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epichem

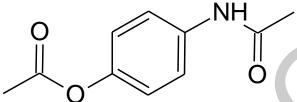
The results of the tests, calibrations and/or measurements included in this document are traceable to Australia/national standards.

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.



	/ 0/ /		
Name	4-(acetylamino)phenyl acetate		
BP Name	Paracetamol Impurity H		
Synonym(s)	N-(4-(acetyloxy)phenyl)acetamide; 4-acetamidophenyl acetate; 4'-acetoxyacetanilide; acetaminophen acetate; 4-acetoxyacetanilide		
Epichem Item #	EPL-AA117 Batch 1		
CAS#	2623-33-8		
Molecular Formula	$C_{10}H_{11}NO_3$		
Molecular Weight	193.20 g/mol		
Appearance	White powder		
<b>Melting Point</b>	153.7-155.2°C		
<b>Combustion Analysis</b>	Required (%): C:62.2; H:5.7; N:7.2. Found (%): C:62.3; H:5.8; N:7.1.		
Purity*	99.6%		
Date of Manufacture	23 May 2012		
<b>Storage Requirements</b>	Protect from heat, light and moisture.		
<b>Special Precautions</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.		
Intended Use	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.		
Retest Date	TBA (Proper Storage and Handling Required)		

<sup>\*</sup> NATA accreditation does not cover the performance of this service

EPL-AA117 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, DMSO-d<sub>6</sub>

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

9/06/2017 2:16:03 PM Acquisition Time (sec) 2.5559 Date Jul 19 2012 Date Stamp Jul 19 2012 File Name Frequency (MHz) Original Points Count 16384 16 Points Count Spectrum Offset (Hz) 16384 2407.3186 s2pul STANDARD Receiver Gain Sweep Width (Hz) 44.00 6410.26 Solvent DMSO-d6
Temperature (degree C) 25.000 Pulse Sequence Spectrum Type -2.51 3.5 8.5 4.5 1.0 10.5 10.0 9.0 2.5 2.0 1.5 4.0 3.0 Chemical Shift (ppm)

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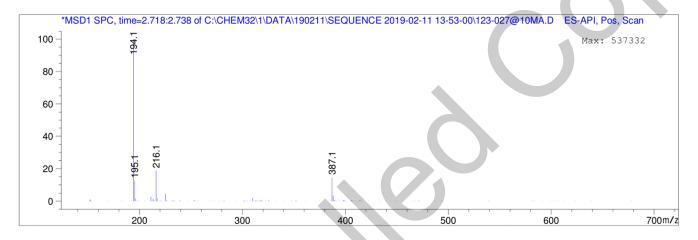
### **Ib.** Mass Spectrum

The mass spectrum of this material was analysed by Liquid Chromatography Mass Spectroscopy (LCMS) using inhouse 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
2.726	4070645	387.10 I 216.10 I 195.10 I 194.15 I



Theoretical value: 194.15 [M+H]+.

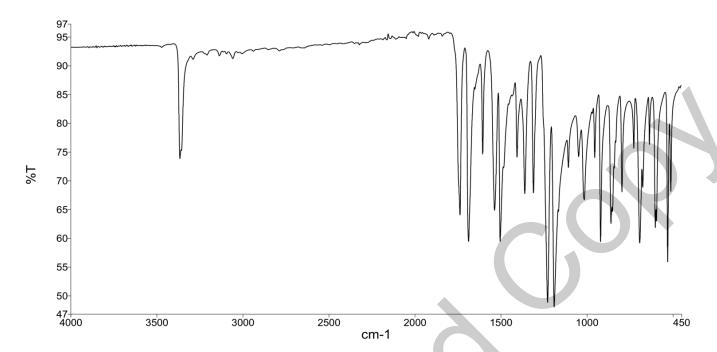
The signal of the Mass Spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

EPL-AA117 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 inhouse EM005.WI09.



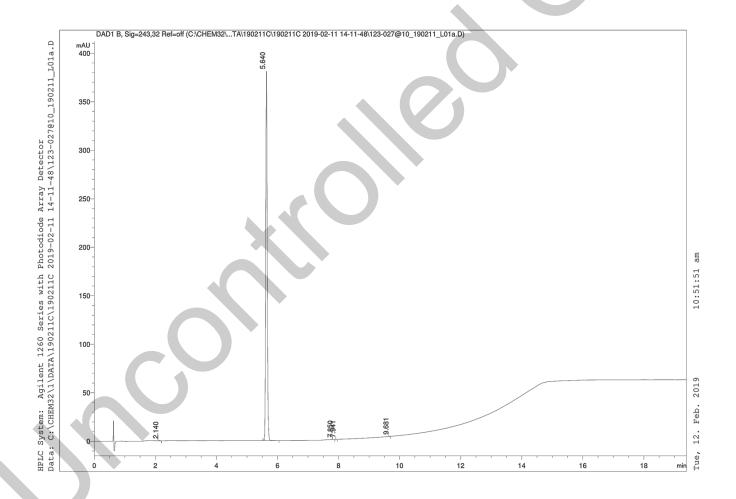
The interpretation of the signals of the Fourier-Transform Infrared Spectrum is consistent with the structural formula.

# 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	25°C DAD Auto					
120 EC-C18 4.6 x 50mm	Time (min)	% Line A (Water + 0.1% (v/v) TFA)	% Line B (Acetonitrile + 0.1% (v/v) TFA)	Flow rate (mL/min)	243nm	1.0 µL 0.4 mg/mL 100% acetonitrile
	0.00	95	5	1.0		accionine
2.7 micron	6.00	77	23	1.0		
	13.20	5	95	1.0		
	18.20	5	95	1.0		
	19.20	95	5	1.0		
	22.20	95	5	1.0		



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### Area Percent Report - Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	2.14	0.87	0.07
2	5.64	1291.30	99.91
3	7.85	0.11	0.01
4	7.94	0.05	0.00
5	9.68	0.10	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 99.9% (average of 10 duplicate runs)



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### **III. Water Content**

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

**Result:** 

Contains 0.1% water.

#### IV. Ash Content

Method: BP2012 Ash

**Result:** 

Contains <0.1% ash.

#### V. Residual Solvents

Method: <sup>1</sup>HNMR

**Result:** 

Contains 0.2% ethanol by <sup>1</sup>HNMR analysis

### VI. Final Result

Chromatographic purity (HPLC)	99.9%
Water content	0.1%
Ash content	<0.1%
Residual solvents	0.2%
Purity*	99.6%

This purity is assessed to be 99.6%.

Product Reviewed By: Product Released By:

John Moursounidis, PhD Boon Tan Head Reference Standards Quality Manager

Release Date: 12 February 2019

The calculation of the purity follows the formula:

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

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