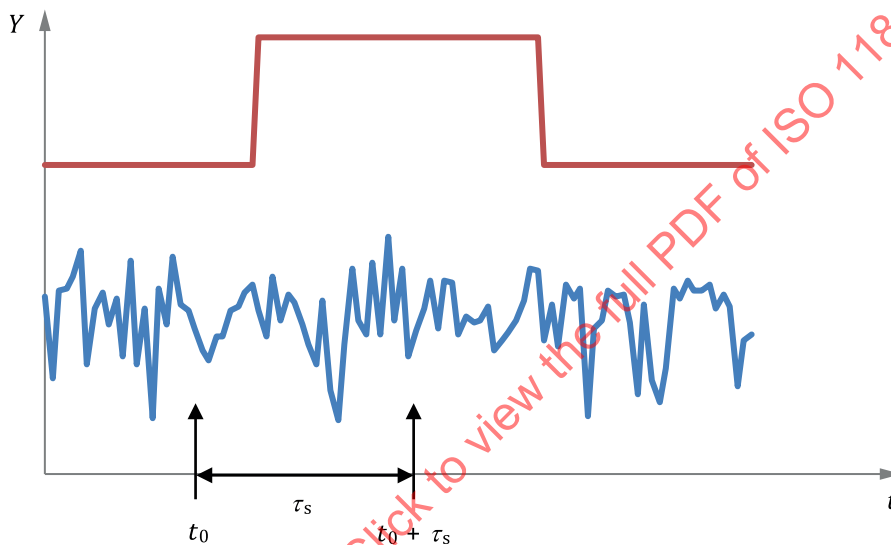


Figure 1 — Signal (upper line) and noise (lower line) with intensity difference

The upper part of [Figure 1](#) depicts the signal as (an approximation to) a rectangular pulse. Noise (constituting background) on the signal is depicted as the oscillatory curve in the lower part of the figure. t_0 denotes a time value on the background portion of the signal and $t_0 + \tau_s$ denotes a time value on the signal itself. The measurement (signal reading) is the difference in intensities at times t_0 and $t_0 + \tau_s$. The value of the signal would be zero at t_0 in the absence of background noise. The signal has a finite value at $t_0 + \tau_s$ when a sample is measured. In the ISO 11843-7 measurement model, the signal and noise are superimposed, and this mixed random process takes the value Y_i at time t_i . The intensities at times t_0 and $t_0 + \tau_s$ are described as Y_{t_0} and $Y_{t_0+\tau_s}$, respectively, and the intensity difference is given by [Formula \(6\)](#).

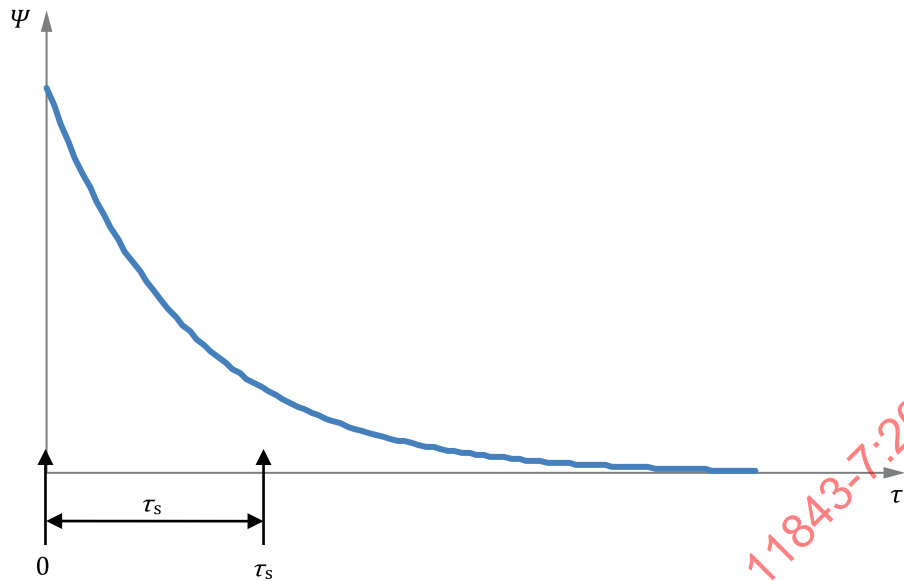


Figure 2 — Auto-covariance function of noise

The difference between the values of the auto-covariance function, $\psi(\tau)$, at 0 and τ_s gives the right side of [Formula \(7\)](#).

