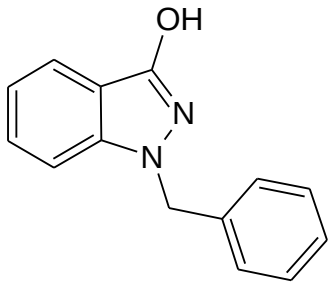
Our Formula. Your Success.

Reference Material Product Information Sheet

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



Name	1-benzyl-1*H*-indazol-3-ol
BP/EP Name	Benzydamine Impurity C
USP Name	Not listed.
Synonym(s)	1-benzyl-3-hydroxy-1*H*-indazole
Epichem Item #	EPL-AA51 Batch 3
CAS #	2215-63-6
Molecular Formula	$C_{14}H_{12}N_2O$
Molecular Weight	224.26 g/mol
Appearance	Beige powder
Melting Point	167.1-170.6°C.
Combustion Analysis	Required (%): C:75.0; H:5.4; N:12.5. Found (%): C:74.8; H:5.4; N:12.3.
Purity*	98.9%
Date of Manufacture	11 September 2018
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*

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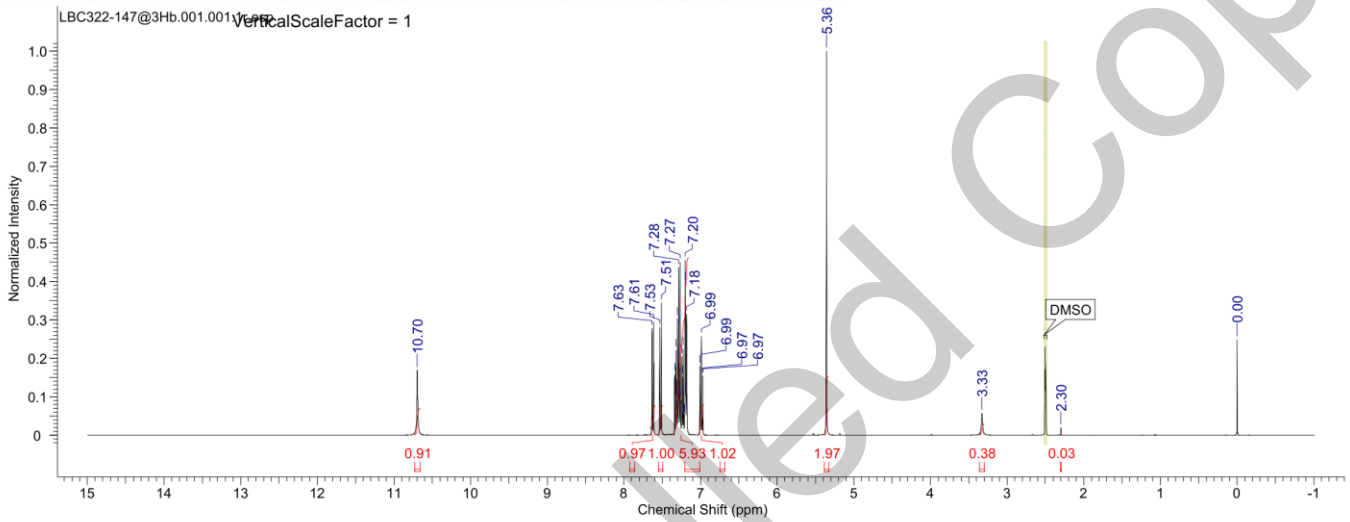
I. Identity

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

Ia. ^{1}HNMR Spectrum

Conditions: 400 MHz, DMSO-d_6
^{1}HNMR spectrum consistent with chemical structure.

Acquisition Time (sec)	3.7547	*Comment*	LBC322-147@3H 1H DMSO {E:\dataexternal\epichem} cygoh 4	*Date*	26 Sep 2018 17:55:12		
Date Stamp	26 Sep 2018 17:55:12			*File Name*	\\naphthalene\company\NMR files\LBC322\LBC322-147@3Hb\1\pdata\1\1r		
Frequency (MHz)	400.13	*Nucleus*	1H	*Number of Transients*	8	*Origin*	spect
Original Points Count	24038	*Owner*	nmr	*Points Count*	32768	*Pulse Sequence*	zg
Receiver Gain	101.00	*SW(cyclical) (Hz)*	6402.05	*Solvent*	DMSO-d6	*Spectrum Offset (Hz)*	2798.9890
Spectrum Type	STANDARD	*Sweep Width (Hz)*	6401.85	*Temperature (degree C)*	26.945		



