Document Number:	TD2021DL	Version Number:	1.0
Written by: Reviewed by:	*WADA* Science / *DL* Working Group *WADA* Laboratory Expert Group	Approved by:	*WADA* Executive Committee
Date:	21 December 2020	Effective Date:	1 April 2021

DECISION LIMITS FOR THE CONFIRMATORY QUANTIFICATION OF EXOGENOUS THRESHOLD SUBSTANCES BY CHROMATOGRAPHY-BASED ANALYTICAL METHODS

1.0 Introduction

The objective of this *Technical Document* (*TD*) is to harmonize the reporting of results for exogenous Threshold Substances (as listed in Table 1) when analyzed in urine *Samples* using chromatography-based quantitative Confirmation Procedures (CP), with particular regard to the *Decision Limits* (*DL*) that shall be applied to determine whether the analytical result indicates an *Adverse Analytical Finding* (*AAF*). It also describes the situations where the *DL* shall be corrected by the specific gravity (SG) of the urine *Sample*, as well as the use of Measurement Uncertainty (MU) information in the establishment of such *DL*.

[Comment: Decision Limits for endogenous Threshold Substances (e.g. human Chorionic Gonadotropin – hCG; human Growth Hormone - hGH) are defined in specific TD [1, 2] or Laboratory Guidelines [3].]

This document provides requirements on the following:

- Target Analytes;
- Threshold (T) and *DL*;
- Maximum values of MU;
- Adjustment of the *DL* for the SG;
- Reporting of quantitative results.

Further guidance is provided in Annex A, including:

- Estimating MU;
- Verification of MU by a Laboratory.

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Table 1

Substance Class	Threshold Substance	Threshold (T)	Maximum Relative Combined Standard Uncertainty at T $u_{c\_Max}(\%)$	*Decision Limit* (*DL*) [a]
S3. Beta-2 Agonists	Salbutamol	**1.00 µg/mL**	10	**1.20 µg/mL**
	Formoterol	**40.0 ng/mL**	15	**50.0 ng/mL**
S6. Stimulants	Cathine	**5.00 µg/mL**	10	**6.00 µg/mL**
	Ephedrine	**10.0 µg/mL**	5.0	**11.0 µg/mL**
	Methylephedrine	**10.0 µg/mL**	5.0	**11.0 µg/mL**
	Pseudoephedrine	**150 µg/mL**	5.0	**170 µg/mL**
S7. Narcotics	Morphine	**1.00 µg/mL**	15	**1.30 µg/mL**
S8. Cannabinoids	Carboxy-THC	**150 ng/mL**	10	**180 ng/mL**

a. The *DL*, expressed to three (3) significant figures, is obtained after adding a guard band *g* to the T, which accounts for the corresponding u_{c_Max} and ensures that any value above the *DL* obtained with the quantitative Analytical Method is higher than (>) the T with a statistical confidence of at least 95% (see Article 3.0).

2.0 Target Analytes

- Quantitative result

The *International Standard* for Laboratories (ISL) [4] requires that results from quantitative CP applied to Threshold Substances shall be based on the mean of three (3) independent determinations. The resulting relative standard deviation (RSD, %) shall be consistent with the quantitative CP method validation data.

The Laboratory shall demonstrate the Fitness-for-Purpose of the quantitative CP through method validation, including the estimation of the MU. Compliance with the criteria presented in Table 1 for $u_{c\_Max}(\%)$ ensures a harmonized reporting of *AAF*s at concentration levels exceeding the applicable *DL*.

- Qualitative result

In one of the three (3) independent determinations, the target Analyte(s) shall be identified in compliance with the prevailing TD IDCR [5].

2.1 Beta-2 Agonists - Formoterol and Salbutamol

The concentration level is based on content of formoterol or salbutamol, defined as the combination of free substance and its glucuronide conjugated forms, expressed as substance equivalent.

If one of these exogenous Threshold Substances is detected in a *Sample* in conjunction with a prohibited diuretic or other masking agent (as specified in the *Prohibited List* [6]), which is identified in compliance with the TD MRPL[7] and the TD IDCR [5], the confirmation of the Threshold Substance requires only the identification of the compound, not its quantification. In such cases, the Laboratory shall:

- Perform the CP for the diuretic/masking agent and report the results as an *AAF* in compliance with the TD MRPL [7] and the TD IDCR [5];

- Perform the (qualitative) CP for the beta-2 agonist and report the results as an *AAF* if identified **at any concentration level** in compliance with the TD IDCR [5].

