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Accreditation Number: 20126

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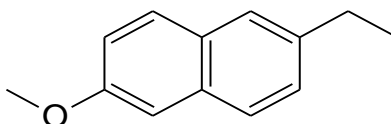
NATA is a signatory to the APLAC Mutual Recognition Arrangement for the mutual recognition of the equivalence of reference materials certificates.



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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.



Name	2-ethyl-6-methoxynaphthalene
BP Name	Naproxen Impurity J
Synonym(s)	Ethylnerolin
Epichem Item #	EPL-AA178 Batch 1
CAS #	21388-17-0
Molecular Formula	C ₁₃ H ₁₄ O
Molecular Weight	186.25 g/mol
Appearance	White powder
Melting Point	60.7-62.0°C
Combustion Analysis	Required (%): C:83.8; H:7.6; N:0.00. Found (%): C:83.7; H:7.6; N:0.00.
Purity*	99.1%
Date of Manufacture	19 November 2015
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 stored under the recommended conditions.
Retest Date	TBA (Proper Storage and Handling Required)

* NATA accreditation does not cover the performance of this service

EPL-AA178 Batch 1

Revision 1

Epichem Pty Ltd, Suite 5, 3 Brodie-Hall Drive, Bentley WA 6102, Australia

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www.epichem.com.au

ABN 80 106 769 902

I. Identity

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

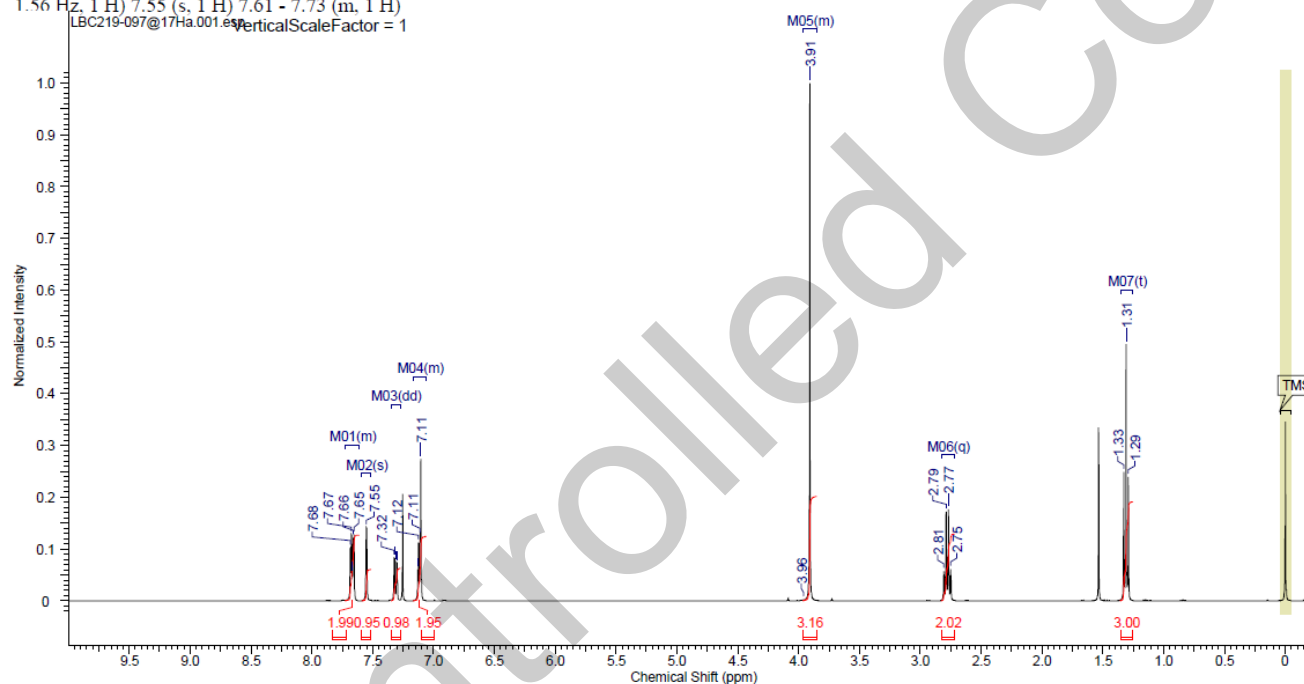
Ia. ¹H NMR Spectrum

Conditions: 400 MHz, CDCl₃

¹H NMR spectrum consistent with chemical structure.

