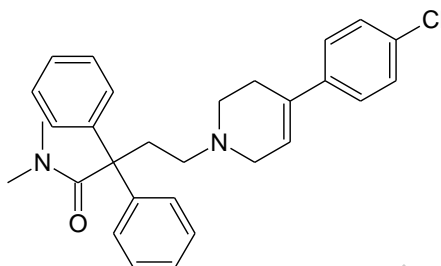


## Reference Material Product Information Sheet

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



<b>Name</b>	dehydro loperamide
<b>BP/EP Name</b>	Loperamide Impurity H
<b>USP Name</b>	Loperamide Impurity H
<b>Synonym(s)</b>	4-(4-Chlorophenyl)-3,6-dihydro-N,N-dimethyl-1,2-diphenylbutanamide, 4-(4-(4-chlorophenyl)-3,6-dihydropyridin-1(2H)-yl)-N,N-dimethyl-2,2-diphenylbutanamide
<b>Epichem Item #</b>	EPL-AA161 Batch 1
<b>CAS #</b>	61299-42-1
<b>Molecular Formula</b>	C <sub>29</sub> H <sub>31</sub> ClN <sub>2</sub> O
<b>Molecular Weight</b>	459.04 g/mol
<b>Appearance</b>	White crystalline powder
<b>Melting Point</b>	134.9-139.2°C.
<b>Combustion Analysis</b>	Required (%): C:75.9; H:6.8; N:6.1. Found (%): C: 75.9; H: 6.9; N: 6.2.
<b>Purity*</b>	99.8%
<b>Date of Manufacture</b>	17 November 2014
<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 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-AA161 Batch 1

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## I. Identity

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

### Ia. <sup>1</sup>HNMR Spectrum

Conditions: 400 MHz, CDCl<sub>3</sub>

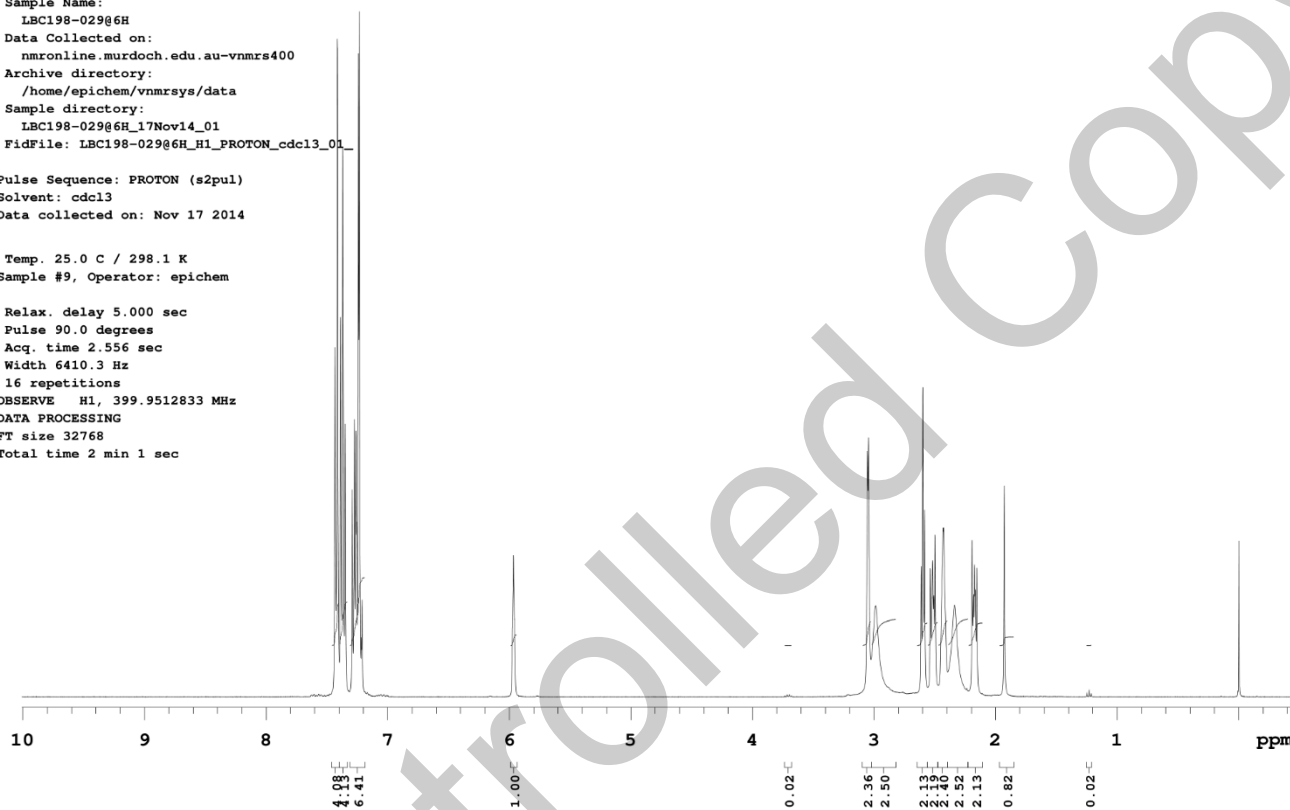
<sup>1</sup>HNMR spectrum consistent with chemical structure.