Near the minimum detectable value, which is dictated by the background fluctuation, intensity difference often applies in analytical optical spectrometry. The definition of intensity difference, e.g. signal reading corrected for background, is [\[2\]\[3\]\[4\]](#):

$$\Delta Y = Y_{t_0 + \tau_s} - Y_{t_0} \quad (6)$$

Here, ΔY corresponds to the response variable Y . The variance of the intensity difference is written as shown in [Formula \(7\)](#) [\[2\]\[3\]\[4\]](#). [For the derivation of [Formula \(7\)](#), see [Annex B](#).]

$$\sigma_{\Delta Y}^2 = 2[\psi(0) - \psi(\tau_s)] \quad (7)$$

[Formula \(7\)](#) is of practical use when the actual auto-covariance functions, $\psi(0)$ and $\psi(\tau_s)$, are known from the observation of background noise as shown in [Figure 2](#). The substitution of [Formula \(7\)](#) for [Formula \(2\)](#) ($\sigma_Y = \sigma_{\Delta Y}$) leads to the minimum detectable value.

Use can be made of the Wiener-Khintchine theorem [\[5\]](#), which relates the auto-covariance function to the power spectral density through the Fourier transform:

$$\psi(\tau_s) = \int_0^\infty S_b(f) |G(f)|^2 \cos(2\pi f \tau_s) df \quad (8)$$

where

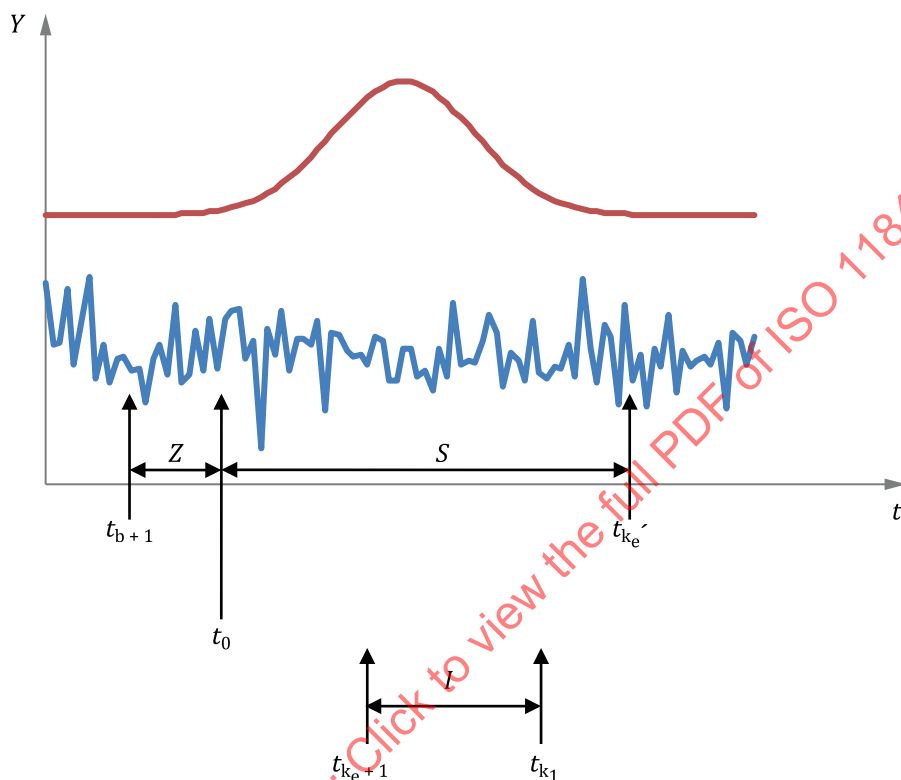
$S_b(f)$ denotes the power spectrum of the observed background noise;

$G(f)$ is the frequency response of the (linear) read-out system.