Acquisition Time (sec)	3.7547	Comment	LBC219-097@17Ha 1H CDCl3 (E:\data\external\epichem) cygh 14		
Date	18 Nov 2015 17:55:12	Date Stamp	18 Nov 2015 17:55:12		
File Name	\NAPHTHALENE\Company\NMR files\LBC219-097@17Ha\1.fid			Frequency (MHz)	400.13
Nucleus	1H	Number of Transients	8	Origin	spect
Owner	nmr	Points Count	32768	Pulse Sequence	zg
SW(cyclical) (Hz)	6402.05	Solvent	CHLOROFORM-d	Receiver Gain	144.00
Spectrum Type	STANDARD	Sweep Width (Hz)	6401.85	Spectrum Offset (Hz)	2788.5776
		Temperature (degree C)	26.836		

¹H NMR (400 MHz, CHLOROFORM-d) δ ppm 1.31 (t, *J*=7.62 Hz, 3 H) 2.78 (q, *J*=7.55 Hz, 2 H) 3.85 - 3.96 (m, 3 H) 7.06 - 7.17 (m, 2 H) 7.31 (dd, *J*=8.40, 1.56 Hz, 1 H) 7.55 (s, 1 H) 7.61 - 7.73 (m, 1 H)



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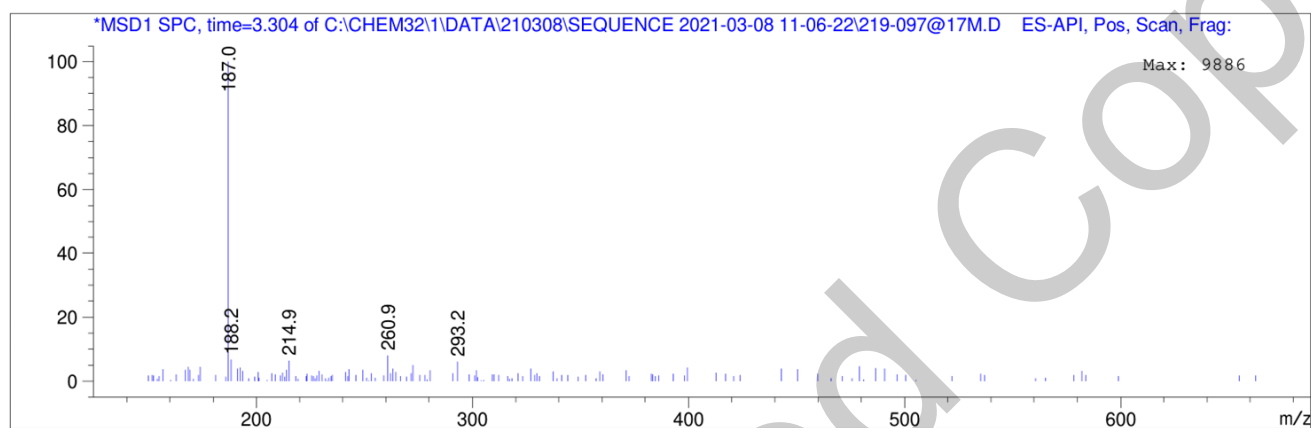
ABN 80 106 769 902

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
3.304	52516	187.00 I



Theoretical value: 187.0 [M+H]⁺.

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

EPL-AA178 Batch 1

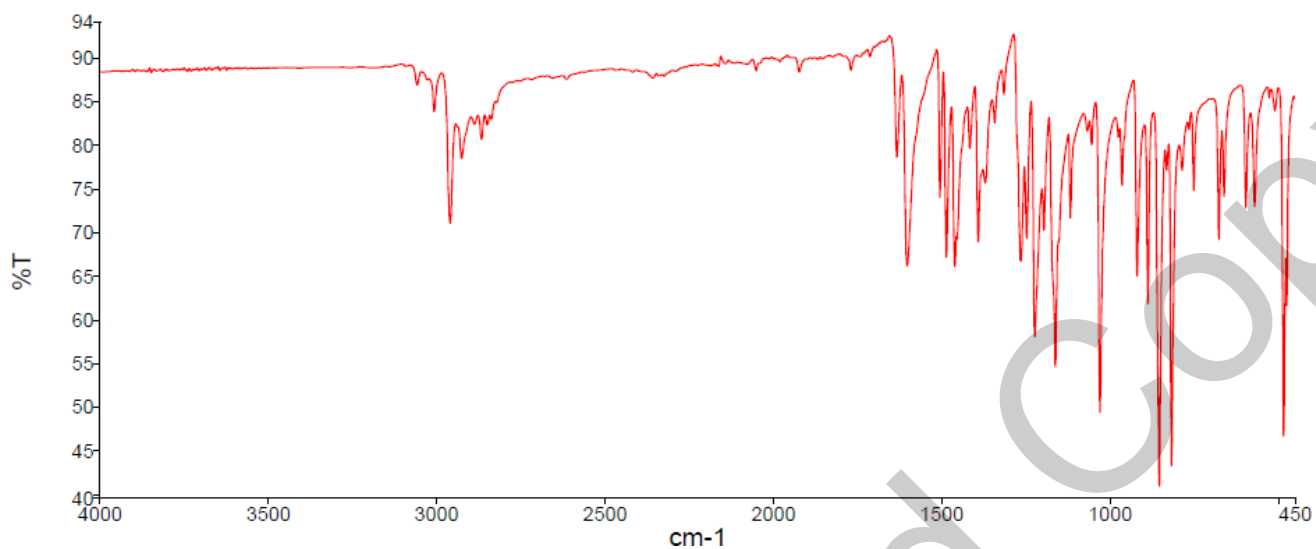
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Ic. IR Spectrum

