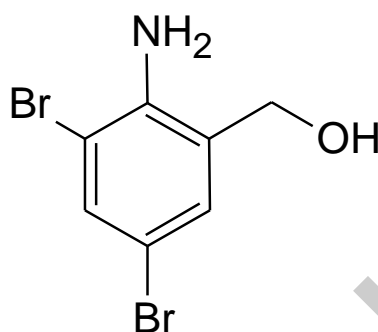


## Reference Material Product Information Sheet

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



<b>Name</b>	(2-amino-3,5-dibromophenyl)methanol
<b>BP Name</b>	Ambroxol Impurity A
<b>Synonym(s)</b>	2-amino-3,5-dibromobenzyl alcohol
<b>Epichem Item #</b>	EPL-AA70 Batch 1
<b>CAS #</b>	50739-76-9
<b>Molecular Formula</b>	C <sub>7</sub> H <sub>7</sub> Br <sub>2</sub> NO
<b>Molecular Weight</b>	280.95 g/mol
<b>Appearance</b>	Cream powder
<b>Melting Point</b>	148.9-149.9°C (decomposition)
<b>Combustion Analysis</b>	Required (%): C:29.9; H:2.5; N:5.0. Found (%): C:30.1; H:2.4; N:4.8.
<b>Purity*</b>	99.0%
<b>Date of Manufacture</b>	8 September 2010
<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-AA70 Batch 1

Revision 4

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

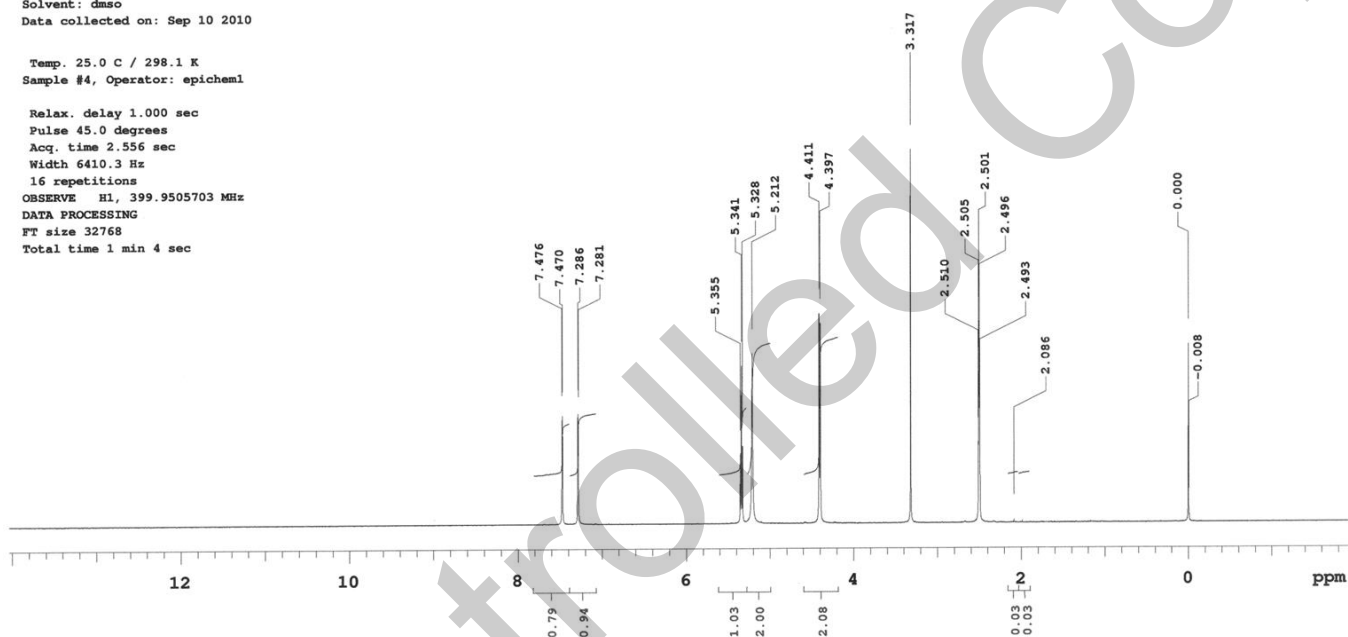
<sup>1</sup>HNMR spectrum consistent with chemical structure.