In cases where a diuretic or masking agent is detected in the *Sample*, the co-presence of any of these beta-2 agonists shall be considered as an *AAF* unless there is an approved *TUE* for the beta-2 agonist itself (see ISL 2021, Article 5.3.6.2.2).

2.2 Stimulants - Cathine, Ephedrine, Methylephedrine and Pseudoephedrine

The concentration level is based on the parent compound of each target Threshold Substance in the free fraction.

If one of these exogenous Threshold Substances is detected in a *Sample* in conjunction with a prohibited diuretic or other masking agent [6], which is identified in compliance with the TD MRPL[7] and the TD IDCR [5], the confirmation of the stimulant requires only the identification of the compound, not its quantification. In such cases, the Laboratory shall:

- Perform the CP for the diuretic/masking agent and report the results as an *AAF* in compliance with the TD MRPL [7] and the TD IDCR [5];

- Perform the (qualitative) CP for the stimulant and report the results as an *AAF* if identified, in compliance with the TD IDCR [5], **at an estimated concentration level greater than (>) the applicable *Minimum Reporting Level* (*MRL*) for stimulants**, as defined in the TD MRPL [7].

In cases where a diuretic or masking agent is detected in the *Sample*, the co-presence of any of these stimulants shall be considered as an *AAF* unless there is an approved *TUE* for the stimulant itself (see ISL 2021, Article 5.3.6.2.2).

The Laboratory shall report cathine as an *AAF* when found at a urinary concentration level greater than (>) the *DL*. However, if pseudoephedrine is also detected in the *Sample* at concentration levels below (<) the *DL*, the concentration level of pseudoephedrine shall also be reported, and a comment shall be made in the Test Report that the cathine finding may have resulted from the administration of pseudoephedrine.

The Laboratories shall refer to TL05 (Oxilofrine) [8] or any other relevant Technical Letter providing guidance on findings related to Threshold Substances classified as stimulants in the *Prohibited List* [6].

2.3 Morphine

The concentration level is based on content of morphine, which is defined as the combination of free substance (free morphine) and its glucuronide conjugated forms (morphine-3-glucuronide and morphine-6-glucuronide), expressed as morphine equivalent.

Occasionally, a morphine finding may result from the administration of a permitted substance such as codeine or ethylmorphine:

- The Laboratories shall refer to the Technical Letter TL22 (Ethylmorphine) [9], which provides details on morphine findings that may be related to the administration of ethylmorphine;
- When codeine is detected in a *Sample*, Laboratories shall report an *AAF* for morphine in cases when both of the following conditions are met:
  - The morphine concentration level in urine is higher than (>) the *DL* or the adjusted *DL* (if SG > 1.018), and
  - The ratio M/C of morphine (M) to codeine (C, defined as the combination of free codeine + codeine-6-glucuronide, expressed as codeine equivalent) is equal to or higher than (≥) 2.00 (expressed truncated to three (3) significant figures), except when C > 5.00 µg/mL, which is indicative of only codeine intake (in this case, the quantification of morphine is not necessary, and the finding shall be reported as a Negative Finding).

    [Comment: The concentration level of C is expressed truncated to three (3) significant figures.]

2.4 Carboxy-THC (11-nor-Δ^9-tetrahydrocannabinol-9-carboxylic acid)

The concentration level is based on the content of carboxy-THC, which is defined as the combination of free substance and its glucuronide conjugated forms, expressed as substance equivalent.

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3.0 Threshold (T) and *Decision Limit* (*DL*)

Where a T has been established for a *Prohibited Substance*, the *DL* represents the value for that *Prohibited Substance* above which it can be decided that the result in a given *Sample*, obtained using a validated measurement procedure, has exceeded the T with a statistical confidence of at least 95%, and hence that an *AAF* is justified. This is illustrated in Figure 1.

