

Certificate of analysis

Certified reference material code: CRM-00-FB2

Name: Certified Reference Material of Fumonisin B2.

Batch: 25-001

Description: This Certified Reference Material (CRM) is a solution of Fumonisin B2 (CAS Number 116355-84-1) in acetonitrile - water, 50% (v/v) solution.

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 ¹H-qNMR in VARIAN 500 WB equipment (Universidad de Santiago de Compostela).

Concentration of Fumonisin B2 (95% Confidence Interval)	(156 ± 10) µmol / kg
	(110 ± 7) µg / g

The starting material is measured by ¹H-qNMR against a Benzoic acid standard for quantitative NMR, certified reference material 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} , 0.80%. The Stability studies after storage of selected units were performed at -20°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} , 2.19%. 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 ± 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, ISO33405.

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-00-FB2, 25-001 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.*



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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 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-00-FB2 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 handled at 15°C to 25°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 five 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: Fumonisin B2 used in the preparation of this CRM was isolated from fungal culture of *F. proliferatum*. The solution was prepared by weighing and mixing FB2 into acetonitrile - water until completely homogenization. Identity of the material was confirmed by ¹H-NMR and LC-MS/MS. Finally, 1.0 mL aliquots of FB2 solution were dispensed into 2-mL amber argon-filled glass ampoules, which were then flame sealed and ready to use for calibration.

Non-certified and non-accredited data:

Taking into account the density of acetonitrile, 50% (v/v) aqueous solution at 20 °C ($\rho_{ACN\ 50\% V:V}^{20^{\circ}C} = 0.9109\ g/mL$), non-certified volumetric concentration values are:

Concentration of Fumonisin B2 (95% Confidence Interval)	(142 ± 9) µM
	(100 ± 6) µg / mL

NOTE: concentration of Fumonisin B2 and impurities are neglected in order to apply solution density value.

Purity analysis was performed by combining ¹H-qNMR and UPLC MS/MS, ≥ 97 % (molar ratio purity). Low amounts of Fumonisin B2 analogues quantified out from principal component are presented, see #Table 2. *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 33405 *Reference materials - Approaches for characterization and assessment of homogeneity and stability.*



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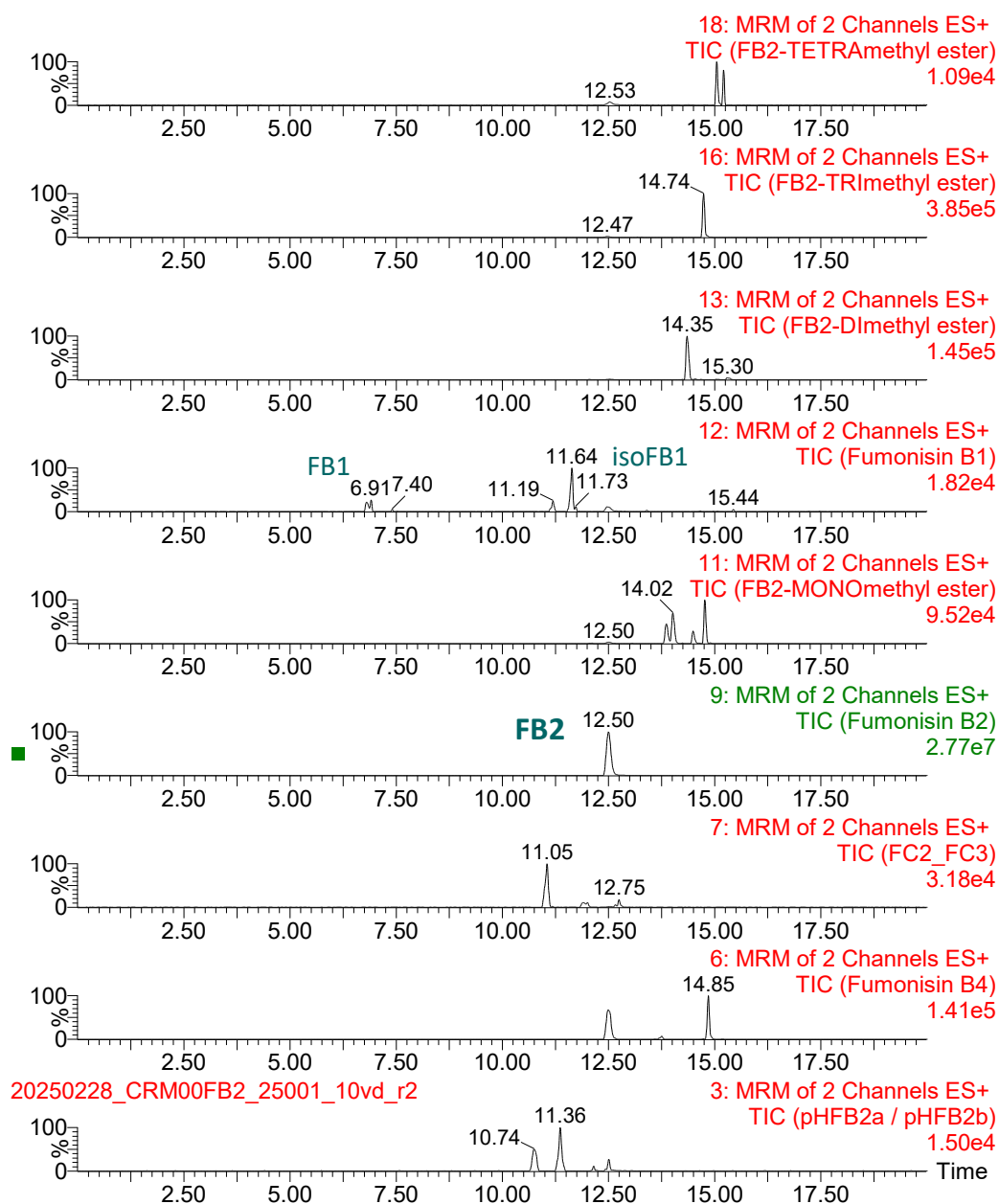