Sample Name:  
LBC198-029@6H  
Data Collected on:  
nmronline.murdoch.edu.au-vmrs400  
Archive directory:  
/home/epichem/vmrsys/data  
Sample directory:  
LBC198-029@6H\_17Nov14\_01  
FidFile: LBC198-029@6H\_H1\_PROTON\_cdc13\_01

Pulse Sequence: PROTON (s2pul)  
Solvent: cdc13  
Data collected on: Nov 17 2014

Temp. 25.0 C / 298.1 K  
Sample #9, Operator: epichem

Relax. delay 5.000 sec  
Pulse 90.0 degrees  
Acq. time 2.556 sec  
Width 6410.3 Hz  
16 repetitions  
OBSERVE H1, 399.9512833 MHz  
DATA PROCESSING  
FT size 32768  
Total time 2 min 1 sec



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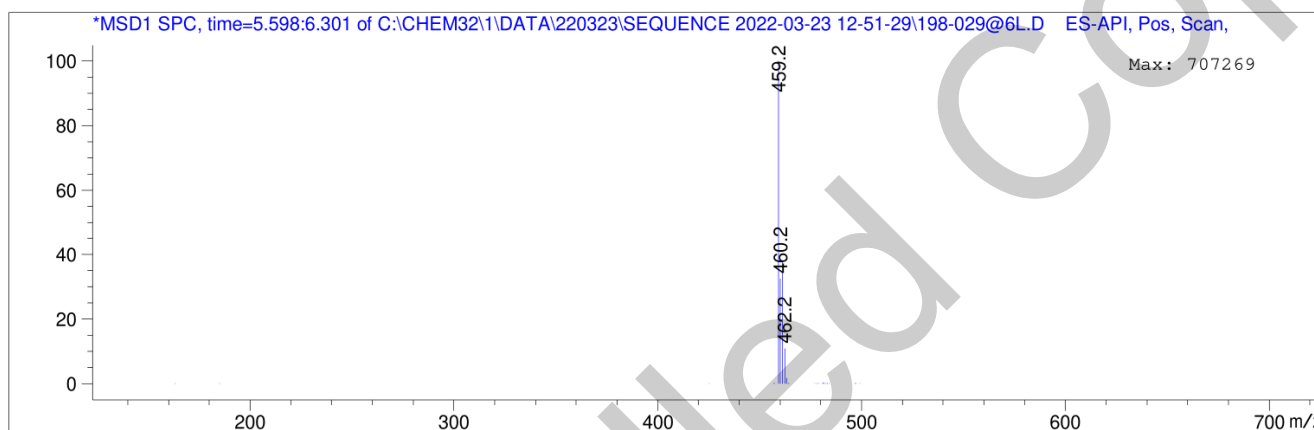
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## Ib. Mass Spectrum

