

Certificate of Analysis

Excellence through measurement

U-[13C₁₇]-AFLATOXIN B1 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] and Eurachem / CITAC Guides [2,3].

2. Description of the Reference Material (RM)

Name: U-[¹³C₁₇]-Aflatoxin B1 in acetonitrile

CAS number: 1162-65-8

Catalog number: B-MYC0210-1.2

Lot #: 114311S Expiry date: 28.07.2015

Starting material: U-[¹³C₁₇]-Aflatoxin B1, Lot #S08224Z,

Romer Labs Diagnostic GmbH

Physical description of RM: Solution of U-[¹³C₁₇]-Aflatoxin B1 in acetonitrile

Packaging and amount of RM: Amber Certan[®] glass ampoules, solution of 1.2 mL

Name and address of the manufacturer: Romer Labs Diagnostic GmbH

Technopark 1, 3430 Tulln, Austria www.romerlabs.com

Name and address of the supplier: LGC Standards GmbH

Mercatorstraße 51, 46485 Wesel, Germany Tel +49(0)2 81 98 87 0, Fax +49(0)2 81/98 87 199

www.lgcstandards.com

2.1 Intended use of the RM

- for laboratory use only
- internal standard [4, 5]

2.2 Instruction for the correct use of the RM

The ampoules should be stored at -20°C in a dark place. Before usage of the RM, the ampoules should be allowed to warm to room temperature. The recommended minimum sub-sample amount for all kinds of application is 100 μ L. 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 handling of this RM are available as safety data sheet.

Hazardous Ingredients Concentration in %

Acetonitrile > 99.9

Pictograms

Signal word Danger Hazard statement(s)

H225, H302, H312, H319, H332

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3. Certified values and their uncertainties

U-[¹³ C ₁₇]-Aflatoxin B1 in acetonitrile			
Compound	Mass concentration ^a		
	Certified value ^b	Uncertainty ^c	
U-[¹³ C ₁₇]-Aflatoxin B1, 99.3 atom % ¹³ C	0.500 μg/mL	± 0.026 μg/mL	

^a Values are based on preparation data and confirmed experimentally by HPLC-UV

3.1 Calculation of uncertainty

After the concentration of the gravimetric prepared solution was confirmed by HPLC-UV, the uncertainty of the calibrant solution was calculated on the basis of preparation [7].

Uncertainty components	Description	Standard uncertainty (u)	
Purity (P) of solid U-[¹³ C ₁₇]-Aflatoxin B1, 99.3 atom % ¹³ C	P = 99 ± 1 %	u (P) = 0.58 %	а
Weighing procedure weighted sample: m _{ws} = 1.262 mg	repeatability: 0.03 mg linearity: 0.012 mg	u (m) = 0.03 mg	b
Dilution procedure	calibration flask 1: 250 mL ± 0.15 mL	u (cal1) = 0.06 mL	С
volumetric flask 1: V _{f1} = 250 mL	repeatability flask 1: 0.03 mL	u (rep1) = 0.03 mL	d
volumetric flask 2: V ₁₂ = 250 mL	volume expansion solvent flask 1	u (Vol. exp.1) = 0.59 mL	е
one-mark glass pipette: V _p = 25 mL		u (V1) = 0.59 mL	f
	calibration flask 2: 250 mL ± 0.15 mL	u (cal2) = 0.06 mL	g
	repeatability flask 2: 0.03 mL	u (rep2) = 0.03 mL	h
	volume expansion solvent flask 2	u (Vol. exp.2) = 0.59 mL	i
		u (V2) = 0.59 mL	j
	calibration pipette: 25 mL ± 0.03 mL	u (cal3) = 0.01 mL	k
	volume expansion solvent pipette	u (Vol. exp.3) = 0.06 mL	I
		u (V3) = 0.06 mL	m

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

 $t_{i,l,m}$ All contributions are combined to give the $u(V) = \sqrt{u(cal)^2 + u(rep)^2 + u(Vol. \exp)^2}$

Calculation of the combined uncertainty \textbf{u}_{c} and the expanded standard uncertainty U

$$c_{Toxin} = \frac{10 \times m_{ws} \times P \times V_p}{V_{f1} \times V_{f2}} = \frac{10 \times 1.262 \times 99 \times 25}{250 \times 250} = 0.500 \ mg \ / L$$

$$\frac{u_{c}(c_{Toxin})}{c_{Toxin}} = \sqrt{\left[\frac{u(P)}{P}\right]^{2} + \left[\frac{u(m)}{m_{ws}}\right]^{2} + \left[\frac{u(V1)}{V_{f1}}\right]^{2} + \left[\frac{u(V2)}{V_{f2}}\right]^{2} + \left[\frac{u(V3)}{V_{p}}\right]^{2}} = \sqrt{\left[\frac{0.58}{99}\right]^{2} + \left[\frac{0.03}{1.262}\right]^{2} + \left[\frac{0.59}{250}\right]^{2} + \left[\frac{0.06}{25}\right]^{2}} = 0.025$$

$$u_c(c_{Toxin}) = c_{Toxin} \times 0.025 = 0.500 \times 0.025 = 0.013 \text{ mg} / L$$

