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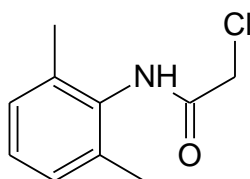
The results of the tests, calibrations and/or measurements included in this document are traceable to Australia/national standards.  
NATA is a signatory to the APLAC Mutual Recognition Arrangement for the mutual recognition of the equivalence of reference materials certificates.



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## Reference Material Product Information Sheet

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



<b>Name</b>	2-chloro- <i>N</i> -(2,6-dimethylphenyl)acetamide
<b>BP Name</b>	Lidocaine Impurity H
<b>Synonym(s)</b>	2',6'-dimethyl-2-chloroacetanilide; <i>N</i> -chloroacetyl-2,6-dimethylaniline
<b>Epichem Item #</b>	EPL-AA98 Batch 2
<b>CAS #</b>	1131-01-7
<b>Molecular Formula</b>	C <sub>10</sub> H <sub>12</sub> ClNO
<b>Molecular Weight</b>	197.67 g/mol
<b>Appearance</b>	White powder
<b>Melting Point</b>	147.2-148.6°C
<b>Combustion Analysis</b>	Required (%): C:60.8, H:6.1, N:7.1. Found (%): C:60.7, H:6.1, N:7.1.
<b>Purity*</b>	100%
<b>Date of Manufacture</b>	6 September 2019
<b>Storage Requirements</b>	Protect from heat, light and moisture.
<b>Special Precautions</b>	<b>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.</b>
<b>Intended Use</b>	This compound is suitable for the identification of impurities and degradants in pharmaceutical materials. The purity assay is considered as relative contribution.
<b>Date of Shipment</b>	TBA This certificate is valid for one year from the date of shipment provided the substance is unopened and stored under the recommended conditions.
<b>Retest Date</b>	TBA (Proper Storage and Handling Required)

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

EPL-AA98 Batch 2

Revision 1

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:

