



CERTIFICATE OF ANALYTICAL DATA FOR MATERIALS INCLUDED IN TECHNICAL ANNEX OF ACCREDITATION N° 02/PMR003

# Certificate of analysis

Certified reference material code: CRM-03-CYN

Name: Certified Reference Material of Cylindrospermopsin Batch: 16-001

**Description:** This Certified Reference Material (CRM) is a solution of Cylindrospermopsin (CAS Number 143545-90-8) in deionized water.

**Intended use:** Calibration of equipment or a measurement procedure. Establishing metrological traceability. Method validation. Quality control of a measurement or measurement procedure. For laboratory use only.

**Certified values and uncertainties:** The certified concentration given below is based on results obtained from the gravimetric preparation of this solution previous quantity determination by <sup>1</sup>H-qNMR in Varian WB 500 MHz equipment (USC).

Concentration of Cylindrospermopsin (95% Confidence Interval)	(25.9 ± 1.2) μmol / kg
	$(10.8 \pm 0.5)  \mu g / g$

The starting material is measured by <sup>1</sup>H-qNMR against a Sucrose NIST certified reference material (SRM 17f) followed by gravimetric preparation using high precision balances calibrated with SI-traceable weights. Consequently the quantified values are traceable to the International System of Units (SI) via an unbroken chain of comparisons through validated methodology.

Homogeneity was evaluated by analysis of variance on 2% of stratified randomly selected ampoules over the entire batch (6 replicate analyses per ampoule). A possible heterogeneity is hidden by the method repeatability. Therefore a maximum possible heterogeneity was calculated as  $u_{bb}$ , 0.81%. The Stability studies after storage of selected units were performed at  $-20\degree$ C for a storage time of 12 months. Using the data from the long-term study, the uncertainty due to possible degradation was calculated as  $u_{lts}$ , 0.99%. Both studies were verified against an independently prepared calibration solution.

The components of Standard Uncertainty include related uncertainties due to characterization, heterogeneity, long term instability, short term instability (dispatch), and bulk assay, Table 1. The results are expressed as the certified value  $\pm$  the expanded uncertainty (calculated by combination of the squared contribution values). Estimated expanded uncertainty with a coverage factor k = 2, corresponding to a level of confidence of about 95 %, as defined in the Guide to the expression of uncertainty in measurement, ISO35.

The certificate is a summary of an extensive program of work involving selection and purification of the material, assessing its suitability and measuring the properties to be certified.

**Expiration of Certification:** This certificate of CRM-03-CYN, 16-001 batch is valid 12 months after purchase date within the measurement uncertainties specified, provided the CRM is handled and stored in accordance with the instructions given in this certificate. This validity may be extended if further evidence of stability becomes available.

Maintenance of Certification: The material will be subjected to regular Laboratorio CIFGA stability monitoring programme to control its further stability over the period of its





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certification. If substantive changes occur that affect the certification before the expiration date, Laboratorio CIFGA will notify the purchaser.

**Instructions for proper handling use:** This material contains toxic material, and should be handled with care. Read MSDS before using. Material Safety Data sheet for information regarding CRM-03-CYN is enclosed in the package. Use proper disposal methods.

Before opening an ampoule it must be guaranteed that the content is a liquid and properly mixed to ensure homogeneity. Sample aliquots for analysis should be withdrawn at 15  $^{\circ}$ C to 25  $^{\circ}$ C immediately after opening the ampoules and should be processed without delay for the certified value to be valid within the stated uncertainty. Ampoule should not remain open longer than few minutes due to solvent volatility and air contamination.

Storage conditions: The original sealed ampoules should be stored at temperatures below/at -20 °C (freezer) until use. Unopened ampoules should be stored upright under normal laboratory conditions inside the original container supplied.

**Source and Preparation of Material:** Cylindrospermopsin was obtained from commercial source, identity of the material was confirmed by <sup>1</sup>H-NMR and LC-MS/MS, Figures 1 and 2. Finally, 0.5 mL aliquots of toxin solution were dispensed into 2-mL amber argon-filled glass ampoules, which were then flame sealed.

\* Non-certified and non-accredited data: Taking into account the density of water at 20 °C ( $\rho^{20} = 0.9982 \text{ g/mL}$ ), non-certified volumetric concentration values are:

Concentration of Cylindrospermopsin (CYN)	(25.9 ± 1.2) μM
(95% Confidence Interval)	(10.8 ± 0.5 μg / mL

Concentration of Cylindrospermopsin and impurities are neglected in order to apply solution density value.

Purity analysis was performed by qNMR and UPLC MS/MS. Both analyses indicated that there was no significant contribution from other analogues, only 7-deoxy-CYN was determined <0.1%. Results of nuclear magnetic resonance (NMR) analysis were consistent with structure and a molar ratio purity of principal components  $\geq$  99 %. No purity correction over certified concentration should be done.

## **Additional lectures:**

- 1. UNE-EN ISO 17034 General Requirements for the competence of reference material producers.
- 2. ISO Guide 35 Reference materials Guidance for characterization and assessment of homogeneity and stability.





