

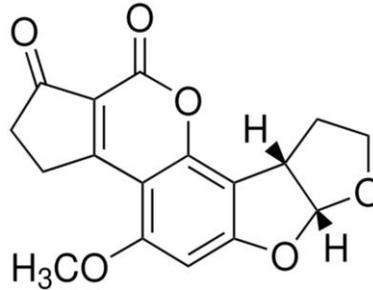
**Reference Material**

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**

**Aflatoxin B2 1 µg/mL in Acetonitrile, Solution of Aflatoxin B2 in Acetonitrile**

**Item Number** VE00008355  
**Lot Number** AB2202503001  
**CAS No.** 7220-81-7  
**Mol. Weight** 314.29 g/mol  
**Mol. Formula** C17H14O6  
**Expiry Date** 04.09.2026  
**Storage Temp** ≤ -18 °C  
**Format** Solution  
**Volume (ml)** 2.5



<b>CERTIFIED</b> Concentration 1,0 µg/mL	<b>CERTIFIED</b> Expanded Uncertainty (U) 0,034 µg/mL
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**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

**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).

**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.

<b>CERTIFIED BY</b>	<b>CERTIFIED ON</b>		<b>VERSION</b>
Dr. D. Steiner	5 Jun 2025		FM-LVA-0516 Rev. 3.0

\* Head of Department