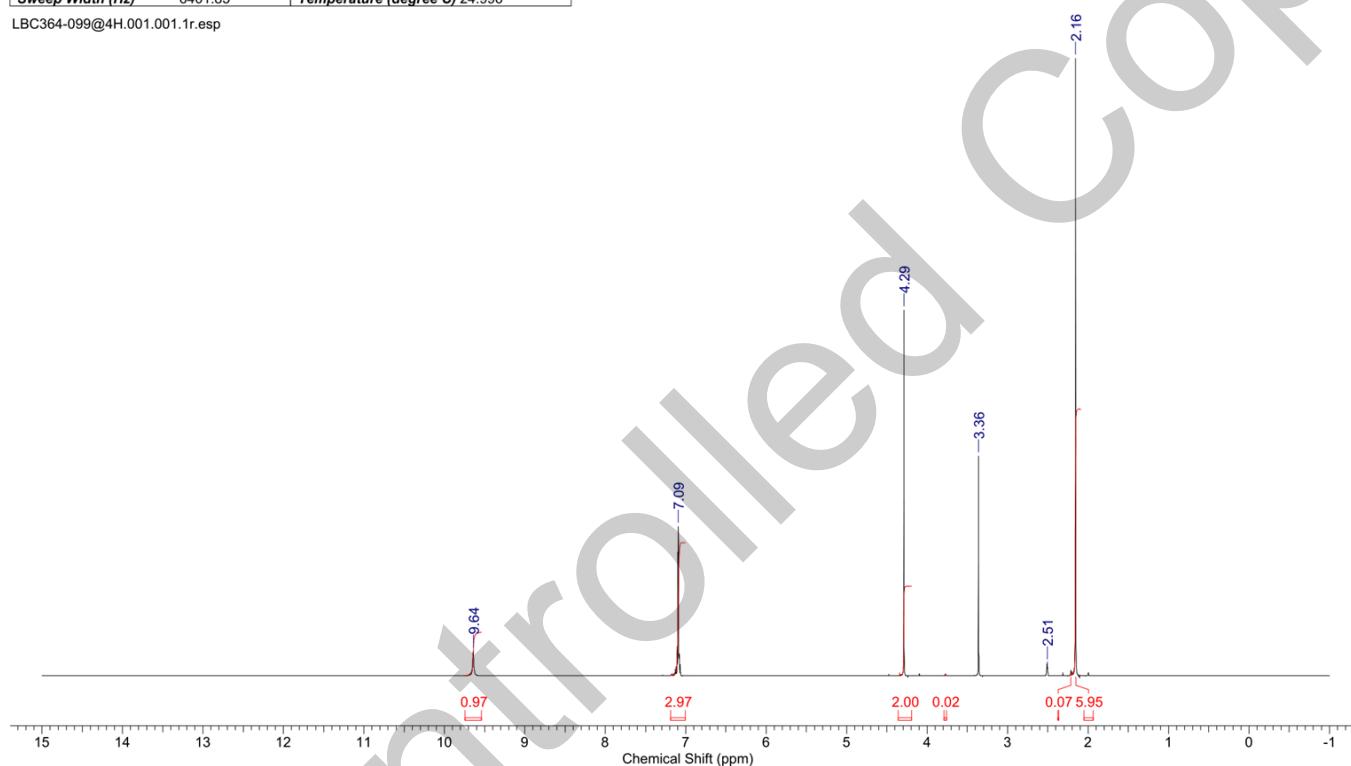
### Ia. <sup>1</sup>H NMR Spectrum

Conditions: 400 MHz, DMSO-d<sub>6</sub>

<sup>1</sup>H NMR spectrum consistent with chemical structure.

Acquisition Time (sec)	3.7547	Comment	LBC364-099@4H 1H DMSO (E:\data\external\epichem) cygoh 16		
Date	24 Jun 2019 17:44:32	Date Stamp	24 Jun 2019 17:44:32		
File Name	\naphthalene\company\NMR files\LBC364\LBC364-099@4H\1\data\1\1r		Frequency (MHz)	400.13	
Nucleus	1H	Number of Transients	8	Origin	spect
Owner	nmr	Points Count	32768	Original Points Count	24038
SW(cyclical) (Hz)	6402.05	Solvent	DMSO-d6	Pulse Sequence	zg
Sweep Width (Hz)	6401.85	Temperature (degree C)	24.996	Receiver Gain	45.20
				Spectrum Offset (Hz)	2800.9097
				Spectrum Type	STANDARD

LBC364-099@4H.001.001.1r.esp



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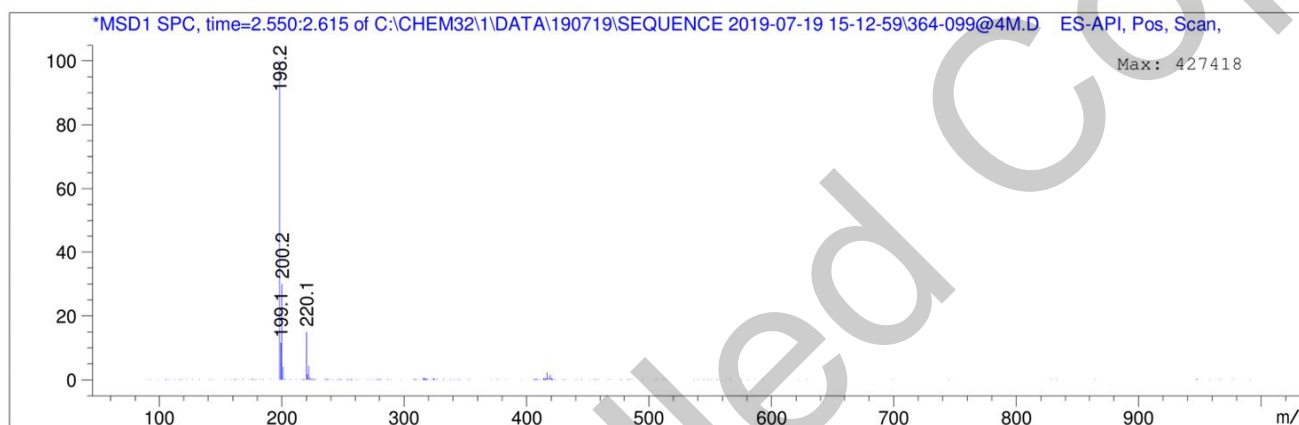
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## Ib. Mass Spectrum