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## **Figures**

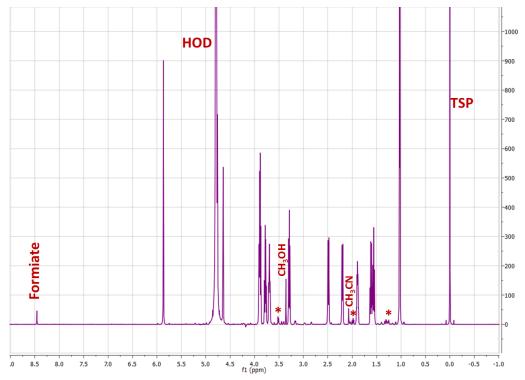


Figure 1.  $^{1}$ H-NMR spectrum of Cylindrospermopsin solution, Varian 500 MHz, solution of CYN (2.8 mM) in  $D_2O$  used in preparation of CRM-03-CYN batch 16-001.  $^{*}$ No identified impurities are highlighted. As the structure of these impurities is not known, their exact amount cannot be calculated. Because the impurities usually are smaller molecules (otherwise, more NMR signals would arise), it can be estimated from the corresponding integrals, but this purity is not relative to close-related impurities to cylindrospermopsin. analogues.

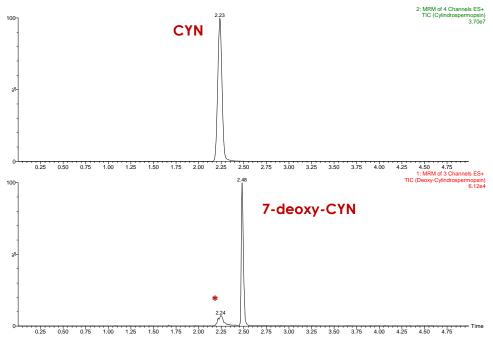


Figure 2. Analysis of the CRM-03-CYN solution by UPLC-MS/MS. Conditions: ACQUITY UPLC HSS T3 1.7μm, 100x2.1 mm, UPLC column. Elution conditions, gradient mode with mobile phases: (A) Water and (B) acetonitrile, both with 0.1% formic acid; Gradient: 0 min 2% B; 1.0 min 2% B; 2.6 min 30% B; 3.6 min 30% B; 3.7 min 2% B; 5.0 min 2% B; flow rate: 0.4 mL/min. Temperature: 35°C. MS detector: Waters Quattro Premier XEVO using MRM transitions of CYN and 7-deoxy-CYN. \* Isotopic shadow from low intensity MRM transitions of CYN.





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Table 1. Uncertainties contribution for the certification of CRM-03-CYN, 16-001. Char: qnmr quantification; ba: bulk assay; bb: homogeneity test; lts: long term stability test; sts: short term stability test.

#### **Uncertainty values**

Uncertainty contribution parameter	CYN
u <sub>char</sub> /%	1.90
$u_{ba}/\%$	0.25
$u_{bb}/\%$	0.81
u <sub>lts</sub> /%	0.99
Expanded uncertainty (k=2) / %	4.61

Short-term stability study, test for slope significance, has been carried out to simulate the conditions of transport. Packaged vials were exposed for 12 days, 9 days, 6 days, 3 days and 0 hr (-80 °C, control), with an isochronous design being used to enable subsequent LC-MS/MS analysis of CRMs to be conducted under repeatability conditions. After 12 days at 37 °C - 40°C temperature results indicated continued stability. With these conditions, no change is expected in the property values of the CRM. The short-term stability uncertainty ( $u_{sts}$ ) was not considered relevant as an uncertainty component and it was not included in the  $u_{CRM}$  uncertainty calculation.

Table 2. Molecular information.

Name	CYLINDROSPERMOPSIN
CAS NUMBER	143545-90-8
Molecular formula	$C_{15}H_{21}N_5O_7S$
Molecular Weight	415.4 g·mol <sup>-1</sup>
Monoisotopic molecular weight	415.1 g·mol <sup>-1</sup>
2D structure	O O O O O O O O O O O O O O O O O O O





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Inquiries concerning this Reference Material should be directed to:

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## RM PRODUCER'S APPROVING OFFICER:

Certification Manager: Álvaro Antelo

This Certificate is only valid if the product was obtained directly from Laboratorio CIFGA S.A. or one of our authorized distributors.

**Certificate Revision History:** July, 29<sup>th</sup> 2016 (Original certificate issue date, revision 1).

February,  $07^{th}$  2018. Minor Editorial changes. Uncertainty update following long term stability testing (certificate rev. 2).

February, 20th 2018. Correction of typographical mistake on the date

(certificate rev. 3).

January, 15<sup>th</sup> 2019. Intended use update. Editorial changes. No attempt was made to re-evaluate the certificate values or any technical data presented on this certificate (certificate rev. 4).

June,  $10^{th}$  2020. Correction of source of material section. No attempt was made to re-evaluate the certificate values or any technical data presented on this certificate. Company address update (certificate rev. 5).

December, 02<sup>nd</sup> 2020. Introduction of the accreditation mark. Editorial changes. No attempt was made to re-evaluate the certificate values or any technical data presented on this certificate (certificate rev. 6).

NOTE: This certificate of analysis shall not be reproduced, except in full, without written approval of Laboratorio CIFGA S.A.