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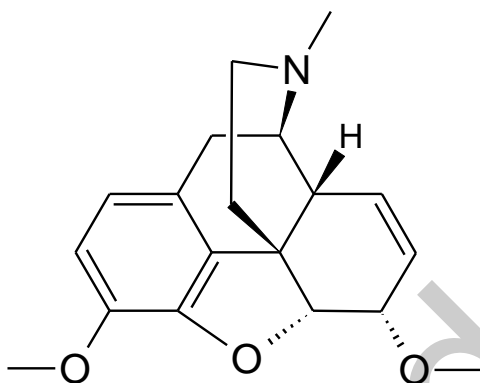
The results of the tests, calibrations and/or measurements included in this document are traceable to Australia/national standards.
NATA is a signatory to the APLAC Mutual Recognition Arrangement for the mutual recognition of the equivalence of reference materials certificates.



Our Formula. Your Success.

Reference Material Product Information Sheet

Epichem's Quality System conforms to ISO9001:2015 as certified by ECAAS Pty Ltd - Certification number 616061.



Name	O-methylcodeine
BP Name	Codeine Impurity A
Epichem Item #	EPL-AA141 Batch 2
CAS #	2859-16-7
Molecular Formula	C ₁₉ H ₂₃ NO ₃
Molecular Weight	313.40 g/mol
Appearance	White powder
Melting Point	141.2-143.0°C
Combustion Analysis	Required (%): C:72.8; H:7.4; N:4.5. Found (%): C:72.9; H:7.4; N:4.4.
Purity*	99.8%
Date of Manufacture	11 September 2018
Storage Requirements	Protect from heat, light and moisture.
Special Precautions	This compound is for laboratory use only. Its toxicological properties may not have been fully established. It should be handled only by suitably qualified personnel.
Intended Use	This compound is suitable for the identification of impurities and degradants in pharmaceutical materials. The purity assay is considered as relative contribution.
Date of Shipment	TBA This certificate is valid for one year from the date of shipment provided the substance is unopened and stored under the recommended conditions.
Retest Date	TBA (Proper Storage and Handling Required)

* NATA accreditation does not cover the performance of this service

EPL-AA141 Batch 2

Epichem Pty Ltd, Suite 5, 3 Brodie-Hall Drive, Bentley WA 6102, Australia
Tel + 61 (0)8 6167 5200 Fax + 61 (0)8 6167 5201 www.epichem.com.au ABN 80 106 769 902