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

Method: Fourier Transform Infrared (FTIR) Spectroscopy



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

EPL-AA178 Batch 1

Revision 1

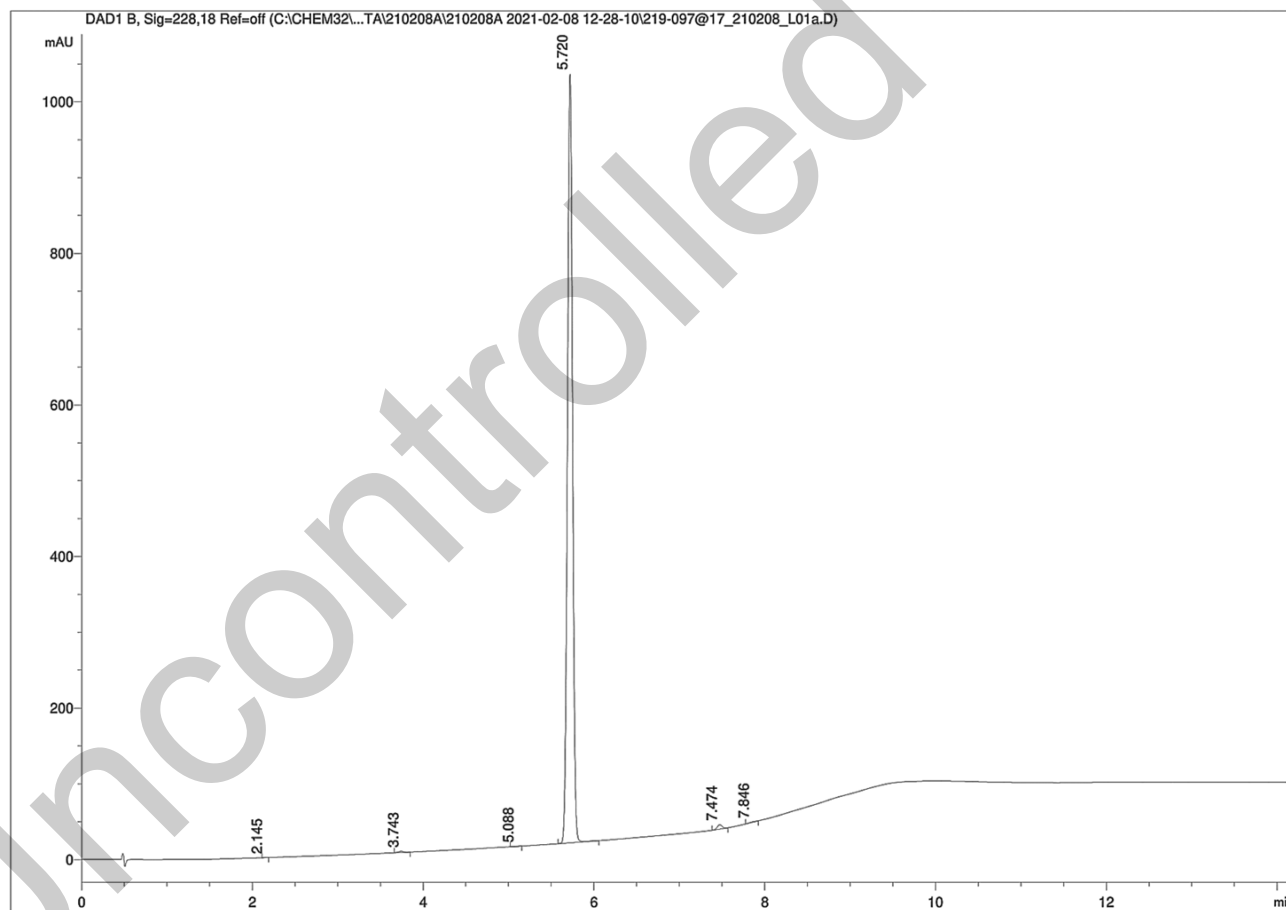
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II. Purity

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 228nm	Auto 1.0 µL 0.20 mg/mL in 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	55	45	1.0		
	6.00	25	75	1.0		
	8.00	5	95	1.0		
	13.00	5	95	1.0		
	14.00	55	45	1.0		
	17.00	55	45	1.0		



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Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	2.14	0.13	0.00
2	3.74	6.06	0.14
3	5.09	0.38	0.01
4	5.72	4183.71	99.26
5	7.47	22.41	0.53
6	7.85	2.02	0.05
Totals			100.0 (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 99.3% (average of duplicate runs)

III. Water Content

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

Results:

Average 0.2%

IV. Ash Content

Method: BP2015 Ash (Appendix XI-J) WS001/26900

Result:

Contains <0.1% ash.

V. Residual Solvents

Method: ¹HNMR

Result:

Contains: <0.1%

VI. Final Result

Chromatographic purity (HPLC)	99.3%
Water content	0.2%
Ash content	<0.1%
Residual solvents	<0.1%
Purity*	99.1%

This purity is assessed to be 99.1%.

Product Reviewed By:

Product Released By:

James Rixson, PhD
Head of Production

Boon Tan
Quality Manager

Release Date: 8 March 2021

**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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