[Formula \(8\)](#) indicates the estimation of the measurement SD, [Formula \(7\)](#), through the noise power spectrum.

5.2 Theory based on power spectrum

A theory based on the power spectrum of the baseline, which is called FUMI (function of mutual information)[6][7][8], provides the SD values of the measured area and height in instrumental analysis. These measured values are the integration of the instrumental output over the integration region, as illustrated in Figure 3. If the signal (shape and size) is invariant, the error of measured area or height, as long as it comes from the noise alone, is equal to the area created by the noise over the integration region. That is, the measurement error is the same as the noise-created area. The SD of the noise-created areas coincides with the SD of measured heights or areas.



Key

- Z zero window
- S signal region
- I integration region

NOTE Additional symbols are explained in Annex A.

Figure 3 — Signal and noise with zero window and integration area

The number of data points over the integration region is $k_f - k_c$.

In the FUMI theory, the noise intensity, Y_i , at point i is described as the mixed random processes of the white noise and AR(1)-process:

$$Y_i = w_i + M_i \quad (9)$$

The purpose of the FUMI theory is to estimate the SD of the noise-created areas, A_F , over the integration region (see Figure 3).

In practice, especially chromatography, different modes of integration are adopted as illustrated in Figure 4. The measurement is of the integrated intensities above the baseline, which is horizontal or oblique within the domain $[k_c + 1, k_f]$. The horizontal baseline is horizontally drawn from the intensity (corrected) at the zero point, and the oblique line is drawn between the intensities at the edges of

the signal region $[0, k_e]$. The latter is often useful for a slowly changing background, called “drift” in chromatography.

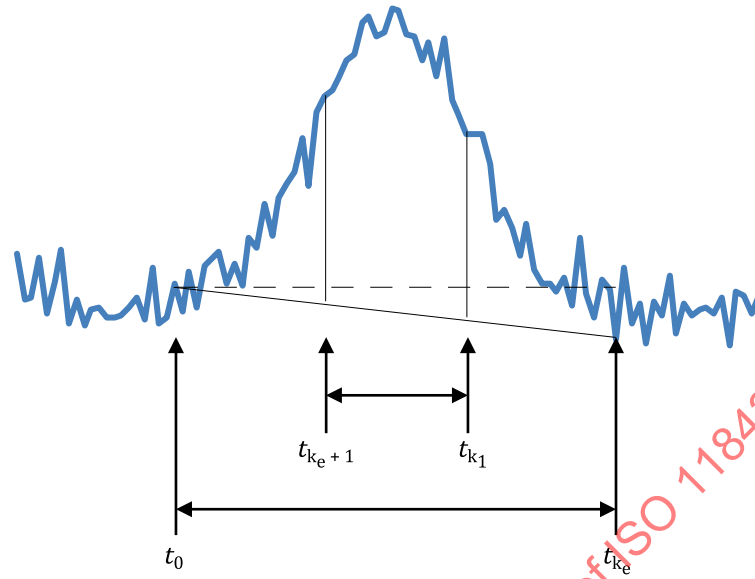


Figure 4 — Integration modes over horizontal line (dashed line) and oblique line (solid line) between t_{k_c+1} and t_{k_f}

The noise-created area, which is the area between the random path and horizontal line over the integration region without signal, can be written as

$$A_F = \sum_{i=k_c+1}^{k_f} Y_i \Delta t \quad (10)$$

where

Y_i the noise intensity is described by [Formula \(9\)](#);

Δt is the time interval between consecutive data.

Here, A_F means the response variable Y .

If the oblique line is used as the baseline, the noise-created area takes the form:

$$A_F = \left(\sum_{i=k_c+1}^{k_f} Y_i \Delta t \right) - A_T \quad (11)$$

where A_T denotes the area of the trapezoid created by the oblique line, horizontal line and vertical lines at the edges of the integration region from $k_c + 1$ to k_f (see [Figure 4](#)). The area of the trapezoid in [Figure 4](#) is taken with a negative sign. The area is taken with a positive sign if the oblique line lies above the horizontal line in the integration region and with a negative sign otherwise.