I. Identity

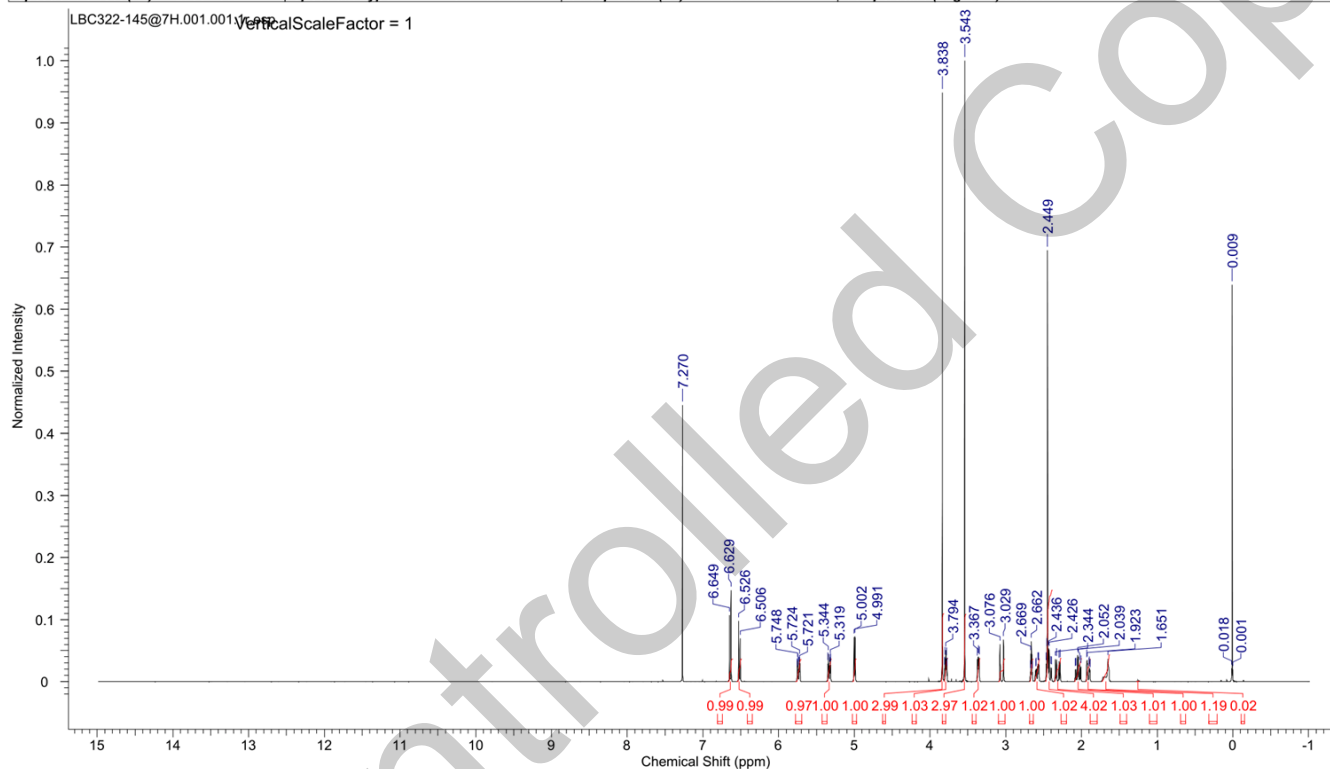
The identity of this product was established using the following analyses:

Ia. ¹H NMR Spectrum

Conditions: 400 MHz, CDCl₃

¹H NMR spectrum consistent with chemical structure.

Acquisition Time (sec)	3.7547	Comment	LBC322-145@7H 1H CDCl3 (E:\data\external\epichem\ cygoh	Date	10 Sep 2018 16:40:32
Date Stamp	10 Sep 2018 16:40:32	File Name	\naphthalene\company\NMR files\LBC322-145@7H\1\data\111r		
Frequency (MHz)	400.13	Nucleus	1H	Number of Transients	8
Original Points Count	24038	Owner	nmr	Points Count	32768
Receiver Gain	144.00	SW(cyclical) (Hz)	6402.05	Solvent	CHLOROFORM-d
Spectrum Offset (Hz)	2795.1392	Spectrum Type	STANDARD	Sweep Width (Hz)	6401.85
				Temperature (degree C)	26.945



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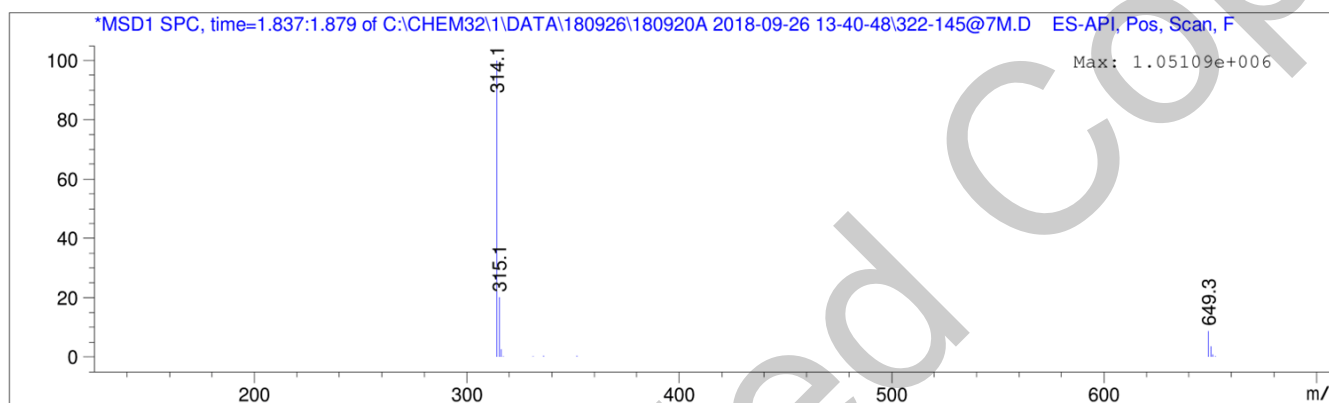
ABN 80 106 769 902