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
5.908	65193672	462.20 I
		461.20 I
		460.20 I
		459.20 I



Theoretical values: 459.2 [M+H]<sup>+</sup>

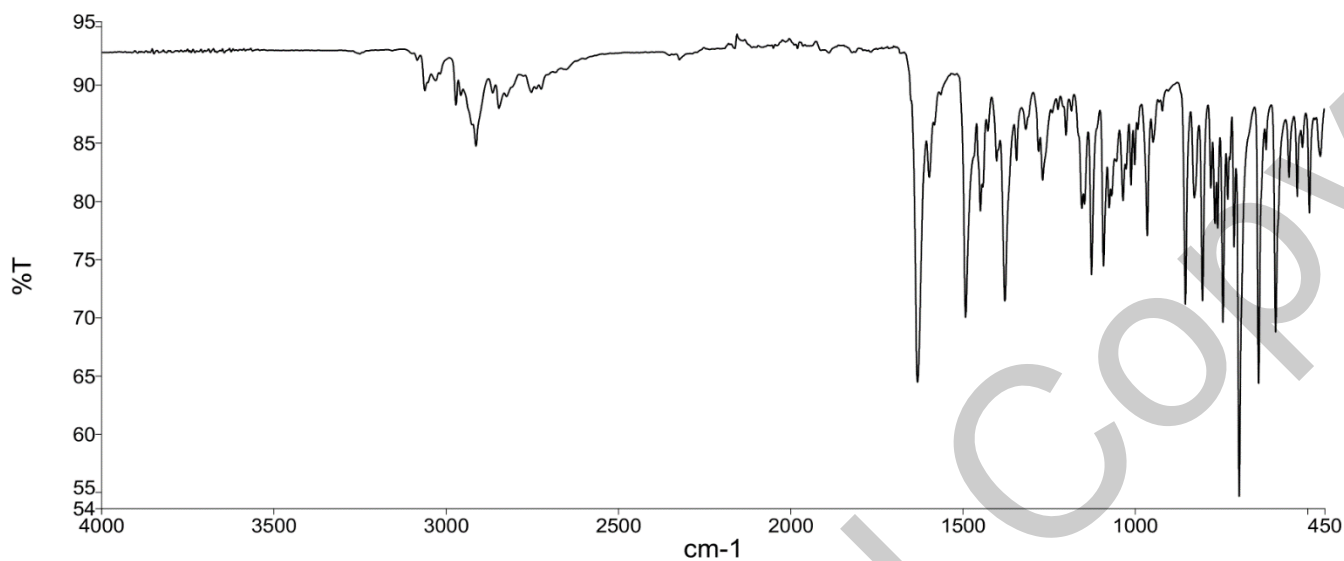
The signal of the Mass Spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

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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.