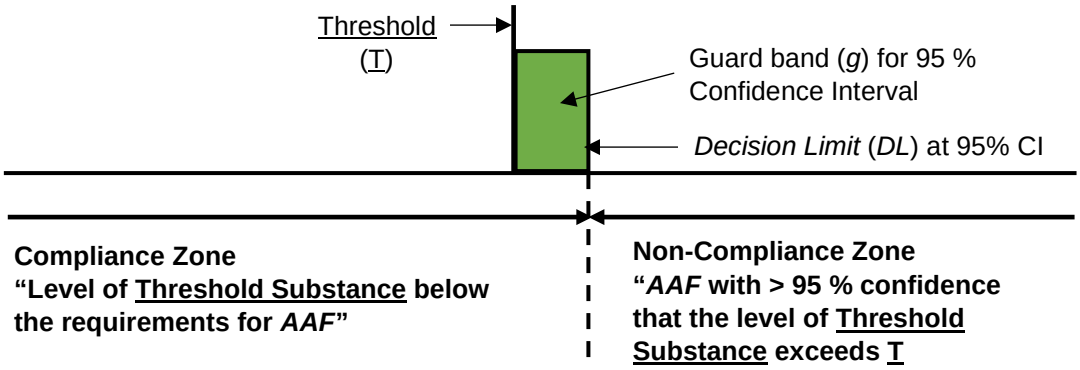

Figure 1: Use of a guard band (*g*) to establish a *DL* relative to a T and to differentiate between compliance and non-compliance zones.

The *DL* value shall be calculated as the sum of the T value and the guard band (*g*), where *g* is calculated based on the relevant *WADA* maximum acceptable value (unit/mL) of the combined standard uncertainty (u_{c_Max}) given in Table 1, using a coverage factor k of 1.645 (95% coverage range, one-tailed normal distribution). The resulting value of the *DL* is then rounded up to the second significant figure.

(Eq. 1) $DL = \mathrm{T} + g$

(Eq. 2) $g = k \cdot u_{c\_Max}$, with k = 1.645

(Eq. 3) $u_{c\_Max} = \mathrm{T} \cdot u_{c_{Max}}(\%)$

(Eq. 4) $AAF > DL$

When a value found in a *Sample* exceeds the T value, but is less than or equal to (≤) the *DL*, the Laboratory shall report this result as a Negative Finding and include a recommendation (*e.g.* in the opinion section of the Test Report) for the *Results Management* Authority to consider this result within its future "target and intelligence" test planning. This result shall not constitute an *AAF* regardless of the value of MU the Laboratory reports for the result.

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4.0 Maximum Levels of Measurement Uncertainty

The maximum acceptable relative combined standard uncertainty (u_{c_Max}, %) represents the minimum requirement to be met by a Laboratory for the uncertainty of the measurement, estimated at levels close to the T value, when reporting a result for the determination of a Threshold Substance. The u_{c_Max} (%) values are set such that a Laboratory can reasonably expect to work within them when applying quantitative CPs for the determination of Threshold Substances.

In most cases, the u_{c_Max} (%) is assigned using robust estimates of method Reproducibility (S_R) obtained from the combined participant Laboratory results from relevant rounds of the External Quality Assessment Scheme (EQAS). In cases where a new Threshold Substance is introduced into this *TD* before EQAS performance data are available, alternative approaches will be used to assign the relevant u_{c_Max} (%). In this case the assignment of u_{c_Max} (%) must be reviewed and approved by the *WADA* Laboratory Expert Group (LabEG). When data obtained from subsequent EQAS rounds becomes available, the u_{c_Max} (%) may be revised to reflect the actual analytical performance of the Laboratories.

The results obtained from the *WADA* EQAS indicate that these minimum requirements are conservative. When setting the target values, the degrees of freedom associated with the MU data are assumed to be large.

- Laboratories shall estimate the relative combined standard uncertainty (u_c, %) for a result at levels close to the T value for each quantitative CP for Threshold Substances;
- The estimated u_c (%) shall be not greater than (≤) the u_{c_Max} (%) value given in Table 1.

> *[Comment: As mentioned above, these u_{c_Max} (%) values are considered to be conservative; therefore, smaller u_c (%) values may be reported by Laboratories.]*

Various approaches to obtain Fit-for-Purpose estimates of u_c (%) associated with the results from a given measurement procedure are given in Annex A.