Ib. Mass Spectrum

The mass spectrum of this material was analysed by Liquid Chromatography Mass Spectroscopy (LCMS) using in-house EM005.WI08.

Method: ACN/water gradient (+ 0.1% formic acid).
ZORBAX SB-C8, 4.6 x 30 mm, 3.5 micron.

Retention Time (MS)	MS Area	Mol. Weight or Ion
1.856	8399350	315.15 I
		314.15 I



Theoretical value: 314.2 [M+H]⁺.

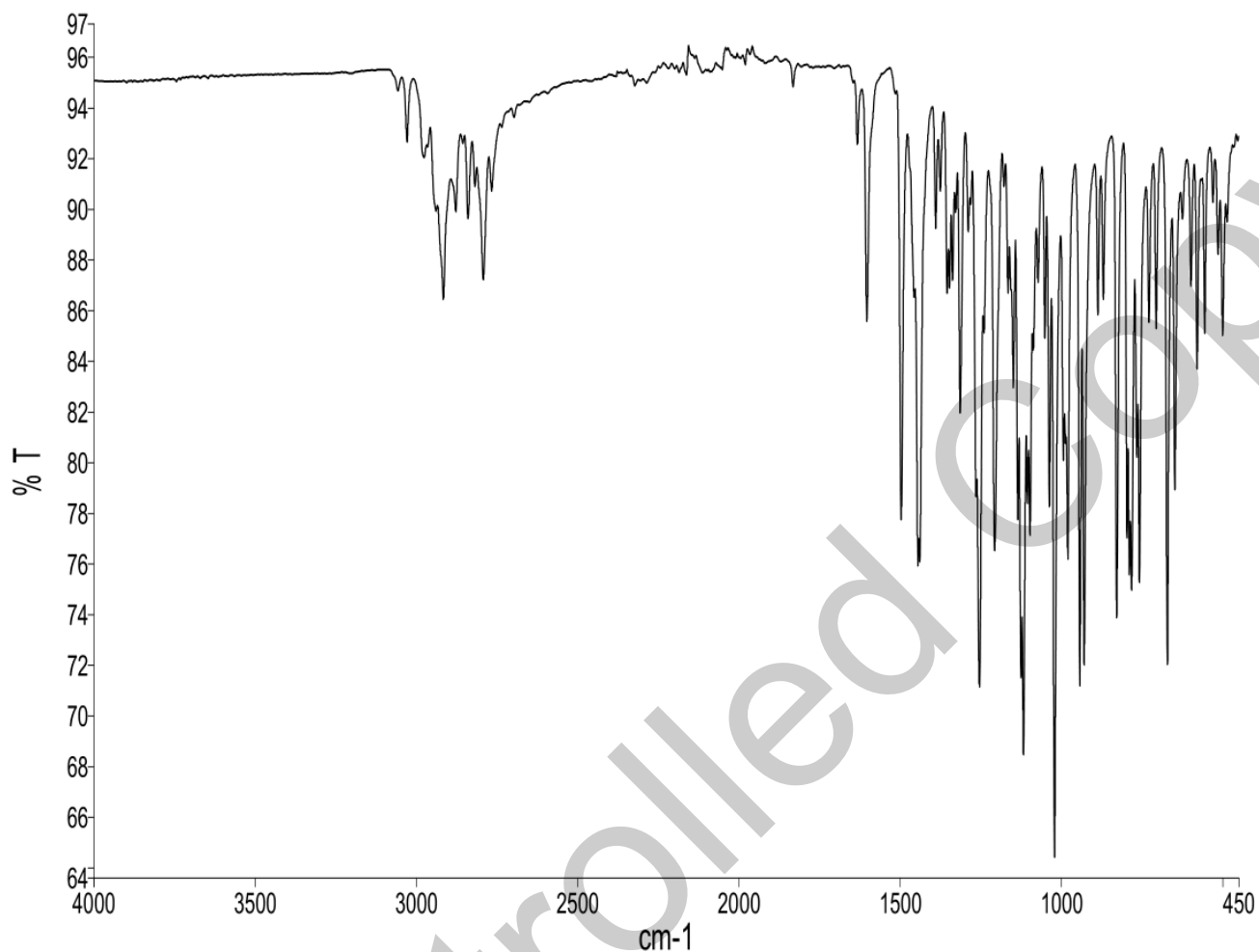
The signal of the Mass Spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