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
2.574	8430058	220.10 I
		200.20 I
		199.10 I
		198.20 I

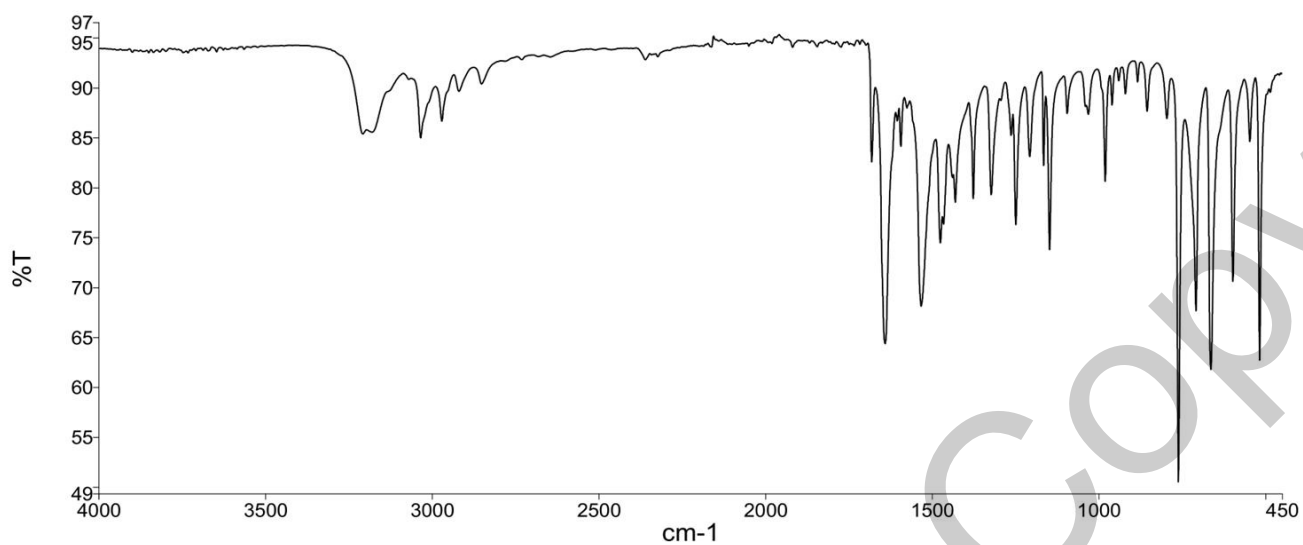


Theoretical value: 198.2 [M+H]<sup>+</sup>.