Figure 1. Analysis of the CRM-00-FB2 (25-001) by UPLC-MS/MS. Conditions: ACQUITY UPLC HSS T3 1.7 μ m, 100x2.1 mm, UPLC column. Elution conditions, gradient mode with mobile phases: (A) Water and (B) Acetonitrile, both with 0.1% formic acid (v/v); Gradient: 0 min 25% B; 2.00 min 25% B; 12.5 min 35% B; 14.0 min 40% B; 17.0 min 90% B; 18.0 min 90% B; 18.1 min 25% B; 20.0 min 25% B; flow rate: 0.4 mL/min. Temperature: 40°C. MS detector: Waters XEVO TQ-MS. FB2 MME isobaric impurities are assigned as MM1-MM4 based on increasing retention time, and the tentative isoFB1 analogues are correspondingly designated isoFB1a to isoFB1c in retention-time order.



Table 1. Uncertainties contribution for the certification of **CRM-00-FB2 (25-001)**. *Char*: *qNMR quantification*; *ba*: *bulk assay*; *bb*: *homogeneity test*; *lts*: *long term stability test*; *sts*: *short term stability test*.

Uncertainty contribution parameter	Uncertainty values
	Fumonisin B2
$u_{char}/\%$	2.06
$u_{ba}/\%$	0.79
$u_{bb}/\%$	0.80
$u_{lts}/\%$	2.19
$u_{sts}/\%$	no significant contribution
Expanded uncertainty ($k=2$) / %	6.42

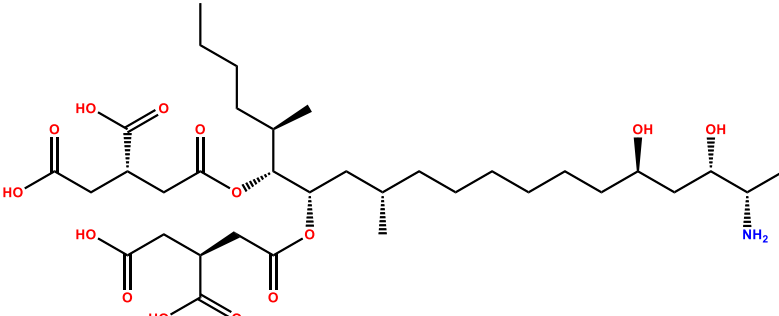
Short-term stability study. The uncertainty due to transport instability was estimated from the test for slope significance at 40 °C for 9 days. The results indicated continued stability, no change is expected in the property values of the CRM within specifications.

Table 2. Non-certified reference values of impurities concentration (μM) determined by UPLC MS/MS. Estimated uncertainty corresponds to a level of confidence in terms of one standard deviation. The impurity concentration was determined using the equimolar response factor, employing the same fragmentation pattern for all components to ensure methodological consistency.

Toxin	Concentration / μM	%
FB2	142	$\geq 97\%$
FB1	-	<0.1%
pHFB2a	-	<0.1%
pHFB2b	-	<0.1%
FB2 monomethylester 1 (FB2 MME1)	0.14 ± 0.01	0.1%
FB2 monomethylester 2 (FB2 MME2)	0.15 ± 0.01	0.1%
FB2 monomethylester 3 (FB2 MME3)	-	0.1%
FB2 monomethylester 4 (FB2 MME4)	0.21 ± 0.02	0.15%
FB2 dimethylester 1 (FB2 DME1)	0.46 ± 0.04	0.32%
FB2 dimethylester 2 (FB2 DME2)	-	<0.1%
FB2 trimethylester (FB2 TME1)	0.77 ± 0.06	0.54%
FB2 tetramethylester (FB2 FME)	-	<0.1%
FB4	1.21 ± 0.10	0.84%
FC2 or FC3	0.21 ± 0.02	0.15%
isoFB1a	-	<0.1%
isoFB1b	0.15 ± 0.01	0.1%
isoFB1c	-	<0.1%



Table 3. Molecular information.

Name	Fumonisin B2
CAS NUMBER	116355-84-1
Molecular formula	C ₃₄ H ₅₉ NO ₁₄
Molecular Weight	705.84 g·mol ⁻¹
Monoisotopic molecular weight	705.39 g·mol ⁻¹
2D structure	

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:

March, 10th 2026 (Original certificate issue date, revision 1).

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



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