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Ic. IR Spectrum

The infra-red spectrum of this material was analysed by Fourier-Transform Infrared Spectroscopy (FTIR) using in-house EM005.WI09.



The interpretation of the signals of the Fourier-Transform Infrared Spectrum is consistent with the structural formula.

EPL-AA141 Batch 2

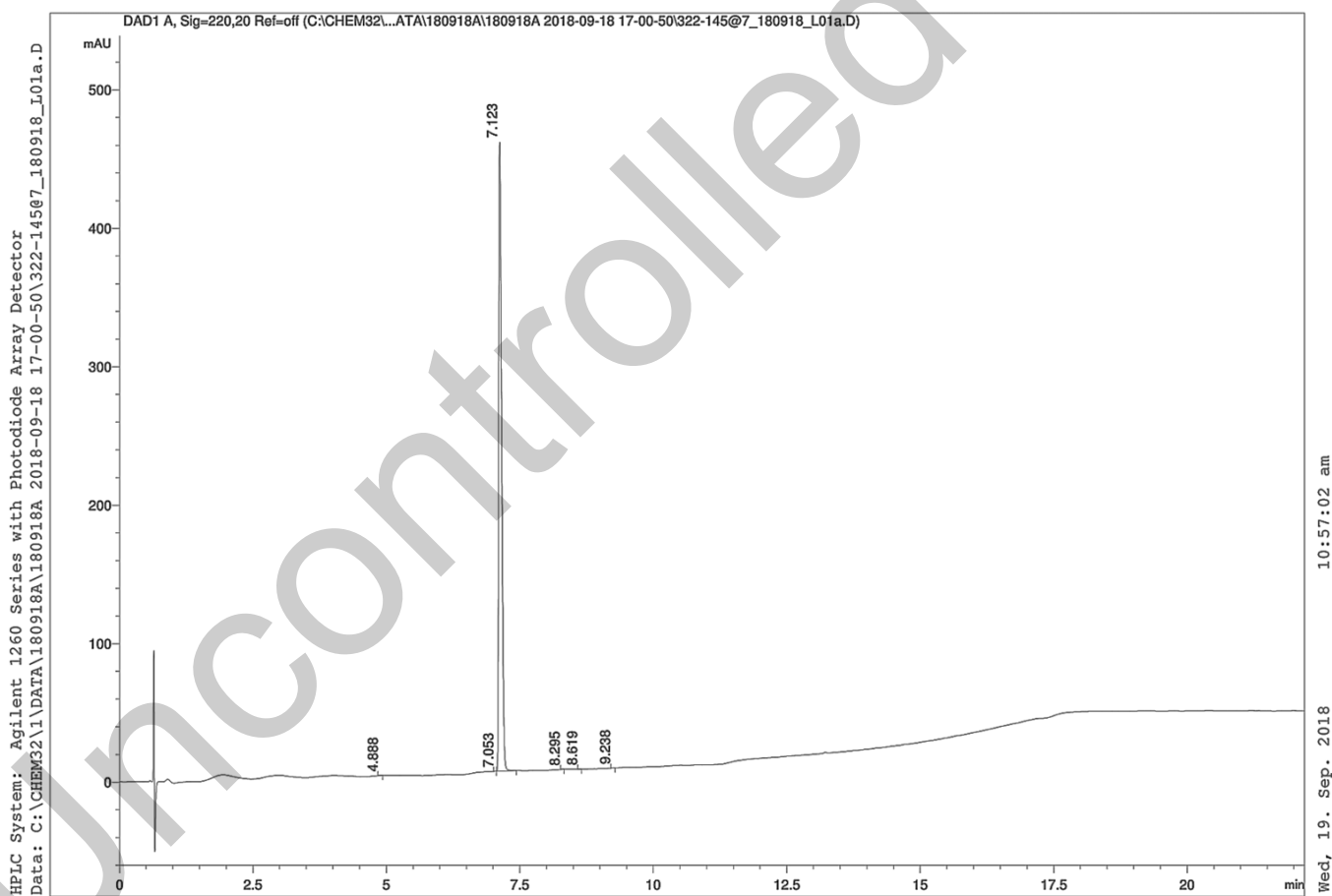
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II. Purity