Calculation of expanded standard uncertainty U using a coverage factor k = 2

$$U(c_{Toxin}) = u_c(c_{Toxin}) \times 2 = 0.013 \times 2 = 0.026 \ mg / L = 0.026 \ \mu g / mL$$

^b Mass concentration based on weighed amount, purity and dilution step

^c Expanded uncertainty U(k=2) of the value u_c according to GUM [6]

^b Estimation of this u-value is based upon the values for repeatability and linearity described in the user manual of the microbalance

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

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

^{e,l,l} Based on the density of 0.7857 g/cm³ at temperature $T = 20^{\circ}$ C and a maximum temperature variation of \pm 3°C, of volume expansion, relative volume expansion coefficient of acetonitrile is 1370 * 10^{6} /°C [8], volume expansion term (rectangular distribution) was divided by $\sqrt{3}$



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4. I sotopic enrichment and isotope pattern

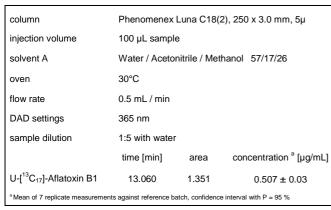
Isotope pattern ^a		
Compound	Isotopic distribution	
[¹³ C ₁₇]-Aflatoxin B1	88.4 %	
[¹³ C ₁₆]-Aflatoxin B1	10.6 %	
[¹³ C ₁₅]-Aflatoxin B1	1.0 %	
Calculated isotopic enrichment level ^a : 99.3 atom % ¹³ C		
^a Approximation based on LC-MS/MS data		

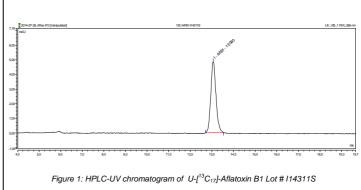
5. Discussion of traceability

This calibrant is certified on the basis of gravimetric preparation [7]. Thus the certified value (mass concentration of U-[13 C₁₇]-Aflatoxin B1, 99.3 atom % 13 C) is based on the weighed amount of the starting material and is therefore traceable to the stated purity of the solid mycotoxin. High purity material represents a practical realization of concentration units, through conversion of mass to molar quantity.

6. Confirmation of certified value by HPLC-UV

The certified concentration of U-[13 C₁₇]-Aflatoxin B1, 99.3 atom % 13 C of the gravimetric prepared solution was confirmed by HPLC-UV against an independently prepared reference batch of unlabeled Aflatoxin B1 calibrant Lot# L14163C.







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7. Further information

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

approved for release by: Dipl. Chem. Ulrich Berger date: 28.07.2014

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References:

- [1] ISO Guide 31, 1-7, (2000), "Reference Materials Contents of Certificates and Labels"
- [2] Eurachem / CITAC Guide, 1-37, (2003), "Traceability in Chemical Measurement"
- [3] Eurachem / CITAC Guide CG-4, 1-133, (2012), "Quantifying Uncertainty in Analytical Measurement "
- [4] G. Häubl, F. Berthiller, R. Krska, R. Schuhmacher, "Suitability of a fully ¹³C isotope labelled internal standard for the determination of the mycotoxin deoxynivalenol by LC-MS/MS without clean-up", Anal. Bioanal. Chem. **384** (3), (2006), 692-696
- [5] G. Häubl, F. Berthiller, J. Rechthaler, G. Jaunecker, E.M. Binder, R. Krska, R. Schuhmacher, (2006), "Characterization and application of isotope-substituted (13C₁₅)-deoxynivalenol (DON) as an internal standard for the determination of DON", Food Addit Contam. 23, (2006), 1187-1193
- [6] International Organization for Standardization (ISO), (1995), "Guide to the Expression of Uncertainty in Measurement", 1st Ed. Geneva, Switzerland
- [7] 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 Aflatoxin B1"
- [8] E.W. Flick, (1996), "Industrial Solvents Handbook", 3rd Ed., Noyes Data Corp. Westwood NJ