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	eference Material Product Information Sheet stem conforms to ISO9001:2015 as certified by ECAAS Pty Ltd - Certification number 616061.			
Name	(RS)-2-(4-((4-chlorophenyl)phenylmethyl)piperazin-1-yl)ethan-1-ol dihydrochloride			
BP Name	Cetirizine Impurity G			
Synonym(s)	( <i>RS</i> )-1-((4-chlorophenyl)(phenyl)methyl)-4-(2-hydroxy-ethyl)piperazine dihydrochloride; ( <i>RS</i> )-4-((4-chlorophenyl)phenylmethyl)-1-piperazineethanol dihydrochloride			
Epichem Item #	EPL-AA28 Batch 2			
CAS #	164726-80-1			
Molecular Formula	C <sub>19</sub> H <sub>23</sub> ClN <sub>2</sub> O.2HCl			
Molecular Weight	403.78 g/mol			
Appearance	Fluffy white solid			
Melting Point	208.2-212.2°C (decomposition)			
Combustion Analysis	Required (%): C:56.5; H:6.2; N:6.9; Cl:26.3. Found (%): C:55.4; H:6.5; N:6.6; Cl:26.4.			
Purity*	97.9%			
Date of Manufacture	23 September 2008			
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 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

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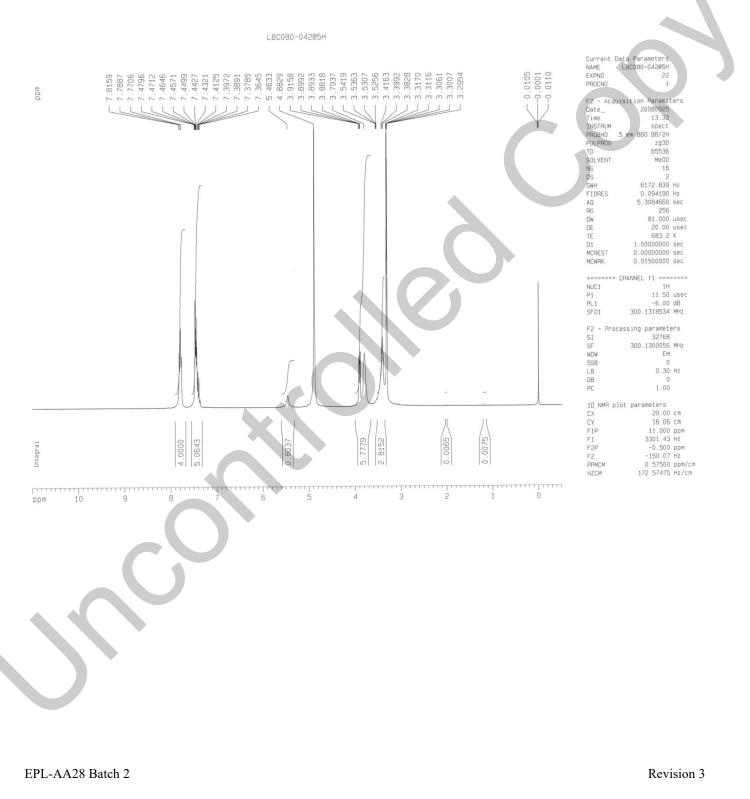
# I. Identity

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

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

Conditions: 300 MHz, CD<sub>3</sub>OD

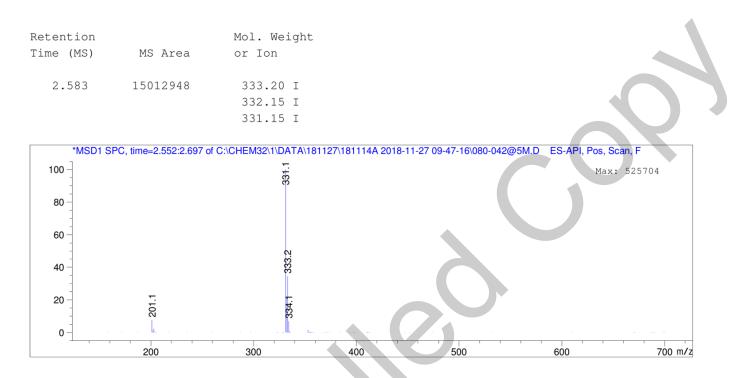
<sup>1</sup>HNMR spectrum consistent with chemical structure.



