



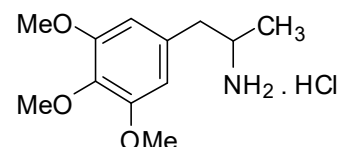
# CERTIFIED REFERENCE MATERIAL CERTIFICATE OF ANALYSIS

## NMIA D778: ( $\pm$ )-3,4,5-Trimethoxyamphetamine hydrochloride

Report ID: D778.2026.01

Chemical Formula:  $C_{12}H_{19}NO_3 \cdot HCl$

Molecular Weight: 261.7 g/mol (HCl) 225.3 g/mol (base)



### Certified value

Batch No.	CAS No.	Purity (mass fraction)
02-D-34	5688-80-2 (HCl) 1082-88-8 (base)	98.5 $\pm$ 0.5%

The uncertainty has been calculated according to ISO Guide 35 and is stated at the 95% confidence limit ( $k = 2$ ).

**IUPAC name:** 1-(3,4,5-Trimethoxyphenyl)-2-propanamine hydrochloride

**Expiration of certification:** The property values are valid till 20 February 2036, ten years from the date of re-certification provided the **unopened** material is handled and stored in accordance with the recommendations below. The material as issued in the unopened container and stored as recommended below should be suitable for use beyond this date, subject to confirmation of batch stability from the issuing body. The expiry date/shelf life does not apply to sample bottles that have been opened. In such cases it is recommended that the end-user conduct their own in-house stability trials.

**Description:** Off-white powder prepared by synthesis, certified for identity and purity by NMI Australia. Packaged in amber glass bottles with a septum and crimped aluminium cap or screw top cap.

**Intended use:** This certified reference material is suitable for use as a primary calibrator.

**Instructions for use:** Equilibrate the bottled material to room temperature before opening.

**Recommended storage:** When not in use this material should be stored at or below 25 °C in a closed container in a dry, dark area.

**Metrological traceability:** The certified purity value is traceable to the SI unit for mass (kg) through Australian national standards via balance calibration. In the mass balance approach all impurities are quantified as a mass fraction and subtracted from 100%.

**Stability:** This material has demonstrated stability over a minimum period of five years. The measurement uncertainty at the 95% coverage interval includes a stability component which has been estimated from annual stability trials. The long-term stability of the compound in solution has not been examined.

**Homogeneity assessment:** The homogeneity of the material was assessed using purity assay by GC-FID on ten randomly selected 1-2 mg sub samples of the material. The material was judged to be sufficiently homogeneous at this level of sampling as the variation in analysis results between samples was not significantly different at a 95% confidence level from that observed on repeat analysis of the same sample.

**Caution:** Treat as a hazardous substance. Use appropriate work practices when handling to avoid skin or eye contact, ingestion or inhalation of dust. Refer to the provided safety data sheet.

S. R. Davies

Dr Stephen R. Davies,  
Team Leader,  
Chemical Reference Materials, NMI.  
3 March 2026

This report supersedes any issued prior to 03 March 2026.

NATA Accreditation No. 198 / Corporate Site No. 14214.

**CIPM MRA notice:** This certificate is consistent with the capabilities that are included in Appendix C of the CIPM MRA drawn up by the CIPM. Under the CIPM MRA, all participating institutes recognize the validity of each other's calibration and measurement certificates for the quantities, ranges and measurement uncertainties specified in Appendix C. The "CIPM MRA Logo" and this statement attest only to the measurement(s) applied for determining the certified values on the certificate (for details see <http://www.bipm.org>).

**Legal notice:** Terms and Conditions associated with the provision of this reference material can be found on the NMIA website.

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### Characterisation Report:

The identity was confirmed by a range of spectroscopic techniques, NMR, IR and MS. The certified purity value was obtained by mass balance from a combination of traditional analytical techniques, including GC-FID, thermogravimetric analysis, Karl Fischer analysis and <sup>1</sup>H NMR spectroscopy. The purity value is calculated as per Equation 1.

$$\text{Purity} = (100 \% - I_{\text{ORG}}) \times (100 \% - I_{\text{VOL}} - I_{\text{NVR}}) \quad \text{Equation 1}$$

$I_{\text{ORG}}$  = Organic impurities of related structure,  $I_{\text{VOL}}$  = volatile impurities,  $I_{\text{NVR}}$  = non-volatile residue.

