Certificate of Analysis_(Ver.2.0)

Aflatoxin Total in Acetonitrile

1. General information

This document is designed and the certified value(s) and uncertainty(ies) are determined in accordance with ISO Guide 31^[1].

2. Description of the Reference Material (RM)

Product Name: Aflatoxin Total in Acetonitrile

Product Number: STD#1089

Lot#: 1101B06

CAS number: Aflatoxin B₁: 1162-65-8; Aflatoxin B₂:7220-81-7; Aflatoxin G₁: 1165-39-5; Aflatoxin G₂:7241-98-7

Forulma: Aflatoxin B₁: C₁₇H₁₂O₆;Aflatoxin B₂:C₁₇H₁₄O₆ ;Aflatoxin G₁:C₁₇H₁₂O₇;Aflatoxin G₂: C₁₇H₁₄O₇

Formula Weight: Aflatoxin B₁: 312.27; Aflatoxin B₂: 314.29 ; Aflatoxin G₁: 328.27; Aflatoxin G₂: 330.29

 $Result \ Concentration: \quad \ \ Aflatoxin \ B_1: 20.08\pm0.28\mu g/mL; \qquad \quad \ \ Aflatoxin \ B_2: 20.11\pm0.28\mu g/mL;$

Aflatoxin G_1 : 20.76±0.29 μ g/mL; Aflatoxin G_2 : 20.27±0.53 μ g/mL

Starting material: Aflatoxin B₁,lot#119906P,Pribolab Pte. Ltd.; Aflatoxin B₂,lot#119911P,Pribolab Pte. Ltd.;

Aflatoxin G₁,lot#l19909P,Pribolab Pte. Ltd.; Aflatoxin G₂,lot#l19818P,Pribolab Pte. Ltd.

Matrix: Acetonitrile, LiChrosolv®, Merck

Amount: 1.2mL

Production date: 06,Jun,2020
Expiry date: 05,Dec,2021

Name of the supplier: Pribolab Pte. Ltd.

2.1 Intended use of the RM

- for laboratory use only
- calibration of analytical instruments

2.2 Instruction for the correct use of the RM

The compound should be stored at -20° C or below in a dark place. Before usage of the RM,the compound should be allowed to warm to temperature($20\pm3^{\circ}$ C). The recommended minimum subsample amount for all kinds of application is 100 µL. The expiry date of this RM is based on the current knowledge and holds only for proper storage conditions in the originally closed flasks/packages.

2.3 Hazardous situation

The normal laboratory safety precautions should be observed when working with this RM.Further details for the handing of this RM are available as safety data sheet.

Hazardous IngredientsConcentration in%PictogramsSignal wordHazard statement(s)Acetonitrile>99.9DangerH225,H302,H312,H319,H332

3. Certified values and their uncertainties

Aflatoxin Total in acetonitrile							
	Mass concentration ^a			Mass concentration ^a			
Compound	Certified value ^b	Uncertainty ^c	Compound	Certified value ^b	Uncertainty ^c		
Aflatoxin B₁	20.08µg/mL	±0.28µg/mL	Aflatoxin G₁	20.76μg/mL	±0.29µg/mL		
Aflatoxin B ₂	20.11μg/mL	±0.28µg/mL	Aflatoxin G ₂	20.27μg/mL	±0.53µg/mL		

- a Mass concentration based on weighed amount, purity and dilution steps
- b Values are based on preparation data and confirmed experimentally by HPLC-DAD
- c Expanded uncertainty U(k=2) of the value u_c according to GUM^[2]

3.1 Calculation of uncertainty

After the concentration of the gravimetric prepared solution was confirmed by HPLC-DAD, the uncertainty of the calibrant was calculated on the basis of preparation^[3].

Uncertainty components	Description	Standard uncertainty _(U)	
Purity(P)of solid Aflatoxin B ₁ Aflatoxin B ₂ Aflatoxin G ₁ Aflatoxin G ₂	P ₁ =99.0±1.0% P ₂ =98.9±1.1% P ₃ =98.9±1.1% P ₄ =98.0±2.0%	u(P ₁)=0.6% u(P ₂)=0.6% u(P ₃)=0.6% u(P ₄)=1.2%	а
Weighing procedure: Weighted sample: m _{Aflatoxin B1} =50.707mg m _{Aflatoxin B2} =50.834mg m _{Aflatoxin G1} =52.477mg m _{Aflatoxin G2} =51.709mg	$U_{(m)}$ =0.0000008g+1.30*10 ⁻⁵ * m_{Toxin} $u_{(m)}$ = $U_{(m)}$ /2	u _(m) =0.0005mg	b
Dilution procedure Volumetric flask1:V _{f1} =250mL Volumetric flask2:V _{f2} =250mL One-mark glass pipette:V _P =25mL	calibration flask1: 250mL±0.15mL repeatability flask1: 0.03mL volume expansion solvent flask1 calibration flask2: 250mL±0.15mL repeatability flask2: 0.03mL volume expansion solvent flask2 Calibration pipette: 25mL±0.03mL volume expansion solvent pipette	u(cal1)=0.06mL u(rep1)=0.03mL u(Vol.exp.1)=0.59mL u(V1)=0.59mL u(cal2)=0.06mL u(rep2)=0.03mL u(Vol.exp.2)=0.59mL u(V2)=0.59mL u(cal3)=0.01mL u(Vol.exp.3)=0.06mL u(v)=0.06mL	c d e f g h i j k l

a Maximum tolerance of purity was divided by $\sqrt{3}$

f,j,m The three contributions are combined to give the $u(V) = \sqrt{u(cal)^2 + u(rep)^2 + u(Volexp)^2}$