5.0 Adjustment of the *DL* for the Urine Specific Gravity (SG)

- For any of the Threshold Substances treated in this document, when the SG of the urine *Sample* (SG_{Sample}) is greater than (>) 1.018, an adjusted *DL* for an individual test result (DL_{adj}) shall be calculated as per Eq. 5 below;

> *[Comment: The SG_{Sample} cut-off value for adjustment of the DL has been set at 1.018 to account for the lower limit of the 95% coverage interval, based on a two-tailed normal distribution, of a reference value of SG at 1.020 for normally hydrated individuals (calculated as 1.020 – U_{Max_SG})].*

- The SG value (SG_{Sample}) to be used in applying Eq. 6 for the calculation of SG_{Sample_Max} is that measured in the Laboratory.

[Comment: The Laboratory shall measure the SG_{Sample} in a single Aliquot during the Initial Testing Procedure (ITP) and the CP, using a method that is included within the Laboratory's ISO/IEC 17025 scope of accreditation, as follows:

- *ITP: In all Samples, using either a digital refractometer or a densitometer;*
- *CP: A digital refractometer shall be used in all "A" and "B" Samples. The adjustment of the DL for the SG is not needed for:*

  (i) "A" and "B" Sample confirmations for those exogenous Threshold Substances that shall not be quantified if detected in the presence of a prohibited diuretic or other masking agent, and

  (ii) "B" Sample confirmations of exogenous Threshold Substances, since in those cases, in accordance with the ISL [4], "B" Sample results shall only confirm the "A" Sample identification (in compliance with the TD IDCR [5]) for the AAF to be valid.

If the SG_{Sample}, as measured by the instrument, reads to ≥ 4 decimal places, the SG_{Sample} is the value obtained after rounding the instrumental value and expressing it to three (3) decimal places (e.g. 1.0223 should be expressed as 1.022; 1.0227 as 1.023. When the measured value finishes in 5, it should be expressed to the nearest higher 3-decimal place value, e.g. 1.0225 should be expressed as 1.023).]

- The SG-adjustment to the *DL* shall be made using the following formula:

$$(\text{Eq. } 5) \quad DL_{\text{adj}} = \frac{(\text{SG}_{\text{Sample Max}} - 1)}{(1.020 - 1)} \,.\, DL$$

Where SG_{Sample_Max} is calculated as:

$$(\text{Eq. } 6) \quad \boldsymbol{SG}_{Sample\_\text{Max}} = \text{SG}_{Sample} + U_{\text{Max\_SG}} = \text{SG}_{Sample} + 0.002$$

$U_{\text{Max\_SG}}$ = 0.002 is the maximum allowed expanded uncertainty ($U_{95\%}$, k = 2) for SG.

- The determined DL_{adj} shall be expressed truncated to three (3) significant figures (trailing zeros (0) shall be considered as significant figures, *e.g.* 1.50; 100) (see Annex B).

6.0 Reporting

The minimum requirements for reporting an *AAF* for a Threshold Substance are:

- The quantitative result (reported as the mean value from triplicate determinations, truncated to three (3) significant figures; trailing zeros (0) shall be considered as significant figures, *e.g.* 13.0; 190);
- A statement that the quantitative result exceeds (>) the relevant *DL* (or DL_{adj}, if SG > 1.018); and
- The u_c (%) associated with a result at levels close to the T value, as determined during the quantitative CP method validation (which shall not be higher than (≤) the corresponding u_{c_Max} (%) specified in Table 1).

Reporting Example for the Test Report:

The concentration level of '*Prohibited Substance* A' in the *Sample* is X.XX (units). This exceeds the *DL* (after adjustment for the SG, if applicable) for A of Y (units). The relative combined standard uncertainty (u_c %) estimated by the Laboratory for a result at the Threshold Z is '*b*' (%). This result constitutes an *Adverse Analytical Finding* for the presence of A in the *Sample*.

7.0 Interpretation Examples

7.1 Ephedrine is detected in a *Sample* with an SG of 1.018 at a concentration level of 11.208 µg/mL using a quantitative Analytical Method where the u_c (%) is 3.6% for a result at the T of 10.0 µg/mL.

In accordance with the reporting rules established in this *TD* (see Article 6.0), this result constitutes an *AAF* since the concentration level of ephedrine in the *Sample*, truncated to three (3) significant figures, is 11.2 µg/mL and exceeds the *DL* for ephedrine of 11.0 µg/mL. The u_c (%) of 3.6 % is lower than the corresponding u_{c_Max} (%) of 5.0. Such a finding shall be reported as follows:

> Test Report: The concentration level of ephedrine in the *Sample* is 11.2 µg/mL. This exceeds the *DL* for ephedrine of 11.0 µg/mL. The relative combined standard uncertainty (u_c %) estimated by the Laboratory for a result at the Threshold (10.0 µg/mL) is 3.6%. This constitutes an *AAF* for the presence of ephedrine in the *Sample*.