Supporting evidence is provided by elemental microanalysis.

GC-FID: Instrument: Agilent 6890N and 8890  
 Column: HP-1, 30 m × 0.32 mm I.D. × 0.25 μm  
 Program: 60 °C (1 min), 10 °C/min to 200 °C, 40 °C/min to 300 °C (5 min)  
 Injector: 200 °C  
 Detector Temp: 320 °C  
 Carrier: Helium  
 Split ratio: 20/1

Relative mass fraction of main component as the free base:  
 Initial analysis: Mean = 99.1%, s = 0.02% (10 sub samples in duplicate, December 2002)  
 Re-analysis: Mean = 98.8%, s = 0.03% (5 sub samples in duplicate, November 2008)  
 Re-analysis: Mean = 98.7%, s = 0.04% (5 sub samples in duplicate, November 2011)  
 Re-analysis: Mean = 99.0%, s = 0.02% (5 sub samples in duplicate, November 2016)  
 Re-analysis: Mean = 98.9%, s = 0.06% (9 sub samples in duplicate, February 2026)

Thermogravimetric analysis: Volatiles content and non-volatile residue < 0.3% mass fraction (December 2002 & November 2005)

Karl Fischer analysis: Moisture content is < 0.2% mass fraction (November 2008, 2011, September 2016, and February 2026)

### Spectroscopic and other characterisation data

GC-MS:	Instrument: HP5890/5971 Column: Zebtron ZB-5, 30 m x 0.25 mm I.D. x 0.25 µm Program: 60 °C (1 min), 10 °C/min to 100 °C, 15 °C/min to 250 °C (3 min) Injector: 200 °C Transfer line temp: 280 °C Carrier: Helium, 1.0 mL/min Split ratio: 20/1
	The retention time of the free base is reported with the major peaks in the mass spectrum. The latter are reported as mass/charge ratios and (in brackets) as a percentage relative to the intensity of the base peak. Free base (11.9 min): 225 (M <sup>+</sup> , 2), 182 (100), 167 (28), 148 (6), 139 (5) <i>m/z</i>
TLC:	Conditions: Kieselgel 60F <sub>254</sub> (Methanol/Hexane/NH <sub>3</sub> ) 20:5:1 Single spot observed, R <sub>f</sub> = 0.3
IR:	Instrument: FT-IR, Biorad WIN FTS3000MX Range: 4000 - 400 cm <sup>-1</sup> , KBr Peaks: 2929, 2715, 2063, 1587, 1512, 1425, 1242, 1125 cm <sup>-1</sup>
<sup>1</sup> H NMR:	Instrument: Bruker DMX-600 Field strength: 600 MHz Solvent: CH <sub>3</sub> OD Spectral data: δ 1.36 (3H, d, <i>J</i> = 6.6 Hz), 2.85 (1H, dd, <i>J</i> = 7.6, 13.6 Hz), 2.95 (1H, dd, <i>J</i> = 6.7, 13.6 Hz), 3.61 (1H, m), 3.78 (3H, s), 3.88 (6H, s), 6.61(2H, s) ppm Residual isopropanol quantified between 0.3 – 0.5% mass fraction (December 2002, May 2006 and February 2026)
<sup>13</sup> C NMR:	Instrument: Bruker DMX-600 Field strength: 150 MHz Solvent: D <sub>2</sub> O Spectral data: δ 18.6, 42.0, 48.9, 49.1, 50.3, 56.7, 61.0, 107.6, 133.4, 138.2, 154.8 ppm
Melting point:	217-219 °C
Microanalysis:	Found: C = 55.3%; H = 7.9%; N = 5.3% Calculated: C = 55.1%; H = 7.7%, N = 5.4% (Calculated for C <sub>12</sub> H <sub>19</sub> NO <sub>3</sub> .HCl)