



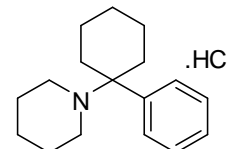
REFERENCE MATERIAL PRODUCT INFORMATION SHEET

NMIA D748b: Phencyclidine hydrochloride

Report ID: D748b.2020.03 (Bottled 170705)

Chemical Formula: C₁₇H₂₅N.HCl

Molecular Weight: 279.9 g/mol (HCl), 243.4 g/mol (base)



Property value

Batch No.	CAS No.	Purity estimate
13-D-25	956-90-1 (HCl) 77-10-1 (base)	99.4 ± 0.9%

IUPAC name: 1-(1-Phenylcyclohexyl)piperidine hydrochloride (1:1).

Expiration of certification: The property values are valid till 22 May 2025, i.e. five 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: White solid prepared by synthesis, and certified for identity and purity by NMIA. Packaged in amber glass bottles with a septum and crimped aluminium cap or screw top cap.

Intended use: This reference material is recommended for qualitative analysis only.

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.

Stability: In the absence of long term stability data the stability of this material has been judged from stability trials conducted on similar materials by NMI Australia over the last ten years. This material has demonstrated stability over a minimum period of three years. 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.

Safety: 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.
13 October 2022

This report supersedes any issued prior to 15 September 2022.

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

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

Characterisation Report:

The identity was confirmed by a range of spectroscopic techniques, NMR, IR and MS. Impurities of related structure were assessed by GC-FID. The purity estimate was obtained by mass balance from a combination of traditional analytical techniques, including GC-FID, thermogravimetric analysis, Karl Fischer analysis and ¹H 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_{ORG} = Organic impurities of related structure, I_{VOL} = volatile impurities, I_{NVR} = non-volatile residue.

Supporting evidence is provided by qualitative headspace GC-MS analysis of occluded solvents and elemental microanalysis.

GC-FID: Instrument: Varian CP-3800
 Column: VF-1MS, 30 m × 0.32 mm I.D. × 0.25 μm
 Program: 100 °C (1 min), 10 °C/min to 220 °C, 30 °C/min to 300 °C (3 min)
 Injector: 200 °C
 Detector Temp: 320 °C
 Carrier: Helium
 Split ratio: 20/1
 Relative peak area of the main component as the free base:
 Initial analysis: Mean = 99.8%, s = 0.08% (10 sub samples in duplicate, October 2013)
 Re-analysis: Mean = 99.8%, s = 0.009% (5 sub samples in duplicate, June 2017)
 Re-analysis: Mean = 99.9%, s = 0.003% (6 sub samples in duplicate, May 2020)

GC-FID: Instrument: Varian CP-3800
 Column: VF-1MS, 30 m × 0.32 mm I.D. × 0.25 μm
 Program: 120 °C (1 min), 10 °C/min to 220 °C, 30 °C/min to 300 °C (3 min)
 Injector: 250 °C
 Detector Temp: 320 °C
 Carrier: Helium
 Split ratio: 20/1
 Relative peak area of the main component as the free base:
 Initial analysis: Mean = 99.7%, s = 0.02% (9 sub samples in duplicate, October 2013)

GC-FID: Instrument: Agilent 6890
 Column: HP-1MS, 30 m × 0.32 mm I.D. × 0.25 μm
 Program: 100 °C (1 min), 10 °C/min to 220 °C, 30 °C/min to 300 °C (3 min)
 Injector: 200 °C
 Detector Temp: 320 °C
 Carrier: Helium
 Split ratio: 20/1
 Relative peak area of the main component as the free base:
 Initial analysis: Mean = 99.5%, s = 0.04% (10 sub samples in duplicate, October 2013)

Karl Fischer analysis: Moisture content 0.1% mass fraction (October 2013 & June 2017)
 Moisture content 0.15% mass fraction (May 2020)

Thermogravimetric analysis: Non-volatile residue < 0.2% mass fraction (October 2013). The volatile content (e.g. organic solvents and/or water) could not be determined because of the inherent volatility of the material and/or degradation at elevated temperatures.

Spectroscopic and other characterisation data

GC-MS:	Instrument:	Agilent 6890/5973
	Column:	TG-1MS, 30 m x 0.25 mm I.D. x 0.25 μ m
	Program:	50 °C (3 min), 5 °C/min to 100 °C, 15 °C/min to 250 °C (5 min), 30 °C/min to 300 °C (2 min)
	Injector:	250 °C
	Split ratio:	20/1
	Transfer line temp:	280 °C
	Carrier:	Helium, 1.0 mL/min
	Scan range:	50-550 <i>m/z</i>
	The retention time of the free base is reported along 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 base peak.	
	Free base (21.9 min):	243 (M^+ , 37), 242 (44), 201 (21), 200 (100), 186 (27), 166 (21), 91 (41) <i>m/z</i>
ESI -MS:	Instrument:	Micromass Quatro LC Micro
	Operation:	Positive ion mode, direct infusion at 10 μ L/min
	Ionisation:	ESI spray voltage at 3.0 kV positive ion
	EM voltage:	650 V
	Cone voltage:	15 V
	Peak:	244.3 ($M+H^+$) <i>m/z</i>
HS-GC-MS:	Instrument:	Agilent 6890/5973/G1888
	Column:	DB-624, 30 m x 0.25 mm I.D. x 1.4 μ m
	Program:	50 °C (5 min), 7 °C/min to 120 °C, 15 °C/min to 220 °C (8.3 min)
	Injector:	150 °C
	Split ratio:	50/1
	Transfer line temp:	280 °C
	Carrier:	Helium, 1.2 mL/min
	Solvents detected:	Ethyl chloride, ethanol, 2-propanol, diethyl ether
TLC:	Conditions:	Kieselgel 60F ₂₅₄ . Hexane/ethyl acetate (3:1) Single spot observed, R_f = 0.24. Visualisation with UV at 254 nm
IR:	Instrument:	Biorad FTS3000MX FT-IR
	Range:	4000-400 cm^{-1} , KBr powder
	Peaks:	2937, 2861, 2480, 1446, 1296, 1193, 1018, 894, 761, 701, 651, 445 cm^{-1}
¹ H NMR:	Instrument:	Bruker Avance III-400
	Field strength:	400 MHz
	Solvent:	MeOH- <i>d</i> ₄ (3.31 ppm)
	Spectral data:	δ 1.12-1.29 (3H, m), 1.35 (2H, qt, J = 13.1, 4.0 Hz), 1.60 (1H, m), 1.72 (1H, m), 1.78-1.92 (6H, m), 1.97 (2H, dt, J = 3.3, 12.9 Hz), 2.35 (2H, m), 3.07 (2H, dm, J = 11.5 Hz), 3.75 (2H, bd, J = 11.5 Hz), 7.49-7.59 (3H, m), 7.66-7.69 (2H, m) ppm Ethanol estimated at 0.4% mass fraction was observed in the ¹ H NMR
¹³ C NMR:	Instrument:	Bruker Avance III-400
	Field strength:	101 MHz
	Solvent:	MeOH- <i>d</i> ₄ (49.0 ppm)
	Spectral data:	δ 23.1, 24.0, 24.6, 26.1, 32.3, 48.9, 72.8, 130.4, 130.96, 131.00, 131.7 ppm
Melting point:		240 °C
Microanalysis:	Found:	C = 72.7%; H = 9.2%; N = 5.1%; Cl = 12.5% (October, 2013)
	Calculated:	C = 73.0%; H = 9.4%; N = 5.0%; Cl = 12.7% (Calculated for C ₁₇ H ₂₅ N.HCl)