7.2 Carboxy-THC is detected in a *Sample* with a SG of 1.022 at a concentration level of 216.7 ng/mL using a quantitative Analytical Method where the u_c is 9.0 % for a result at the Threshold of 150 ng/mL. The DL_{adj} calculated according to Eq. 5 and expressed to three (3) significant figures is 216 ng/mL (see Annex B).

In accordance with the reporting rules established in this *TD* (see Article 6.0), this result does not constitute an *AAF*, since the concentration level of carboxy-THC in the *Sample*, truncated to three (3) significant figures, is 216 ng/mL and does not exceed the DL_{adj} for carboxy-THC of 216 ng/mL.

Since the concentration level of carboxy-THC does not exceed the adjusted *DL*, the Laboratory shall report this result as a Negative Finding and include a recommendation (*e.g.* in the opinion section of the Test Report) for the *Results Management* Authority to consider this result within its Test Distribution Plan.

> *[Comment: When the result for a Threshold Substance in a Sample scantily exceeds the DL, the confidence interval [mean ± expanded uncertainty $U_{95\%}$ (k = 2)] for the Laboratory result may extend below the DL. It is important to note that this shall not invalidate an AAF. For appropriate statistical comparison, the u_c with a single-tailed distribution coverage factor (k = 1.645) is taken into consideration when the Laboratory result is compared to the T to demonstrate that the result obtained for the Threshold Substance exceeds the T at greater than (>) 95% confidence.]*

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ANNEX A

1. Estimating Measurement Uncertainty (MU)

The International Vocabulary of Metrology (ISO/IEC Guide 99:2007) [10] formally defines MU as a parameter characterizing the dispersion of quantity values attributed to a measurand.

More simply stated, the combined standard MU of a result [$u_c(y)$] is equivalent to an estimate of the standard deviation (SD) associated with the result (y) that would have been obtained for the sample under analysis if repeated several times. Multiplication of $u_c(y)$ by a coverage factor (k) gives the expanded MU (U) associated with result (y). For a given sample, the combination of the result (y) and its associated U specifies a range describing the dispersion of the values that can reasonably be attributed to the measurand at a stated level of statistical confidence. For *Doping Control* purposes, a value of U corresponding to a 95% coverage range is applied.

Accreditation to ISO/IEC 17025 [11], as well as compliance with the ISL [4], requires that Laboratories evaluate the MU associated with their results at levels close to the Threshold, and report the uncertainty where applicable. The ISO/IEC Guide to the Expression of Uncertainty in Measurement (GUM) establishes general rules for evaluating and expressing uncertainty in measurement that are applicable to ISO/IEC 17025 accredited laboratories [12].

The examples cited in the GUM concentrate on one method, referred to elsewhere as the "analytical", "modelling" or "bottom-up" approach, for uncertainty evaluation. The basic GUM principles also allow for more global approaches for estimating the sources of MU, generally referred to as "top-down" or "empirical" approaches, using data derived from intra- or inter-laboratory method validation studies, internal quality control procedures or the results of EQAS. These approaches are all potentially compliant with the GUM principles provided the MU estimate obtained is suitable for the intended purpose of the measurement. Various references are available which give worked examples of both the "bottom-up" and "top-down" approaches to MU estimation [13, 14, 15, 16, 17].

Different approaches may be applied for the estimation of the combined standard measurement uncertainty $u_c(y)$ associated with an individual result (y). They use:

A. A modelling approach based on the principles described in the GUM;

B. Intra-laboratory approach: "In-house" method validation data combined with quality control data;

C. Inter-laboratory approach: Data derived from inter-laboratory collaborative trials or from EQAS.

The strategy used for uncertainty estimation does not have to follow one exclusive model and in practice the combination of data obtained from two or more different approaches can be employed.

All of these approaches are considered acceptable. Any of these approaches may be employed by a Laboratory to estimate the MU associated with their measurement results, provided the Laboratory

estimate does not exceed the maximum acceptable (target) MU associated with the determination of specific Threshold Substances that have been established by *WADA*. These maximum acceptable MU are conservative estimates derived from EQAS performance data.