The purity of this material was analysed by high performance liquid chromatography (HPLC) using in-house EM005.WI07.

HPLC Conditions:

Column	Conditions				Detector	Injector
Agilent Poroshell 120 EC-C18 4.6 x 50mm 2.7 micron	25°C				DAD 220nm	Auto 1.0 µL 0.65 mg/mL 100% acetonitrile
	Time (min)	% Line A (Water + 0.1% (v/v) TFA)	% Line B (Acetonitrile + 0.1% (v/v) TFA)	Flow rate (mL/min)		
	0.00	95	5	1.0		
	5.00	85	15	1.0		
	10.00	65	35	1.0		
	16.00	5	95	1.0		
	21.00	5	95	1.0		
	22.00	95	5	1.0		
	25.00	95	5	1.0		



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Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	4.89	0.52	0.03
2	7.05	0.36	0.02
3	7.12	1696.71	99.88
4	8.29	0.21	0.01
5	8.62	0.32	0.02
6	9.24	0.61	0.04
Totals			100 (rounded)

For the calculation the system peaks were ignored. The content of the analyte was determined as a ratio of the peak area of the analyte and the cumulative areas of the purities, added up to 100%.

Results:

Average 99.8% (average of 10 duplicate analyses)

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III. Water Content

Method: Karl-Fischer titration using in-house EM005.WI04.

Results:

Average <0.1%

IV. Ash Content

Method: BP2018 Ash (Appendix XI J) Method II

Result:

Contains <0.1% ash.

V. Residual Solvents

Method: ¹HNMR

Result:

No significant impurities detected by ¹H NMR analysis.

VI. Final Result

Chromatographic purity (HPLC)	99.8%
Water content	<0.1%
Ash content	<0.1%
Residual solvents	<0.1%
Purity*	99.8%

This purity is assessed to be 99.8%.

Product Reviewed By:

Product Released By:

John Moursounidis, PhD
Head Reference Standards

Boon Tan
Quality Manager

Release Date: 16 October 2018

*NATA accreditation does not cover the performance of this service.

The calculation of the purity follows the formula:

$$\text{Purity(\%)} = \frac{((\text{Chromatographicpurity[HPLC]}) \times (100 - (\text{watercontent} + \text{ashcontent} + \text{volatilecontents})))}{100}$$

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