Australian Government
Department of Industry, Science and Resources

National Measurement Institute

NATA
WORLD RECOGNISED ACCREDITATION

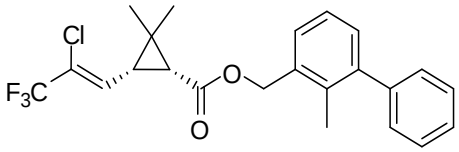
CERTIFIED REFERENCE MATERIAL CERTIFICATE OF ANALYSIS

NMIA P1804: Bifenthrin (cis form)

Report ID: P1804.2025.01

Chemical Formula: $C_{23}H_{22}ClF_3O_2$

Molecular Weight: 422.9 g/mol



Certified value

Batch No.	CAS No.	Purity (mass fraction)
10-AV-01	**82657-04-3**	**99.6 ± 0.3%**

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

IUPAC name: (2-Methyl-3-biphenylyl)methyl (1SR,3SR)-3-[(1Z)-2-chloro-3,3,3-trifluoro-1-propen-1-yl]-2,2-dimethylcyclopropanecarboxylate (1R, 3R is shown).

Note: The absolute stereochemistry and enantiomeric purity have not been determined.

Expiration of certification: The property values are valid till 23 April 2035, 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. The material will be re-tested on an annual basis to ensure that the property values are still valid. In the event a product fails the stability trial, notification will be sent to all impacted customers.

Description: White powder sourced from an external supplier and 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 4 ºC 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 5 years. The measurement uncertainty at the 95% confidence 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 five 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.
16 July 2025

This report supersedes any issued prior to 16 July 2025.

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

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 ^{1}H NMR spectroscopy. The purity value is calculated as per Equation 1.

Purity = (100 % - I_{ORG}) x (100 % - I_{VOL} – I_{NVR}) Equation 1

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

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

Note: Bifenthrin in the *trans* configuration is present in 0.3-0.4% mass fraction.

GC-FID:	Instrument:	Agilent 6890 or 8890		
	Column:	DB-5, HP-5 or VF-1MS, 30 m × 0.32 mm I.D. × 0.25 μm		
	Program:	150 °C (1 min), 20 °C/min to 230°C (8 min), 20 °C/min to 310 °C (2 min)		
	Injector:	250 °C	Detector Temp:	320 °C
	Carrier:	Helium	Split ratio:	20/1
	Relative mass fraction of the main component:			
	Initial analysis:	Mean = 99.5%, s = 0.005% (10 sub samples in duplicate, September 2010)		
	Re-analysis:	Mean = 99.6%, s = 0.02% (10 sub samples in duplicate, September 2010)		
	Re-analysis:	Mean = 99.6%, s = 0.02% (5 sub samples in duplicate, June 2011)		
	Re-analysis:	Mean = 99.6%, s = 0.03% (5 sub samples in duplicate, April 2012)		
	Re-analysis:	Mean = 99.6%, s = 0.01% (5 sub samples in duplicate, January 2016)		
	Re-analysis:	Mean = 99.6%, s = 0.01% (5 sub samples in duplicate, November 2020)		
	Re-analysis:	Mean = 99.6%, s = 0.01% (5 sub samples in duplicate, April 2025)		
Karl Fischer analysis:		Moisture content 0.1% mass fraction (August 2010, June 2011 & May 2012) Moisture content < 0.1% mass fraction (February 2016, November 2020, and April 2025)		
Thermogravimetric analysis:		Non volatile residue < 0.1% mass fraction (September 2010). The volatile content, organic solvents and/or water, could not be analysed accurately because of the inherent volatility of the material.		
QNMR:	Instrument:	Bruker Avance-400		
	Field strength:	400 MHz		
	Solvent:	$CDCl_3$ (7.26 ppm)		
	Internal standard:	Dimethyl sulfone (100% mass fraction)		
	Initial analysis:	Mean = 100.2%, s = 0.24% (4 sub samples, September 2010)		

Spectroscopic and other characterisation data

GC-MS:	Instrument:	Agilent 6890/5973
	Column:	DB-5MS, 30 m x 0.25 mm I.D. x 0.25 μm
	Program:	150 °C (1 min), 25 °C/min to 230 °C (8 min), 25 °C/min to 310 °C (2 min)
	Injector:	250 °C
	Split ratio:	20/1
	Transfer line temp:	280 °C
	Carrier:	Helium
	Scan range:	50-550 *m/z*
	The retention time of the parent compound is reported with the major peaks in the mass spectra. The latter are reported as mass/charge ratios and (in brackets) as a percentage relative to the base peak.	
	Parent (10.80 min):	422 (M^+, < 1), 181 (100), 166 (27), 165 (27), 152 (3), 141 (3) *m/z*
HS-GC-MS:	Instrument:	Agilent 6890/5973/G1888
	Column:	DB-624, 30 m x 0.25 mm I.D. x 1.4 μm
	Program:	50 °C (5 min), 7 °C/min to 120 °C, 15 °C/min to 220 °C (8.3 min)
	Injector:	150 °C
	Transfer line temp:	280 °C
	Carrier:	Helium, 1.2 mL/min
	Split ratio:	50/1
	Solvents detected:	Hexane
TLC:	Conditions:	Kieselgel 60F_{254}. Hexane/acetone (98:2) Single spot observed, R_f = 0.6. Visualisation with UV at 254 nm.
IR:	Instrument:	Biorad FTS3000MX FT-IR
	Range:	4000-400 cm^{-1}, KBr powder
	Peaks:	3093, 3059, 3004, 2970, 1720, 1141, 1358, 1296, 1274, 1197, 1148, 952, 889, 786, 704 cm^{-1}
^{1}H NMR:	Instrument:	Bruker Avance-400
	Field strength:	400 MHz
	Solvent:	$CDCl_3$ (7.26 ppm)
	Spectral data:	δ 1.31 (3H, s) 1.32 (3H, s), 2.07(1H, d, *J* = 8.4 Hz), 2.19 (1H, t, *J* = 9.0 Hz), 2.22 (3H, s), 5.19 (1H, d, *J* = 12.6 Hz), 5.23 (1H, d, *J* = 12.6 Hz), 6.96 (1H, m), 7.23-7.27 (2H, m), 7.28-7.32 (2H, m), 7.34-7.39 (2H, m), 7.40-7.44 (2H, m) ppm
	Hexane observed in the HS-GC-MS was below the limit of detection by ^{1}H NMR.	
^{13}C NMR:	Instrument:	Bruker Avance-400
	Field strength:	100 MHz
	Solvent:	$CDCl_3$ (77.0 ppm)
	Spectral data:	δ 14.9, 16.2, 28.4, 28.7, 30.9, 32.9, 65.4, 120.4 (q, *J* = 272 Hz), 121.8 (q, *J* = 38 Hz), 125.6, 126.9, 128.1, 128.4, 129.3, 130.0 (q, *J* = 4.5 Hz), 130.4, 134.2, 134.4, 141.8, 143.0, 170.1 ppm
Melting point:		69-70 °C
Microanalysis:	Found:	C = 65.4%; H = 5.3%; Cl = 8.8%; F = 13.4% (July, 2010)
	Calculated:	C = 65.3%; H = 5.2%; Cl = 8.4%; F = 13.5% (Calculated for $C_{23}H_{22}ClF_3O_2$)