The general expression of the SD, σ_F , of noise-created areas is:

$$\sigma_F = \left(E \left[A_F^2 \right] \right)^{1/2} \quad (12)$$

where $E[\cdot]$ denotes the ensemble mean of a random variable inside the square brackets. It should be noted that $E[A_F] = 0$, since, by definition, the ensemble mean of the noise-created area over the

horizontal line [the first term in the right side of [Formula \(11\)](#)] is zero and the ensemble mean of the trapezoid, A_T , is zero (also see [Annex C](#)).

The above derivation is based on the assumption that there exists no uncertainty of the zero level, i.e. $Y_0 = 0$. In practice, however, this type of uncertainty exists and should be taken into account. The measurement in chromatography is usually performed relative to the zero level, which is the average of background intensities over a region referred to here as a zero window (see [Figure 3](#)).

The squared SD, σ_Y^2 , of measured areas (noise-created areas) within the zero window takes the form [\[6\]\[7\]\[8\]](#):

$$\sigma_Y^2 = \sigma_Z^2 + \sigma_F^2 \quad (13)$$

where σ_Z^2 denotes the variance originating from the zero window and σ_F^2 is the variance from the measured area [σ_F given by [Formula \(12\)](#)]. The summary of the derivation of [Formulae \(14\)](#) and [\(15\)](#) is given in [Annex C](#). The minimum detectable value can be obtained by the substitution of [Formula \(13\)](#) for [Formula \(2\)](#).

The variance, σ_Z^2 , can be described as [\[6\]\[7\]\[8\]](#)

$$\sigma_Z^2 = \frac{(k_f - k_c)^2}{b} \tilde{w}^2 + \frac{(k_f - k_c)^2}{b^2 (1 - \rho)^2} \left(b - 2\rho \frac{1 - \rho^b}{1 - \rho} + \rho^2 \frac{1 - \rho^{2b}}{1 - \rho^2} \right) \tilde{m}^2 \quad (14)$$

and the influence of the signal integration over the signal region takes the form [\[6\]\[7\]\[8\]](#):

$$\sigma_F^2 = (k_f - k_c) \tilde{w}^2 \quad (\text{first term, 15})$$

$$+ \frac{1}{(1 - \rho)^2} \left[k_f - k_c - 2\rho \frac{1 - \rho^{k_f - k_c}}{1 - \rho} + \rho^2 \frac{1 - \rho^{2(k_f - k_c)}}{1 - \rho^2} \right] \tilde{m}^2 \quad (\text{second term, 15})$$

$$+ \rho^2 \frac{1 - \rho^{2k_c}}{1 - \rho^2} \left(\frac{1 - \rho^{k_f - k_c}}{1 - \rho} \right)^2 \tilde{m}^2 \quad (\text{third term, 15})$$

$$+ a^2 \tilde{w}^2 \quad (\text{fourth term, 15})$$

$$+ \left\{ \alpha^2 \frac{1 - \rho^{2k_e}}{1 - \rho^2} - 2\alpha \left[\left(\frac{1 - \rho^{k_f - k_c}}{1 - \rho} \right) \rho^{k_e + k_c - 1} \left(\frac{1 - \rho^{-2k_c}}{1 - \rho^{-2}} \right) + \sum_{i=1}^{k_f - k_c} \left(\frac{1 - \rho^{k_f - k_c + 1 - i}}{1 - \rho} \right) \rho^{k_e - k_c - i} \right] \right\} \tilde{m}^2 \quad (\text{fifth term, 15})$$

where

$$\alpha = \frac{(k_f - k_c)(k_f + k_c + 1)}{2k_e} \quad (16)$$

- $\tilde{w}$ denotes the SD of the white noise;
- $\tilde{m}$ is the SD of the white noise included in the autoregressive process of first order;
- $\tilde{\rho}$ is the constant parameter of the autoregressive process of first order;
- k_c, k_f and k_e are defined in [Figures 3](#) and [4](#);
- b denotes the number of consecutive data points in the zero window $[-b + 1, 0]$.

