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	eference Material Product Information Sheet tem conforms to ISO9001:2015 as certified by ECAAS Pty Ltd - Certification number 616061.				
HO NO2					
Name	4-nitrophenol				
BP Name	Paracetamol Impurity F				
Synonym(s)	4-hydroxynitrobenzene; <i>p</i> -nitrophenol				
Epichem Item #	EPL-AA115 Batch 1				
CAS #	100-02-7				
Molecular Formula	C <sub>6</sub> H <sub>5</sub> NO <sub>3</sub>				
Molecular Weight	139.11 g/mol				
Appearance	Pale yellow powder				
Melting Point	111.3-114.2°C				
<b>Combustion Analysis</b>	Required (%): C:51.8; H:3.6; N:10.1. Found (%): C:52.0; H:3.8; N:9.8.				
Purity*	99.6%				
Date of Manufacture	26 September 2011				
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	ТВА				
	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-AA115 Batch 1

Revision 1

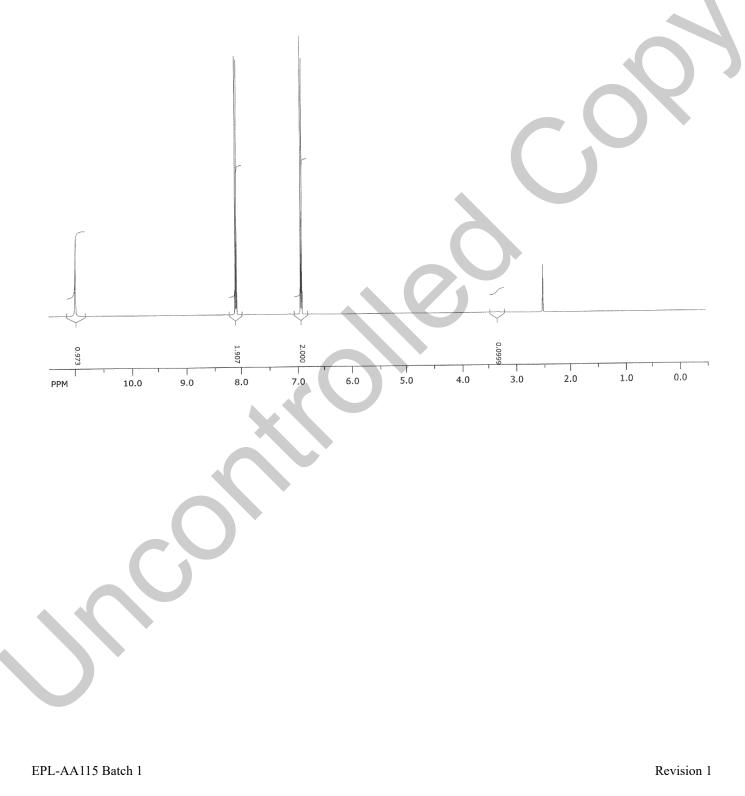
# I. Identity

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.



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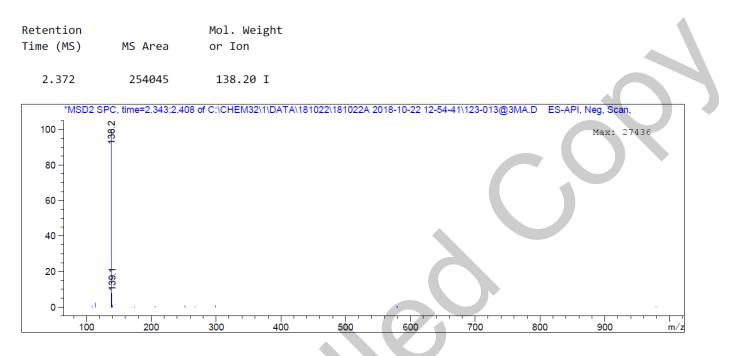
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 ABN 80 106 769 902

#### **Ib. Mass Spectrum**

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

Method: ACN/water gradient (+ 0.1% formic acid).

ZORBAX SB-C8, 4.6 x 30 mm, 3.5 micron.



Theoretical value: 138.2 [M-H]<sup>-</sup>.

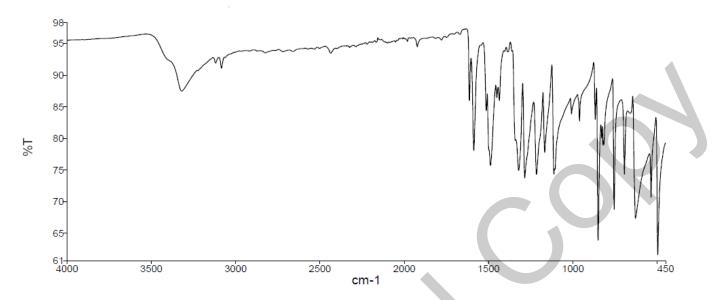
The signal of the Mass Spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

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### Ic. IR Spectrum

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



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

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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	Conditi	ons	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)	318nm	1.0 μL 0.25 mg/mL in
	0.00	90	10	1.0		100% acetonitrile
2.7 micron	5.00	70	30	1.0		(NO MODIFIERS)
	11.50	5	95	1.0	-	
	16.50	5	95	1.0		
	17.50	90	10	1.0		
	20.50	90	10	1.0		
	64 Ref=off (C:\0	CHEM32\ATA\210113A\210113A	2021-01-13 12-39-33\123-013@3_21011	3_L01a.D)		
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50-	C					
50-	C		6	ŝ	226	
50	C		7.479	640 640	12.556	
50-	C		7.479		12.556	

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### Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	5.34	661.85	99.98
2	7.48	0.01	0.00
3	12.01	0.05	0.01
4	12.56	0.05	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% (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: BP 2012 Ash

### **Result:**

Contains 0.3% ash.

## **V. Residual Solvents**

Method: <sup>1</sup>HNMR

#### **Result:**

No significant impurities detected by <sup>1</sup>H NMR analysis.

# VI. Final Result

Chromatographic purity (HPLC)	100%
Water content	0.1%
Ash content	0.3%
Residual solvents	<0.1%
Purity*	99.6%

This purity is assessed to be 99.6%.

Product Reviewed By:

Product Released By:

James Rixson, PhD Head of Production

Carol Worth, PhD Quality Manager Release Date: 24 June 2022

\**NATA accreditation does not cover the performance of this service.* The calculation of the purity follows the formula:

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

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