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

### Ic. IR Spectrum

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



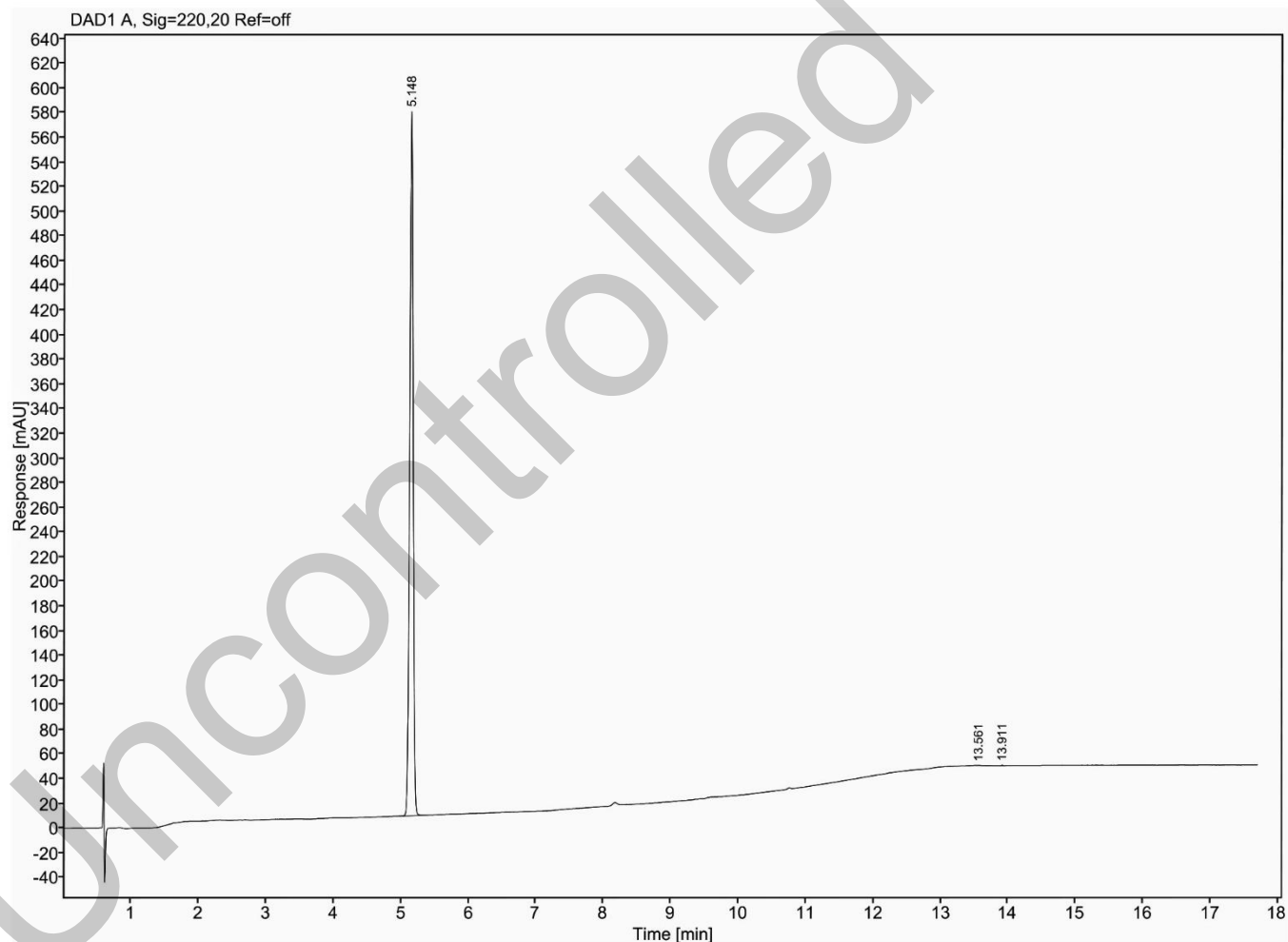
The interpretation of the signals of the Fourier-Transform Infrared Spectrum is consistent with the structural formula.

## 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	45°C				DAD 220nm	Auto 1.0 µL  0.9 mg/mL in 50% acetonitrile 50% water (NO MODIFIERS)
	Time (min)	% Line A (Water + 0.1% (v/v) TFA)	% Line B (Acetonitrile + 0.1% (v/v) TFA)	Flow rate (mL/min)		
	0.00	90	10	1.0		
	6.00	60	40	1.0		
	11.50	5	95	1.0		
	16.50	5	95	1.0		
	17.50	90	10	1.0		
	20.50	90	10	1.0		



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### Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	5.15	2074.19	99.78
2	13.56	4.15	0.20
3	13.91	0.44	0.02
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)

### III. Water Content

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

#### Results:

Average <0.1%

### IV. Ash Content

Method: BP 2019 Ash Appendix XIJ Method II

#### Result:

Contains <0.1% ash.

### V. Residual Solvents

Method: <sup>1</sup>HNMR

#### Result:

<0.1% by <sup>1</sup>H 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:

Jacob Heppell  
Chemist

Carol Worth, PhD  
Quality Manager

Release Date: 28 July 2022

\*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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