The five terms in [Formula \(15\)](#) denote the following stochastic contributions to the measurement uncertainty:

- first term: the error from the white noise in the integration domain ($k_f - k_c$ data points);
- second term: the error from the AR(1)-process in the integration domain ($k_f - k_c$ data points);
- third term: the influence of k_c data points between the zero point and the starting point of integration;
- fourth term: the effect of the white noise in the oblique baseline;
- fifth term: the effect of the AR(1)-process in the oblique baseline.

NOTE The applicability of the FUMI theory is rather wide, but there are two typical situations where it is not.

One is that the predominant error source is not background noise. In mass spectroscopy, if the ionization process produces much more error than the noise, the FUMI theory underestimates the SD of measured areas.

The other situation is where actual instrumental noise includes the noise that cannot successfully be approximated by the mixed processes of the white noise and AR(1)-process. An example is spike noise of high intensity.

Baseline shifts of fixed patterns which are beyond the stationary assumption are often observable in gradient chromatography, but the integration mode of the oblique baseline can assure, though restricted, the robustness of the approach.

6 Practical use of FUMI theory

6.1 Estimation of noise parameters by Fourier transform

All the parameters necessary for applying the FUMI theory, i.e. [Formulae \(13\)](#) to [\(16\)](#), can be uniquely determined from the experimental data. The signal parameters (b, k_c, k_f, k_e) can be set according to the shape of a target peak, as shown in [Figure 3](#). On the other hand, the noise parameters ($\tilde{w}, \tilde{m}, \rho$) are automatically determined from the power spectral density of the noise, as described below.

The noise power spectral density results from the Fourier transform of noise data, Y_i . The Fourier and inverse Fourier transforms are:

$$\hat{Y}_k = \sum_{i=0}^{N-1} Y_i W^{ki} \quad (17)$$

$$Y_i = \frac{1}{N} \sum_{k=0}^{N-1} \hat{Y}_k W^{-ki} \quad (18)$$

where

N denotes the number of data points involved in the region of the Fourier transform;

$W = \exp[-j(2\pi/N)]$;

j is the imaginary unit.

The power spectral density, $P(k)$, of the random process, Y_i , is defined as

$$P(k) = \frac{\hat{Y}_k \bar{\hat{Y}}_k}{N} \quad (19)$$

where $\bar{\hat{Y}}_k$ is the conjugate number of $\hat{Y}_k$.

If the model noise described by [Formula \(9\)](#) is adopted, the power spectral density of [Formula \(19\)](#) can be described as [\[6\]\[7\]\[8\]](#)

$$P(k) = \frac{2\tilde{m}^2}{(1+\rho)\left[(1-\rho)^2 + 4\pi^2(k/N)^2\right]} + \tilde{w}^2 \quad (20)$$

The noise parameters ($\tilde{w}$, $\tilde{m}$ and $\tilde{\rho}$) needed for the FUMI theory can be determined by the nonlinear least squares fitting of [Formula \(20\)](#) to the power spectral density of the actual background noise.

The first term of the Fourier transform, $\hat{Y}_0$, included in [Formulae \(19\)](#) and [\(20\)](#), is the dc component, and the corresponding spectral density, $P(0)$, is neglected. The frequency, k , in [Formula \(20\)](#), which is the integer here, ranges from 1 to $N/2$ where $N/2$ is the Nyquist frequency. A typical example of [Formula \(20\)](#) is illustrated in [Figure 5](#).

In practice, frequency, k , is often expressed as Hertz. Let Δt be the time interval between consecutive data points (the intervals of an analogue-to-digital converter used for the acquisition of data, Y_i). The frequency which corresponds to k ($= 1, 2, \dots, N/2$) is $k/(\Delta t N)$.