A. Modeling Approach

In this case, the Laboratory develops a measurement equation or model in which result (y) is a function of independent input parameters x_1, x_2, x_3….x_n that all influence the measurement result.

If the mathematical model is a combination of addition/subtraction and multiplication/addition operations, then an appropriate quadratic combination is used to calculate the $u_c(y)$. This approach is also referred to as the "bottom-up" or "GUM" approach.

The GUM approach is based on the propagation of uncertainties where the estimated standard deviation associated with the measurement result (y) is named $u_c(y)$ and is determined from the estimated standard deviations associated with each input estimate (x_i). These uncertainty components from the input quantities are then combined to give the combined standard uncertainty $u_c(y)$.

When the input quantities are independent, the $u_c(y)$ is given as:

$$(\text{Eq. } 7) \qquad u_c(y) = \sqrt{\sum_{i=1}^{N} \left(\frac{\partial f}{\partial x_i}\right)^2 u^2(x_i)}$$

Where f is the function that defines the measurand.

More details on the application of this method and the implications in cases where two or more of the input quantities are correlated can be found in the GUM and elsewhere in the literature [12, 15].

> *[Comment: The uncertainty budget derived using this approach indicates the relative magnitude of the various sources of uncertainty but carries the risk of missing a contributing factor which may significantly affect the overall estimate of MU. Nonetheless, it is a valuable means of establishing where the major sources of uncertainty are found in a quantitative CP and for identifying where efforts should be focused if a reduction is desired in the overall MU of results obtained through use of the quantitative CP.]*

B. Intra-Laboratory Data Approach

This approach assumes that the quantitative CP has undergone intra-Laboratory validation including an estimation of the Intermediate Precision (also referred to as the within-Laboratory reproducibility or imprecision). It is based on a three-component measurement model:

$$(\text{Eq. } 8) \qquad y = m + B + e$$

The result (y) is the sum under Intermediate Precision conditions of the measurement method mean (m), an estimate of method bias (B) and a random error contribution (e) and the $u_c(y)$ associated with the result is given by:

$$(\text{Eq. } 9) \qquad u_c(y) = \sqrt{u(m)^2 + u(B)^2 + u(e)^2}$$

The estimate of within-Laboratory Intermediate Precision of results, usually obtained from intra-Laboratory QC and method validation data, can be expressed as a standard deviation (s_w). It provides a Fit-for-Purpose estimate of the uncertainty contribution from the $u(m)$ and $u(e)$ terms and the "internally visible" bias component (B_{Int}).

$$(\text{Eq. } 10) \qquad S_w \sim \sqrt{u(m)^2 + u(e)^2 + u(B_{Int})^2}$$

If (y) is the result of a single analysis, the equation for calculating the standard uncertainty associated with the result simplifies to:

$$(\text{Eq. } 11) \qquad u_c(y) = \sqrt{{s_w}^2 + u(B_{Ext})^2}$$

When (y) is the average of n replicate analyses:

$$(\text{Eq. } 12) \qquad u_c(y) = \sqrt{\frac{{s_w}^2}{n} + u(B_{Ext})^2}$$

In both cases, B_{Ext} is an estimate for bias not accounted for by intra-Laboratory studies and the uncertainty due to bias [u_{bias} or $u(B_{ext})$] can be estimated by using the following equations [13]:

$$(\text{Eq. } 13) \quad u_{bias} = \sqrt{{\Delta_i}^2 + \frac{s^2}{n} + u_{ref}^2}$$

where:
n - number of replicate measurements of the sample used as reference (CRM, QC or EQAS sample) prepared at a specified dilution level;
s - standard deviation (SD) under Repeatability conditions of the results obtained for the replicate measurements of the reference sample at a specified dilution level;
u_{ref} - uncertainty of the reference sample, and;
$\Delta_i = C_{lab,i} - C_{ref,i}$

Where information is available from n_{bias} separate bias determinations, then the u_{bias} shall be expressed as the root mean square of the bias (RMS_{bias}).

$$(\text{Eq. } 14) \quad u_{bias} = RMS_{bias} = \sqrt{\frac{\Sigma u_{bias\_i}^2}{n_{bias}}}$$

where:
n_{bias} - number of independent bias determinations.