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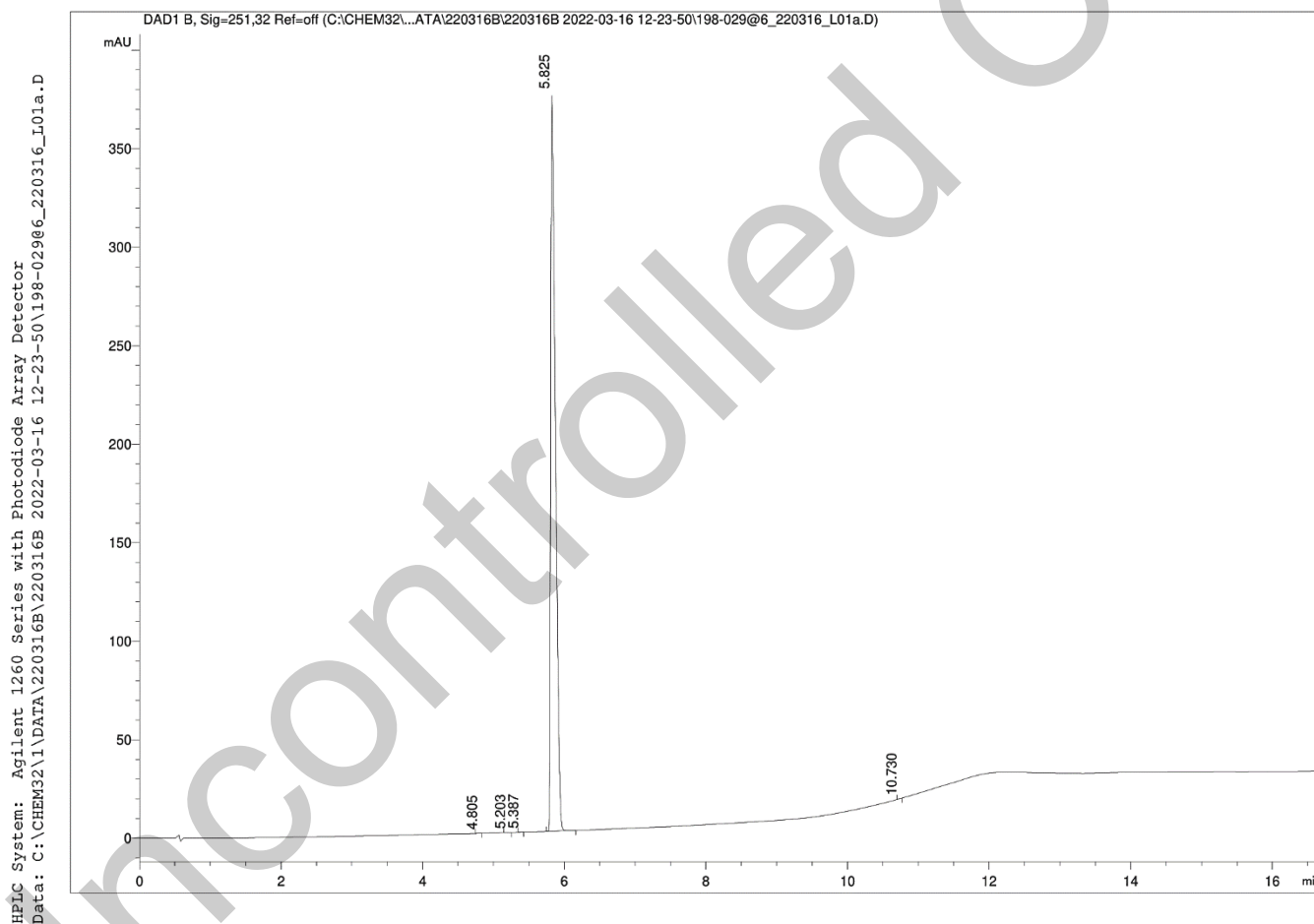
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## II. Purity

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  251nm	Auto  1.0 µL  1.0 mg/mL in 100% acetonitrile (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	70	30	1.0		
	8.00	30	70	1.0		
	10.50	5	95	1.0		
	15.50	5	95	1.0		
	16.50	70	30	1.0		
	19.50	70	30	1.0		



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### Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	4.81	0.10	0.01
2	5.20	0.18	0.01
3	5.39	0.05	0.00
4	5.83	1744.34	99.97
5	10.73	0.18	0.01
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 100.0% (average of 10 duplicate runs)

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### III. Water Content

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

**Results:**

Average 0.1%

### IV. Ash Content

Method: BP 2015 Ash Appendix XI J

**Result:**

Contains <0.1% ash.

### V. Residual Solvents

Method: <sup>1</sup>H NMR

**Result:**

0.1% ethanol detected by <sup>1</sup>H NMR analysis.

### VI. Final Result

Chromatographic purity (HPLC)	100.0%
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:

Jacob Heppell, PhD  
Chemist

Product Released By:

Carol Worth, PhD  
Quality Manager  
Release Date: 22 March 2022

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

The calculation of the purity follows the formula:

$$\text{Purity(\%)} = \frac{((\text{Chromatographic purity [HPLC]}) \times (100 - (\text{water content} + \text{ash content} + \text{volatile contents})))}{100}$$

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