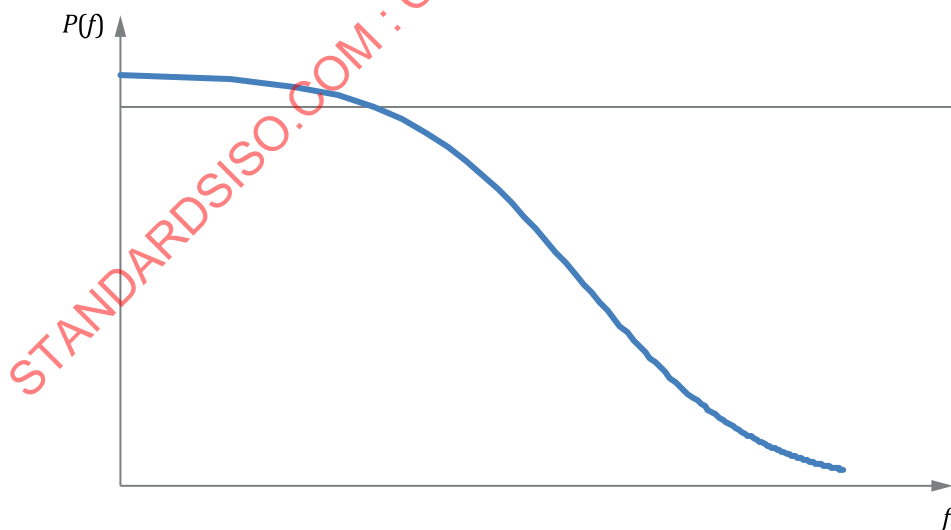


Figure 5 — Power spectral density of model noise

In practice, the power spectral density would be less smooth than the theoretical behaviour in [Figure 5](#). If it is an observed one, it might contain small oscillations about a smooth signal.

6.2 Estimation of noise parameters by autocovariance function

Another method based on recently published literature^[9] is introduced here for estimating the noise parameters, $\tilde{w}$, $\tilde{m}$, $\tilde{\rho}$, necessary for the precision evaluation.

An estimate of ρ can be made as

$$\tilde{\rho} = \frac{\sum_{t=1}^J \hat{\rho}_Y(t+1) / \hat{\rho}_Y(t)}{J} \quad (21)$$

where J is an integer ($2 \leq J \leq N - 1$) and

$$\hat{\rho}_Y(i) = \frac{\sum_{t=1}^{N-i} (Y_t - \bar{Y})(Y_{t+i} - \bar{Y})}{(N-i) \hat{\sigma}_Y^2} \quad (22)$$

$$\hat{\sigma}_Y^2 = \frac{\sum_{t=1}^n [Y(t) - \bar{Y}]^2}{n-1} \quad (23)$$

and

$$\bar{Y} = \frac{\sum_{t=1}^N Y_t}{N} \quad (24)$$

The other parameters can be estimated as

$$\tilde{m} = \sqrt{1 - \rho^2} \hat{\sigma}_r \quad (25)$$

and

$$\tilde{w} = \sqrt{\hat{\sigma}_Y^2 - \hat{\sigma}_r^2} \quad (26)$$

where

$$\hat{\sigma}_r^2 = \hat{\sigma}_Y^2 \frac{\hat{\rho}_Y(1)}{\rho} \quad (27)$$

An advantage of this method as compared to that of [6.1](#) is that although the number of data points used for the noise parameter estimation is limited to the n th power of 2 in the latter method, there is no such a limitation for the former.

We should note that the method in this clause is applicable to the background noise which is made up of the mixed random processes of the white noise and AR(1)-process or the AR(1)-process alone, and not the white noise alone.

6.3 Procedures for estimation of SD

The FUMI theory requires the instrumental output to estimate the SD of measured areas, which in turn leads to the minimum detectable value. The complete set of steps is illustrated in [Figure 6](#).

Digital instrumental output such as chromatograms and spectra is required for extracting both the signal and noise parameters. The signal parameters involve the signal region ranging from point 0 to point k_e , the integration region from point $k_c + 1$ to point k_f , and the zero window from point $-b + 1$ to

0 (see [Figure 3](#)). The signal region is made up of $k_e + 1$ data points, the integration region of $k_f - k_c$ data points, and the zero window b data points.