[Comment: When appropriately applied, this approach, as with the other empirical approaches, is as valid as the modeling approach, and should provide a conservative but pragmatic estimation of MU.]

C. Inter-Laboratory Method Performance or EQAS Approach

Where a Laboratory has participated in an inter-Laboratory comparison to evaluate a quantitative CP, or has demonstrated appropriate implementation of a literature method validated using such an approach, the inter-Laboratory Reproducibility of the method (s_R), calculated from the results of the comparison and expressed as SD, can be used as an estimate of the u_c of an individual result obtained using the method:

$$(\text{Eq. } 15) \qquad u_c(y) = \frac{s_R}{\sqrt{n}} \qquad (y \text{ is the average of } n \text{ replicate analyses})$$

This approach is applicable, in practice, only when the validation study includes a multi-centre, inter-Laboratory trial conducted to a pre-defined experimental protocol.

> *[Comment: The major sources of variability can be assessed by inter-Laboratory studies and provide estimates of Repeatability standard deviation (s_r), Reproducibility (s_R) and Bias (B) of the method (with respect to a known reference value). The Reproducibility (s_R) can be used as an estimate of the u_c associated with an individual measurement result obtained using this quantitative CP Procedure.]*

Data obtained from ongoing participation in an EQAS also allows, in some cases, for the calculation of a performance characteristic of the ensemble of methods used by participants that can serve, in the absence of a properly constituted inter-Laboratory study, as a conservative estimate of the Reproducibility (s_R) of the quantitative CP used by an individual Laboratory. It is mostly in the latter sense that the term s_R is used in the current draft. This estimate is only valid when:

- The values reported by participants in the EQAS round (after exclusion of outliers) fall into a normal Gaussian distribution;
- The intra-Laboratory Repeatability (s_r) for the method is smaller than (<) the variation of the participants' results ($s_r < s_R$);
- Uncertainty contributions from instability or heterogeneity of the EQAS sample are negligible;
- The matrices utilized correspond closely to those encountered in routine analytical conditions (*i.e.* "representative" matrices are used to prepare the EQAS materials);
- The target values of the study fall within the range of application of the method;
- The Laboratory obtains satisfactory results in a minimum number of consecutive rounds.

In this case the SD of the participants' results after exclusion of outliers or as calculated from robust statistics can be used as an estimate of the u_c associated with a result obtained by the method. This value can then be applied as described for the s_R estimate above.

> *[Comment: As noted before, the Reproducibility (s_R) estimate can be used as a conservative estimate of the u_c associated with a result. Moreover, a Laboratory can, by its participation in the WADA EQAS, verify and demonstrate the validity of its chosen approach to estimate the MU.]*

2. Verification of Measurement Uncertainty

Regardless of the approach employed by a Laboratory to estimate the MU for the results it obtains using a particular quantitative CP, it is important that this MU estimate be validated, and its veracity continuously monitored. This can be accomplished by regular comparison with an appropriate QC sample, preferably a Certified Reference Material (CRM), if available, and/or through evaluation of method performance using EQAS data.

The MU for a particular quantitative CP, estimated by a Laboratory can also be checked by comparison to data generated from an appropriate EQAS by employing the E_n number.

$$\text{(Eq. 16)} \qquad E_n = \frac{x - x_a}{\sqrt{U(x)^2 + U(x_a)^2}}$$

Where x_a is the assigned value for the EQAS study, x is the Laboratory result, and $U(x_a)$ and $U(x)$ are respectively the expanded uncertainties associated with each result.

Monitoring the $|E_n|$ values over time provides the Laboratory an important tool to evaluate the agreement between its MU estimation for a quantitative procedure and the actual performance of that procedure. Provided that the estimated MU is less than or equal to (≤) the u_{c_Max} required by *WADA*, it is considered that when $|E_n|$ is distributed:

- Around one (1): then the estimated MU is in good agreement with the Laboratory's EQAS performance*;*
- Repeatedly at levels considerably smaller than (<<) one (1): then the MU could be overestimated. This shows that the historical Laboratory performance in the EQAS compared to the inter-Laboratory consensus values is better than its estimated MU. The Laboratory should evaluate the need for re-assessing the MU for this particular quantitative CP;
- Repeatedly greater than (>) one (1): the MU could be underestimated as the Laboratory's performance in the EQAS is worse than its estimated MU. In this case the reason for the high E_n value should be re-assessed. If necessary, steps should be taken to re-evaluate the MU.

