

**CERTIFICATE OF ANALYSIS - Certified Reference Material**

ISO 17034

Reference Material

This certificate has been created in compliance with ISO/IEC 17034. The reference material has been developed, manufactured, and validated in accordance with ISO/IEC 17025, ISO/IEC 17034, and a certified quality management system. LVA GmbH is accredited by the Austrian accreditation body as a recognized reference material producer in line with ISO/IEC 17034.

Product Name	Item Number	Lot Number	Format	Volume	Expiry Date	Storage Temp
Fusarium Mix in Acetonitrile	VE00008364	FRM202503001	Multicomponent Solution	2,5 mL	4 Sep 2026	≤ -18 °C

CERTIFIED							
Compound Name	Concentration (µg/mL)	Expanded Uncertainty U (µg/mL)	CAS Number	Lot Number	Purity (%)	Amount (mg)	RT (min)
Deoxynivalenol	100,0	3,83	C15H20O6	DONLVA01	93,0	10,8	2,1
Zearalenon	100,0	3,39	C18H22O5	ZONLVA01	98,5	10,2	8,8
HT-2 Toxin	100,2	3,09	C22H32O8	HT2LVA02	94,5	10,6	7,6
T-2 Toxin	100,4	3,01	C24H34O9	T2LVA01	98,2	10,2	8,4

Metrological Traceability

This mycotoxin standard is produced gravimetrically for precise concentration determination, calculated by dividing the mass of the raw material, adjusted for purity, by the solvent volume to yield a final concentration in µg/mL. Raw materials, with a minimum purity of 90% verified via q-NMR by an external ISO/IEC 17025-accredited lab, are weighed on ISO/IEC 17025-compliant precision balances using traceable weights. Solvents are stabilized at room temperature monitored by traceable thermometers. All calibration and measurement protocols are documented to ensure traceable and reproducible results, with final processing at the LVA site in Klosterneuburg.

Associated Uncertainty

The mathematical representation of the u_{CRM} calculation is as follows:

$$u_{CRM} = \sqrt{u_{prep}^2 + u_{bb}^2 + u_{lts}^2 + u_{sts}^2}$$

u_{CRM} is calculated from the uncertainty contributions of the gravimetric preparation (u_{prep}), the measurement between units (u_{bb}), the long-term stability (u_{lts}), and the short-term stability (u_{sts}). This combined uncertainty is then expanded by a factor of $k = 2$, resulting in the expanded uncertainty U , which accompanies the certified value as a \pm indication. The summation of uncertainty values is done in absolute terms (in µg/mL).

Health and Safety Information

All chemical reference materials should be treated as potentially hazardous and utilized exclusively by qualified laboratory personnel. Please consult the Safety Data Sheet for comprehensive details regarding any hazards and the necessary precautions to be observed.

CERTIFIED BY

Dr. D. Steiner

CERTIFIED ON

5 Jun 2025

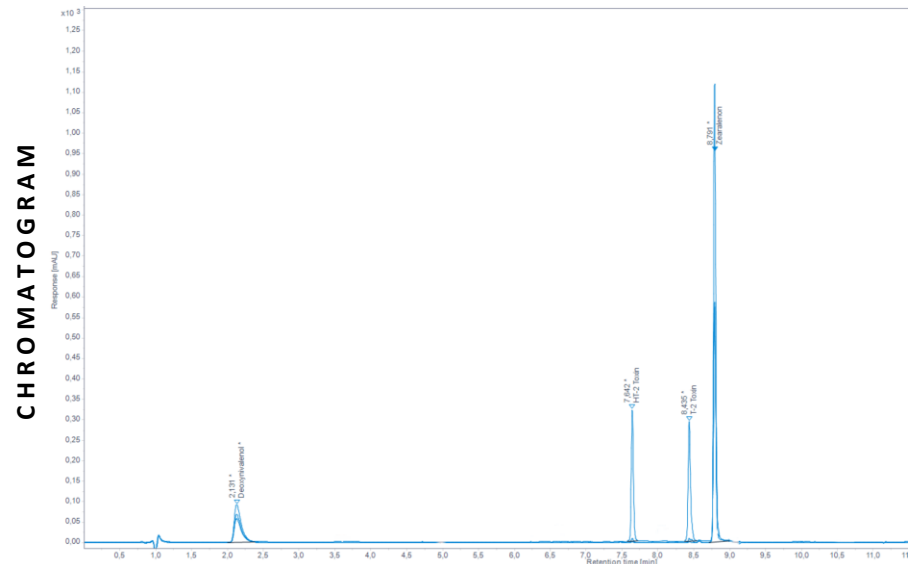
VERSION

FM-LVA-0519
Rev. 3.0

* Head of Department



LVA GmbH
Magdeburggasse 10, 3400 Klosterneuburg, AUSTRIA
referenzmaterialien@lva.at



Instrument

HPLC-System Agilent 1200

Detection

DAD

Column

Kinetex 2,6 μ m, 100 x 3 mm

Method Details

Eluent A: H₃PO₄ 0,1%

Eluent B: Acetonitrile

Inj.-Vol.

20 μ l

Flow

0,5 ml/min

Measurement Method

The assigned value is derived from a purity determination and gravimetrically prepared value. Purity was determined by an external ISO/IEC 17025 laboratory following q-NMR analysis.

Batch Information

Solvent: Acetonitrile, Lot No. 24I104031, 100 mL.

Method Details:

Eluent A: H₃PO₄ 0,1%

Eluent B: Acetonitrile

Eluent A->B: 14 min

Oven 35 °C

Sample dilution 1:10 with water

DAD setting

23°C room temperature

Homogeneity Assessment

Homogeneity was evaluated following the guidelines outlined in ISO 33405:2024. Random stratified sampling protocol was employed to sample completed units. The chemical analysis results were subsequently subjected to Single Factor Analysis of Variance (ANOVA). The uncertainty attributed to homogeneity was determined from the ANOVA and no heterogeneity was observed.

Stability Assessment

The stability assessment includes both short-term and long-term evaluations, simulating transportation and storage conditions for the sales units. The short-term stability study was conducted at room temperature over a period of 5 weeks, while the long-term stability was assessed according to the specified storage conditions until the expiry date. Additionally, long-term monitoring is conducted for each released batch throughout its shelf life.

Intended Use

Designed for utilization in a laboratory setting as a standard for calibration and quality control, or for method development in analytical techniques. Not intended for drug, household, or other applications.

Storage

The reference material should be kept in its original sealed container at the specified temperature.

Instructions for Handling and correct Use

The internal pressure of the container may vary slightly from the atmospheric pressure at the user's location. Open the container slowly and with caution to prevent material dispersion.

Procedure in Scope of Accreditation

This certificate has been issued in accordance with the procedure documented in DK-LVA-0471 "Referenzmaterialproduktion", with the issue date of 27 November 2024, as defined within the scope of accreditation.