The signal parameters are arbitrary, but the following instruction will be useful in practice. For a Gaussian signal like that in [Figure 3](#), the signal region may be $\pm 3\sigma$ or $\pm 4\sigma$ around the signal centre, where σ denotes the standard deviation (width) and the signal centre is the mean of the Gaussian signal. In many cases, the peak height or entire area is the typical measurement. For the peak height measurement, $k_f = k_c + 1$. For the entire area measurement, $k_f = k_e - 1$ and $k_c = 0$. In either measurement, the number of data points in the integration region is $k_f - k_c$. The integration over a part of the signal region, as shown in [Figure 3](#), is also effective from the viewpoint of precision. The zero window may be set empirically as a region even narrower than the signal region, as shown in [Figure 3](#).

Unlike the signal parameters, the noise parameters ($\tilde{w}$, $\tilde{m}$ and $\tilde{p}$) are uniquely determined according to the stochastic properties of background noise. Some examples in liquid chromatography are given in [Table 1](#). The first step is to identify a region from the background noise that has no significant signal contribution.

The second step is the Fourier transform of the background noise in this region to give the power spectral density according to [Formula \(20\)](#). If the number, N , of data points in [Formula \(20\)](#) is limited to the n th power of 2 in the fast Fourier transform (FFT) used, 512 or 1 024 data points will be appropriate for the FFT region for the signal region of 50 or 100 data points.

The third step is the least-squares fitting of the theoretical curve [[Formula \(20\)](#)] to the power spectral density obtained as mentioned above. A small number of data points in the FFT region (e.g. 32) have a tendency, in some situations, to bias the exact estimation of the noise parameters.

The signal and noise parameters determined as mentioned above are substituted for [Formulae \(13\)](#) to [\(16\)](#) to provide the final value of the FUMI theory (precision).

The minimum detectable value can be found graphically from the precision profile as the net state variable at which the CV of the net state variable is 30 %. This method is also applicable for the CV of the response variable, since the precision profile of the net state variable is identical to that of the response variable if the calibration function is linear [[10](#), [11](#)]. In the case of nonlinear calibration, however, the precision profiles of the response variable and the net state variable are different. In this case, the method given in ISO 11843-5 applies to the transformation of the different precision profiles and the minimum detectable value can be estimated as well.

Table 1 — Example of noise parameters estimation in liquid chromatography^[6]

	$\tilde{w}$	$\tilde{m}$	$\tilde{p}$
Experiment A	14	3,7	0,99
Experiment B	12	9,0	0,94
Experiment C	14	5,6	0,99

