

Certificate of Analysis Reference Material

Lipomed Document QC-CA-94 Version: 007-03.Jun.2020

Supersedes: 006-14.Nov.2017

Chemical name: d,I-MDMA.HCI

(RS)-1-(Benzo[d][1,3]dioxol-5-yl)-N-methylpropan-2-amine.hydrochloride

Lot No: 94.3B11.1 Release date: May 25, 2020 Art. No: MDM-94-HC Last testing date: N/A

Retest date: May 2025

Chemical formula: $C_{11}H_{15}NO_2$ Molwt: 193.25

Hydrochloride 229.70

CAS Registry No: 64057-70-1

TEST	SPECIFICATION	RESULT	
1. Appearance	white to off-white crystalline powder	conforms	
2. Purity HPLC > 98.5 % 99.950 ± 0.050 % The purity is calculated from the distribution of 6 HPLC analyses with a 95% level of confidence.			
3. Free base content (Corrected from purity and water	> 81.9 % er)	83.8 %	
4. Optical rotation	N/A (racemic mixture)	N/A	
5. Water content	Karl Fischer < 1 %	0.3 %	
6. Calculated hydrochloride content		15.9 %	

Storage Conditions: For maximum stability store air-tight at 2 - 8°C in a dark location.

Lipomed certifies that this standard meets the specifications stated in this certificate and warrants this product to meet the stated acceptance criteria through the retest date when stored unopened as recommended.

FOR ANALYTICAL PURPOSES ONLY: NOT FOR HUMAN OR ANIMAL USE!

Issued by Dr. L. Prévot

Date sign: Arlesheim,

June 03, 2020

LIPOMED AG Fabrikmattenweg 4 4144 Arlesheim Switzerland Tel. +41 61 702 02 00 Fax +41 61 702 02 20 HPOMED GmbH Hegenheimer Str. 2 79576 Weil am Rhein Germany

+49 7621 1693 474

+49 7621 1693 473

LIPOMED Inc. 150 Cambridgepark Drive, Suite 705 Cambridge, MA 02140 U.S.A. +1 (617) 577 7222

www.lipomed.com +1 (617) 577 1776 lipomed@lipomed.com









Certificate of Analysis Reference Material

Lipomed Document QC-CA-94 Version: 007-03.Jun.2020

Supersedes: 006-14.Nov.2017

Chemical name: d,I-MDMA.HCI

(RS)-1-(Benzo[d][1,3]dioxol-5-yl)-N-methylpropan-2-amine.hydrochloride

Lot No: 94.3B11.1 Release date: May 25, 2020 Art. No: MDM-94-HC Last testing date: N/A

Retest date: May 2025

Chemical formula: C₁₁H₁₅NO₂ Molwt: 193.25

Hydrochloride 229.70

CAS Registry No: 64057-70-1

TEST	SPECIFICATION	RESULT
7. Identity	IR	IR corresponds
	UV: in methanol $ \lambda_{\text{max},1} = 203.0 \pm 1.0 \text{ nm} \\ \epsilon_{\text{mol},1} = 32000 \pm 3000 $	UV: in methanol $\lambda_{\text{max},1} = 203.9 \text{ nm}$ $\epsilon_{\text{mol},1} = 31473$
	$\begin{array}{l} \lambda_{\text{max,2}} = 236.0 \pm 1.0 \text{ nm} \\ \epsilon_{\text{mol,2}} = 4200 \pm 500 \end{array}$	$\lambda_{\text{max},2} = 236.0 \text{ nm}$ $\epsilon_{\text{mol},2} = 4010$
	$\begin{array}{l} \lambda_{\text{max,3}} = 287.0 \pm 1.0 \text{ nm} \\ \epsilon_{\text{mol,3}} = 4200 \pm 500 \end{array}$	$\begin{array}{l} \lambda_{\text{max,3}} = 286.1 \text{ nm} \\ \epsilon_{\text{mol,3}} = 4020 \end{array}$
	Melting point: 149 \pm 3 °C	151 - 152 °C
	Proton NMR	corresponds to structure
	Mass Spectroscopy (ESI*)	MH ⁺ corresponds

+1 (617) 577 1776









GENERAL INFORMATION

Quality Documentation:

This certificate is designed in accordance with ISO Guide 31 (Reference Materials - Contents of Certificates and Labels) and ISO Guide 35 (Reference Materials - General and Statistical Principles for Certification).

Quality Standards for Arlesheim production site:

ISO 9001 Quality Management System. Manufacturing, analysis, packaging and distribution of

Analytical Reference Materials and Pharmaceuticals. IQNet/SQS Certification: 37199

ISO/IEC 17025 General requirements for the competence of Testing Analytical Reference Standards.

ANAB Certificate number: AT-1760

ISO 17034 General requirements for the competence of Reference Material Producer.

ANAB Certificate number: AR-1761

Quality Control Assessment:

The product quality is controlled by regularly performed quality control tests (retests).

Intended Use:

The product covered by this certificate is designed for calibration or for use in quality control procedures for the specified chemical compound listed page 1. This product can be used for quantification and/or identification. If dilution is required use only diluent compatible with all certified analyses in this preparation. All solutions should be thoroughly mixed prior to

Expiration/Retest dates:

Expiration date/Retest date of the unopened flask stored at the recommended storage condition is the last day of the month listed page 1.

A retest will be performed 6 months prior to the stated retest date. Upon successful retesting, a new retest date or expiration date is set for the product. The certificate of analysis is then updated and made available on our website. For our products, the extension of the shelf life is capped to 10 years after release as the maximum period.

Gravimetric preparation:

All balances are calibrated annually by an ISO/IEC 17025 accredited calibration service. Calibration verification is performed weekly with certified traceable weights. Each balance has been assigned a minimum weighing.

Purity:

- Purity and/or chemical identity are determined by one or more of the following techniques: HPLC/UV, GC/FID, LC/MS, IR, UV, 1H NMR, Karl Fischer, melting point and optical rotation if applicable
- Purity of isomeric compounds is reported as the sum of the isomers
- Purity values are rounded to the last decimal place given

Uncertainty Statistics and Confidence limits:

The uncertainties are determined in accordance with ISO 17034 and ISO/IEC 17025. Uncertainty is given for a minimum injection volume of 1 μl. The certified uncertainty value (including characterization uncertainty, homogeneity between ampoules uncertainty, storage stability uncertainty and shipping stability uncertainty) is combined using the following formula:

$$U(y) = \sqrt{U_{characterization}^2 + U_{homogeneity}^2 + U_{storage \, stability}^2 + U_{shipping \, stability}^2}$$

The packaged amount is the minimum sample size for which uncertainty is valid.

Stability:

The manufacturer guarantees the stability of this product throughout its intended shelf life, when handled and stored accordingly to the given storage conditions.

Legal and Safety Notice:

This product is for routine laboratory analysis and research purposes only. Due to the hazardous nature, only trained personnel should handle this product. The General Terms and Conditions of Lipomed apply.

lipomed@lipomed.com

LIPOMED AG Fabrikmattenweg 4 4144 Arlesheim Switzerland Tel. +41 61 702 02 00 Fax +41 61 702 02 20 LIPOMED GmbH Hegenheimer Str. 2 79576 Weil am Rhein Germany

+49 7621 1693 473 +49 7621 1693 474

LIPOMED Inc. 150 Cambridgepark Drive, Suite 705 Cambridge, MA 02140 +1 (617) 577 7222 www.lipomed.com

+1 (617) 577 1776