Sample Name:  
LBC104-175@7H  
Data Collected on:  
VnmrJ-vnmrs400  
Archive directory:  
/home/service/data/epichem1  
Sample directory:  
LBC104-175@7H\_10Sep10\_01  
FidFile: 10Sep10\_LBC104-175@7H\_HI\_PROTON\_dms0\_01

Pulse Sequence: PROTON (s2pul)  
Solvent: dms0  
Data collected on: Sep 10 2010

Temp. 25.0 C / 298.1 K  
Sample #4, Operator: epichem1

Relax. delay 1.000 sec  
Pulse 45.0 degrees  
Acq. time 2.556 sec  
Width 6410.3 Hz  
16 repetitions  
OBSERVE H1, 399.9505703 MHz  
DATA PROCESSING  
FT size 32768  
Total time 1 min 4 sec



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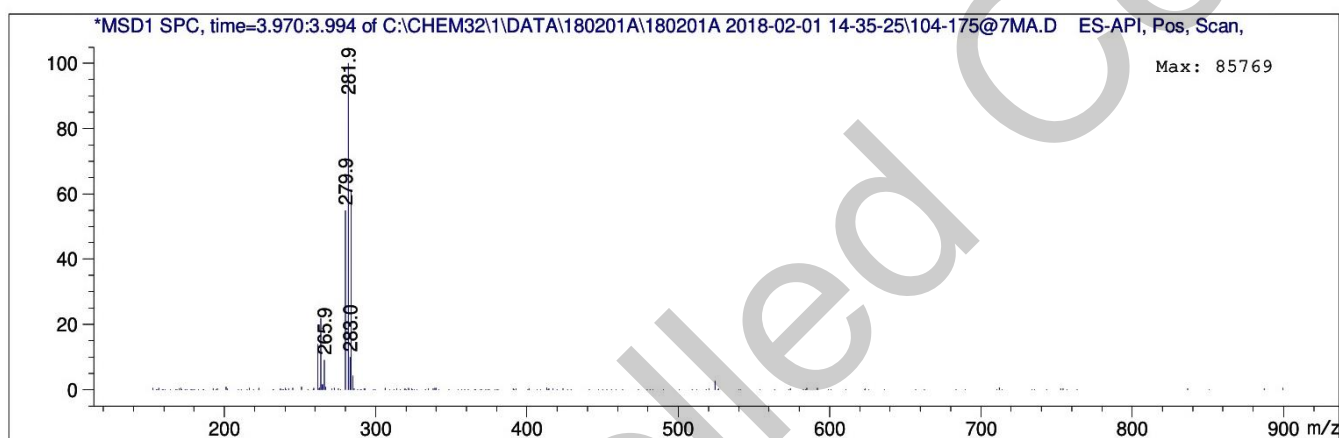
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## Ib. Mass Spectrum

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

Method: 5% to 100% ACN in water gradient (+0.1% formic acid)  
Zorbax Eclipse XDB-C8, 3.0 x 100 mm, 3.5 micron

Retention Time (MS)	MS Area	Mol. Weight or Ion
3.975	1095131	283.95   281.90   279.90   263.90   261.90

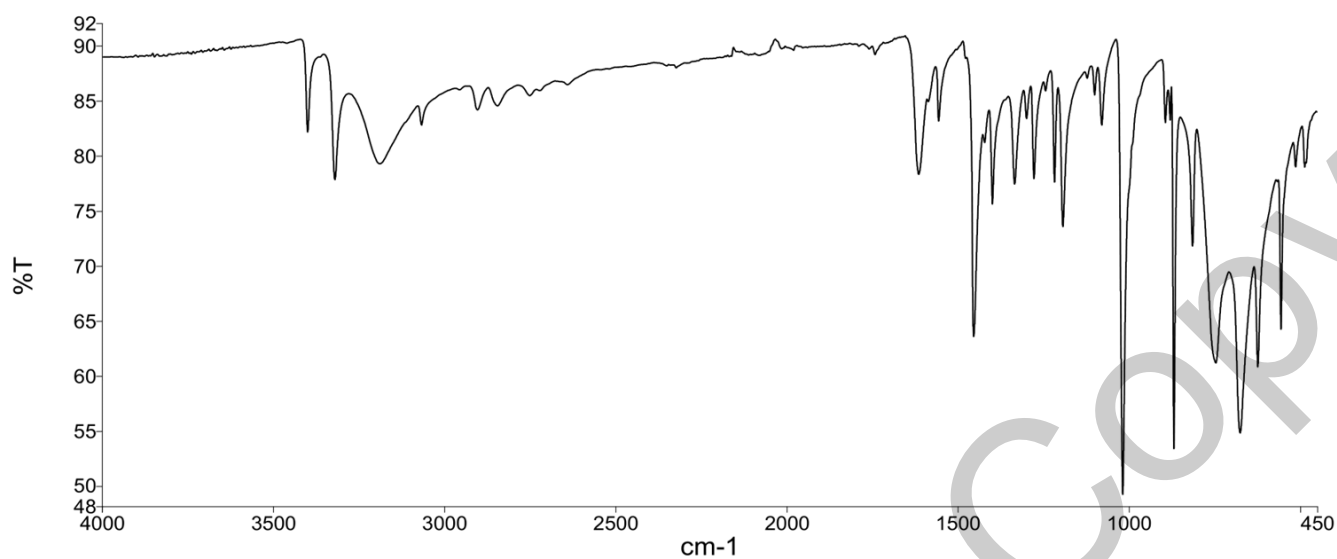


Theoretical value: 281.9 [M+H]<sup>+</sup>.