### **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: 331.1 [M+H]<sup>+</sup>.

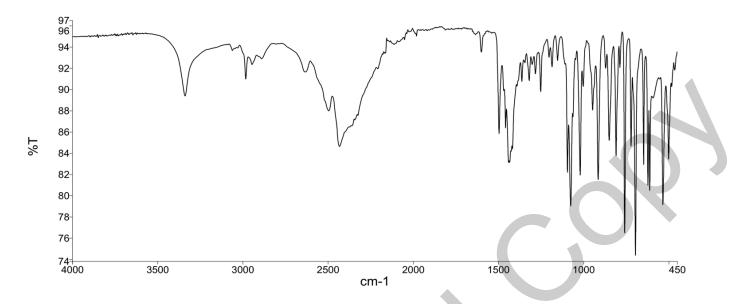
The signal of the Mass Spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

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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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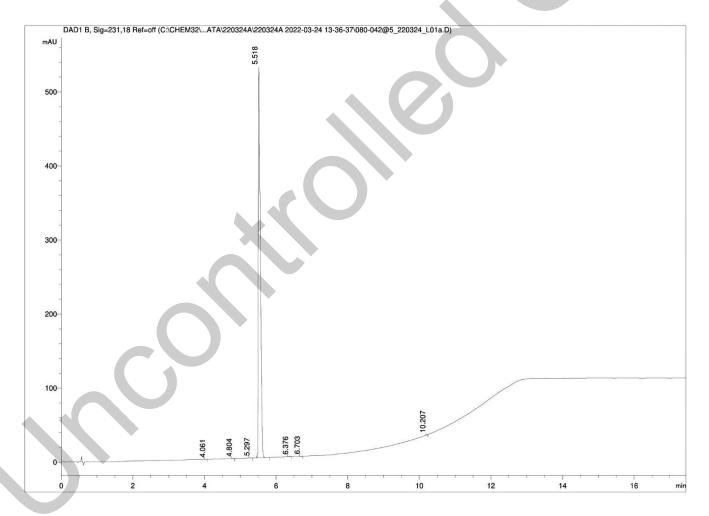
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# **II.** Purity

The purity of this material was analysed by high performance liquid chromatography (HPLC) using in-house EM005.W107.

# **HPLC Conditions:**

Column	Conditi	Conditions				Injector
Agilent Poroshell 120 EC-C18	25°C				DAD	Auto
	Time	% Line A (Water +	% Line B (Acetonitrile	Flow rate	231nm	1.0 μL
4.6 x 50mm 2.7 micron	(min)	0.1% (v/v) TFA)	+ 0.1% (v/v) TFA)	(mL/min)	-	1.25 mg/mL in
	0.00	80	20	1.0		50% acetonitrile
	6.25	55	45	1.0		50% water (+0.1% TFA)
	11.25	5	95	1.0		
	16.25	5	95	1.0		
	17.25	80	20	1.0		
	20.25	80	20	1.0		



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## Area Percent Report – Sorted by Signal

Retention Time (rounded)	Area	Area % (rounded)
4.06	0.10	0.00
4.80	0.24	0.01
5.30	0.40	0.02
5.52	2101.19	99.90
6.38	0.76	0.04
6.70	0.20	0.01
10.21	0.47	0.02
		100 (rounded)
	4.06   4.80   5.30   5.52   6.38   6.70	4.06     0.10       4.80     0.24       5.30     0.40       5.52     2101.19       6.38     0.76       6.70     0.20

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 analyses)

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# **III. Water Content**

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

### **Results:**

Average 1.5%

# IV. Ash Content

Method: Combustion adjuvant added.

### **Result:**

Contains 0.5% ash.

# **V. Residual Solvents**

Method: <sup>1</sup>HNMR

## **Result:**

Contains <0.1% acetonitrile and <0.1% ethanol by <sup>1</sup>H NMR analysis.

# VI. Final Result

Chromatographic purity (HPLC)	99.9%
Water content	1.5%
Ash content	0.5%
Residual solvents	<0.1%
Purity*	97.9%

This purity is assessed to be 97.9%.

Product Reviewed By:

Product Released By:

James Rixson, PhD Head of Production Carol Worth Quality Manager Release Date: 14 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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