It is important to highlight that individual $|E_n|$ values greater or lower than one (1) may not necessarily justify actions to be taken by the Laboratory. Rather, the history of values and their trends should be monitored.

Whenever there is a change in the quantitative CP (extraction step, derivatization conditions, internal standard, etc.) a re-validation of the procedure and a re-assessment of the MU of results obtained using the altered procedure is required. It is necessary to check that the quantitative CP is still Fit-for-Purpose (*e.g.* the MU estimated by the Laboratory for a particular quantitative CP is below the acceptable u_{c_Max} given in Table 1 above).

Document Number:	TD2021DL	Version Number:	1.0
Written by: Reviewed by:	*WADA* Science / *DL* Working Group *WADA* Laboratory Expert Group	Approved by:	*WADA* Executive Committee
Date:	21 December 2020	Effective Date:	1 April 2021

ANNEX B Adjusted *Decision Limits*

Table 2. Adjusted *DL*s calculated for SG > 1.018 as per Eq. 5 and expressed truncated to three (3) significant figures

		Salbutamol	Formoterol	Cathine	Ephedrine	MethylE	PSE	Morphine	C-THC
SG	SG_{Max}	**1.20**	**50.0**	**6.00**	**11.0**	**11.0**	**170**	**1.30**	**180**
1.019	1.021	1.26	52.5	6.30	11.5	11.5	178	1.36	189
1.020	1.022	1.32	55.0	6.60	12.1	12.1	187	1.43	198
1.021	1.023	1.38	57.5	6.90	12.6	12.6	195	1.49	207
1.022	1.024	1.44	60.0	7.20	13.2	13.2	204	1.56	216
1.023	1.025	1.50	62.5	7.50	13.7	13.7	212	1.62	225
1.024	1.026	1.56	65.0	7.80	14.3	14.3	221	1.69	234
1.025	1.027	1.62	67.5	8.10	14.8	14.8	229	1.75	243
1.026	1.028	1.68	70.0	8.40	15.4	15.4	238	1.82	252
1.027	1.029	1.74	72.5	8.70	15.9	15.9	246	1.88	261
1.028	1.030	1.80	75.0	9.00	16.5	16.5	255	1.95	270
1.029	1.031	1.86	77.5	9.30	17.0	17.0	263	2.01	279
1.030	1.032	1.92	80.0	9.60	17.6	17.6	272	2.08	288
1.031	1.033	1.98	82.5	9.90	18.1	18.1	280	2.14	297
1.032	1.034	2.04	85.0	10.2	18.7	18.7	289	2.21	306
1.033	1.035	2.10	87.5	10.5	19.2	19.2	297	2.27	315
1.034	1.036	2.16	90.0	10.8	19.8	19.8	306	2.34	324
1.035	1.037	2.22	92.5	11.1	20.3	20.3	314	2.40	333
1.036	1.038	2.28	95.0	11.4	20.9	20.9	323	2.47	342
1.037	1.039	2.34	97.5	11.7	21.4	21.4	331	2.53	351
1.038	1.04	2.40	100	12.0	22.0	22.0	340	2.60	360
1.039	1.041	2.46	102	12.3	22.5	22.5	348	2.66	369
1.040	1.042	2.52	105	12.6	23.1	23.1	357	2.73	378

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Date:	21 December 2020	Effective Date:	1 April 2021

8.0 References

[Current versions of WADA ISL, Technical Documents and Laboratory Guidelines may be found at https://www.wada-ama.org/en/what-we-do/science-medical/laboratories]

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[3] *WADA* Laboratory Guidelines on human Growth Hormone Biomarkers Test for *Doping Control* Analyses.

[4] The World Anti-Doping *Code International Standard* for Laboratories (ISL).

[5] *WADA Technical Document* TD IDCR: Minimum Criteria for Chromatographic-Mass Spectrometric Confirmation of the Identity of Analytes for *Doping Control* Purposes.

[6] The World Anti-Doping *Code International Standard Prohibited List.*

[7] *WADA Technical Document* TD MRPL: Minimum Required Performance Levels for Detection and Identification of Non-Threshold Substances by Chromatographic-Mass Spectrometric Analytical Methods.

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