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

### Ic. IR Spectrum

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



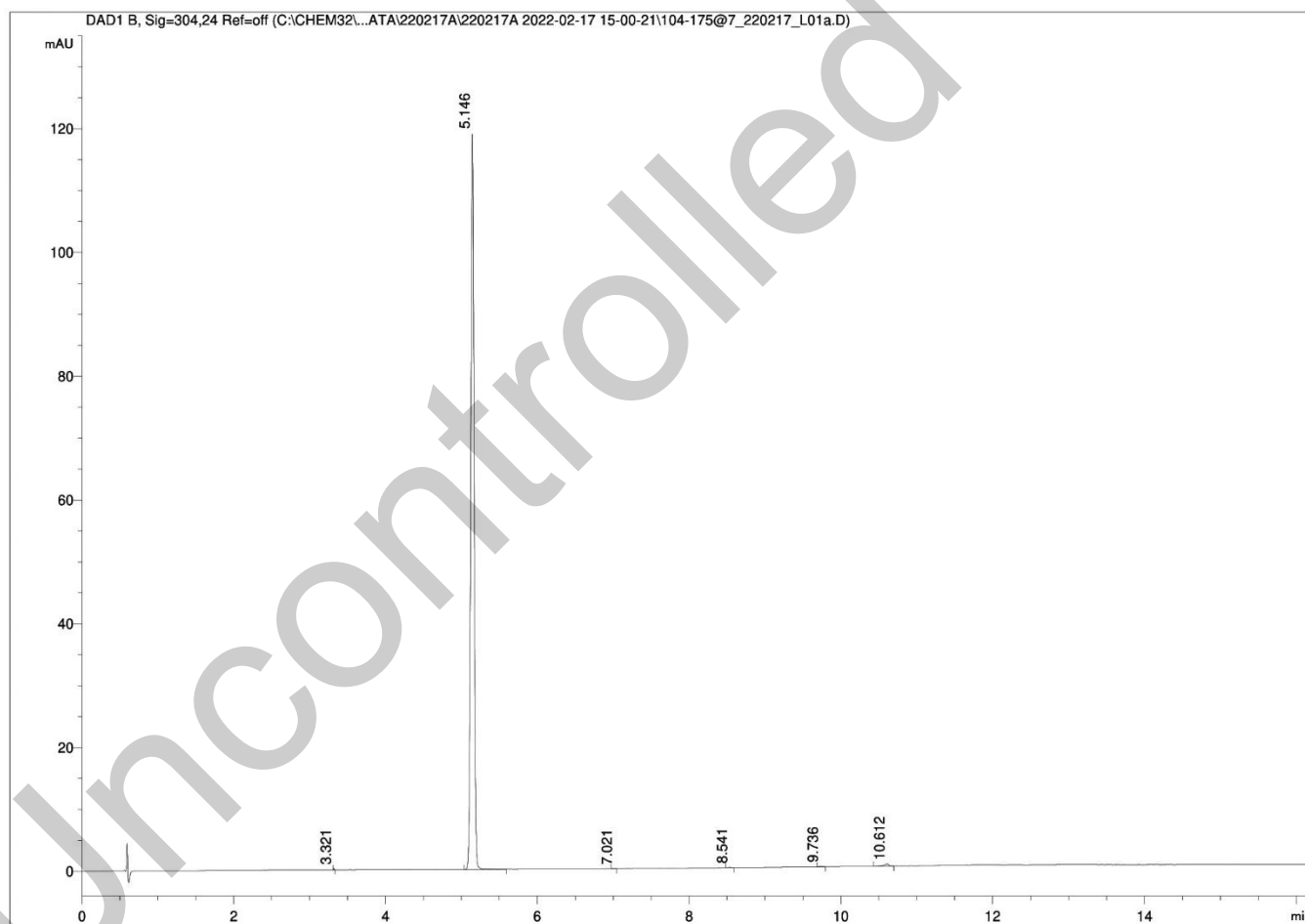
The interpretation of the signals of the Fourier-Transform Infrared Spectrum is consistent with the structural formula.

## 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 304nm	Auto 1.0 µL  0.8 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	80	20	1.0		
	5.00	50	50	1.0		
	10.00	5	95	1.0		
	15.00	5	95	1.0		
	16.00	80	20	1.0		
	19.00	80	20	1.0		



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### Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	3.32	0.15	0.04
2	5.15	350.41	99.35
3	7.02	0.05	0.01
4	8.54	0.23	0.07
5	9.74	0.24	0.07
6	10.61	1.63	0.46
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 99.3% (average of 10 duplicate runs)

### III. Water Content

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

#### Results:

Average 0.1%

### IV. Ash Content

Method: BP 2010 Ash

#### Result:

Contains 0.2% 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)	99.3%
Water content	0.1%
Ash content	0.2%
Residual solvents	<0.1%
Purity*	99.0%

This purity is assessed to be 99.0%.

Product Reviewed By:

Product Released By:

James Rixson, PhD  
Head of Production

Carol Worth, PhD  
Quality Manager

Release Date: 23 June 2022

\*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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