Calculation of the combined uncertainty uc and the expanded standard uncertainty U

b Calculation of this u-value is based upon the uncertainty formula for the weighed amount as given in the calibration report from annual balance calibration

c,g,k A triangular distribution(division by $\sqrt{6}$)was chosen for the calculation of u(cal)

d,h Based on a series of ten fill and weigh experiments on a typical 250mL flask; the value was used directly as a standard deviation

e,i,I Based on the density of 0.7857 g/cm³ at temperature T=20 $^{\circ}$ C and a maximum temperature variation of $\pm 3^{\circ}$ C, of volume expansion, relative volume expansion coefficient of acetonitrile is 1370*10-6/ $^{\circ}$ C[7],volume expansion term(rectangular distribution)was divided by $\sqrt{3}$

$$\mathbf{C}_{\text{Toxin}} = \frac{10 \times \mathbf{m}_{\text{ws}} \times P1 \times V_{P}}{V_{f_{1}} \times V_{f_{2}}} = \frac{10 \times 50.707 \times 99.0 \times 25}{250 \times 250} = 20.08 \, \text{mg} \, / L$$

$$\frac{u_{\cdot} C_{\text{Toxin}}}{C_{\text{Toxin}}} = \sqrt{\left[\frac{u(P)}{P}\right]^{2} + \left[\frac{u(m)}{m_{\text{ws}}}\right]^{2} + \left[\frac{u(V1)}{V_{f1}}\right]^{2} + \left[\frac{u(V2)}{V_{f2}}\right]^{2} + \left[\frac{u(V3)}{V_{P}}\right]^{2}} = \sqrt{\left[\frac{0.6}{99.0}\right]^{2} + \left[\frac{0.0004}{50.707}\right]^{2} + \left[\frac{0.59}{250}\right]^{2} + \left[\frac{0.06}{25}\right]^{2} = 0.007}$$

$$\mathbf{U}_{\mathbf{C}} \left(\mathbf{C}_{\text{Toxin}}\right) = \mathbf{C}_{\text{Toxin}} \times 0.007 = 20.08 \times 0.007 = 0.140 \, \text{mg} \, / L$$

$$\mathbf{Calculation of expanded standard uncertainty U using a coverage factor k=2}$$

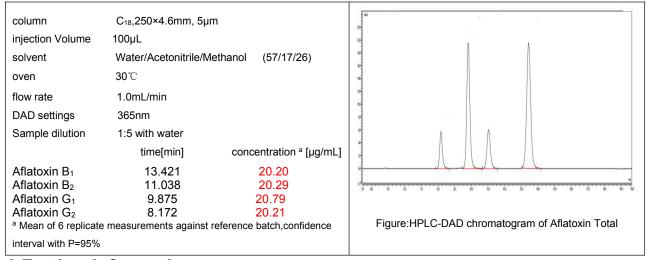
$$\mathbf{U}_{\mathbf{C}} \left(\mathbf{C}_{\text{Toxin}}\right) = \mathbf{U}_{c} \left(\mathbf{C}_{\text{Toxin}}\right) \times 2 = 0.140 \times 2 = 0.28 \, \text{mg} \, / L = 0.28 \, \mu \, \text{g} \, / \, \text{mL}$$

4. Discussion of traceability

This calibrant is certified on the basis of gravimetric preparation^[4]. Thus the certified value(mass concentration of Aflatoxin Total is based on the weighed amount of the starting material and is therefore traceable to the stated purity of the solid raw material. High purity material represents a practical realization of concentration units, through conversion of mass to molar quantity.

5. Confirmation of certified value by HPLC-DAD

The certified concentration of Aflatoxin Total of the gravimetric prepared solution was confirmed by HPLC-DAD against an independently prepared reference batch of Aflatoxin Total.



6. Further information

The purchaser must determine the suitability of this product for its particular use. Pribolab makes no warranty of any kind, express or implied, other than its products meet all quality control standards set by Pribolab. We do not guarantee that the product can be used for a special application.

Inspected by

Quality System Specialist

References:

- [1] ISO Guide 31, 1-7, (2000), "Reference Materials Contents of Certificates and Labels"
- [2] International Organization for Standardization (ISO), (2008), "Guide to the Expression of Uncertainty in Measurements", (GUM 1995 with minor corrections) 1st Ed. Geneva, Switzerland
- [3] R.D. Josephs, R. Krska, S. MacDonald, P. Wilson, H. Pettersson, J. AOAC Int. 86, 50-60. (2003), "Preparation of a Calibrant as Certified Reference Material for Determination of the Fusarium Mycotoxin, Zearalenone"
- [4] E.W. Flick, (1998), "Industrial Solvents Handbook",5rd Ed., Noyes Data Corp. Westwood NJ