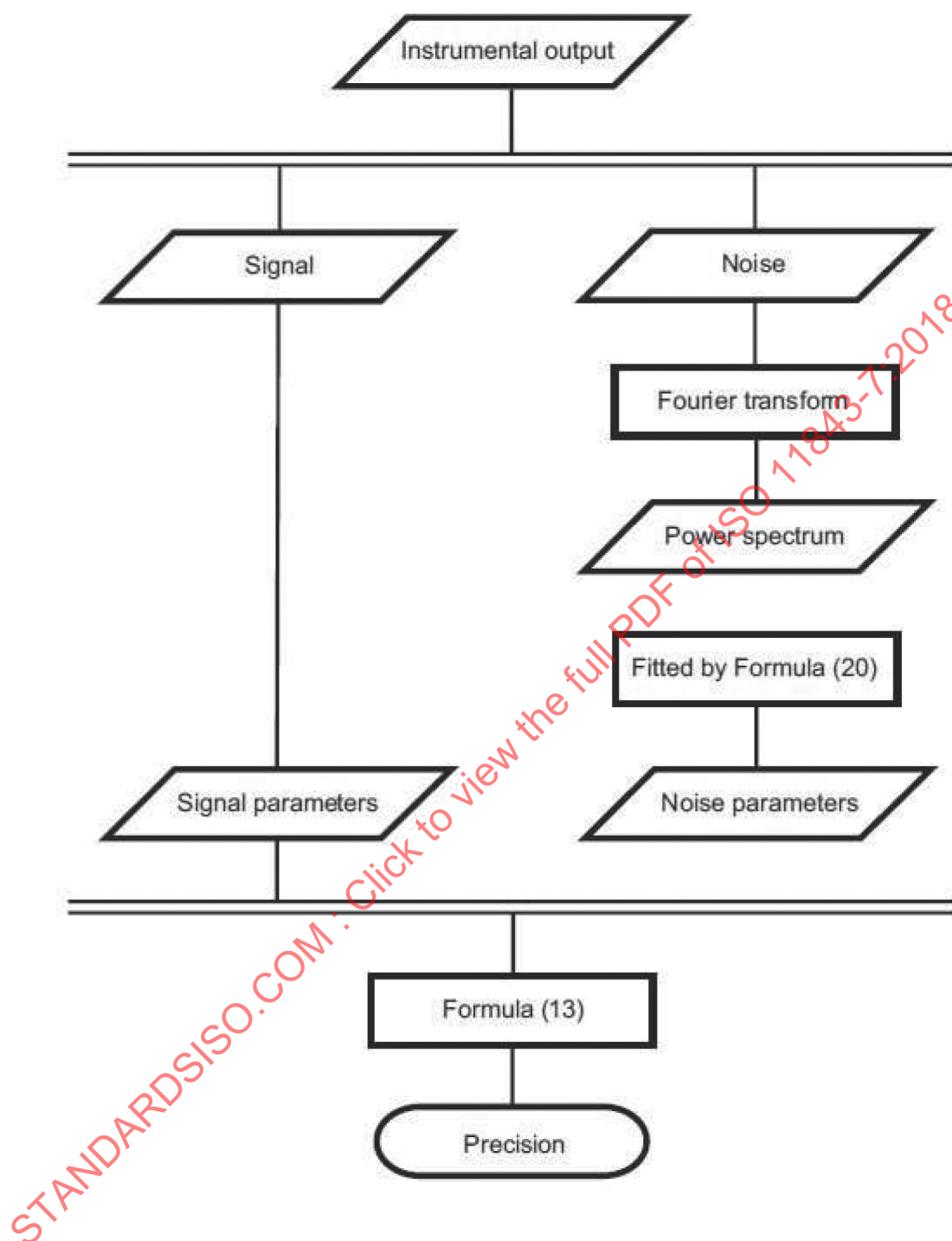


Figure 6 — Procedures of FUMI theory

Annex A (informative)

Symbols and abbreviated terms used in this document

SD	standard deviation
CV	coefficient of variation (SD divided by the mean)
X	net state variable
Y	response variable (measured area or height)
t_i	one of a set of uniformly spaced time values $t_1, t_2, \dots$
Δt	$t_{i+1} - t_i$, length of time interval for analogue-to-digital conversion in data acquisition
Y_i	intensity of instrumental output at time t_i , (sum of Y_i means Y)
Y_t	intensity of instrumental output at time t (sum of Y_t means Y)
x_d	minimum detectable value of net state variable
k_c	coefficient to specify the probability of an error of the first kind
k_d	coefficient to specify the probability of an error of the second kind
$\sigma_Y(X)$	SD of response variable as a function of X
$\sigma_X(X)$	SD of net state variable as a function of X
$ dY/dX $	absolute value of derivative of calibration function
$E[\cdot]$	mean of a random variable inside the square brackets
$\psi(\tau)$	auto-covariance function with lag τ , defined by Formula (5)
w_i	random variable at point i of the white noise with mean zero and SD $\tilde{w}$
$\tilde{w}$	SD of the white noise w_i
M_i	random variable of the AR(1)-process at point i
m_i	white noise included in AR(1)-process at point i , defined by Formula (4)
$\tilde{m}$	SD of the white noise m_i included in AR(1)-process
$\tilde{p}$	ratio for retaining the previous state, defined by Formula (4)
A_F	noise-created area, which is the area created by the noise alone
FUMI	abbreviation of function of mutual information