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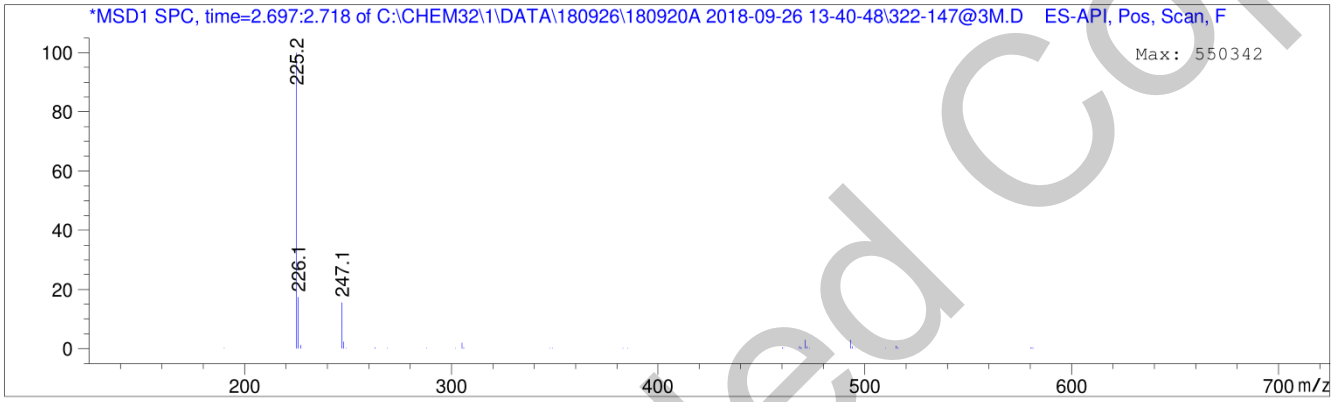
Ib. Mass Spectrum

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

Method: 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
2.705	3398690	247.10 I
		226.10 I
		225.20 I



Theoretical values: 225.2 $[M+H]^+$

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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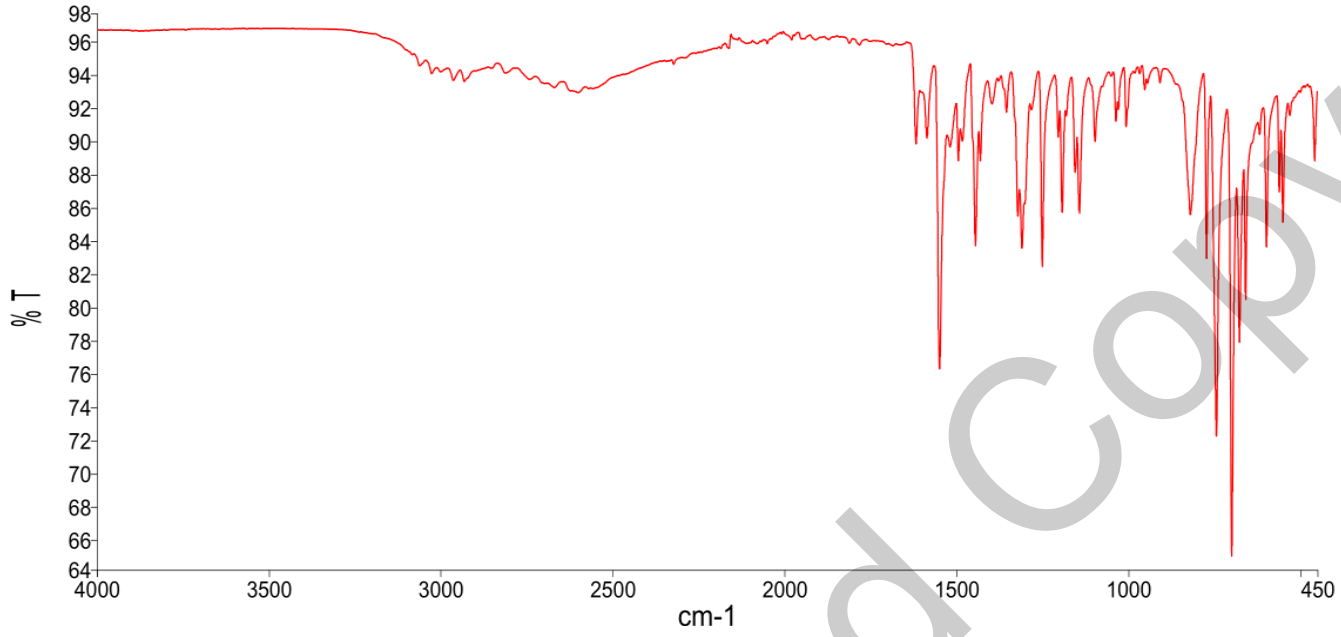
Ic. IR Spectrum

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



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

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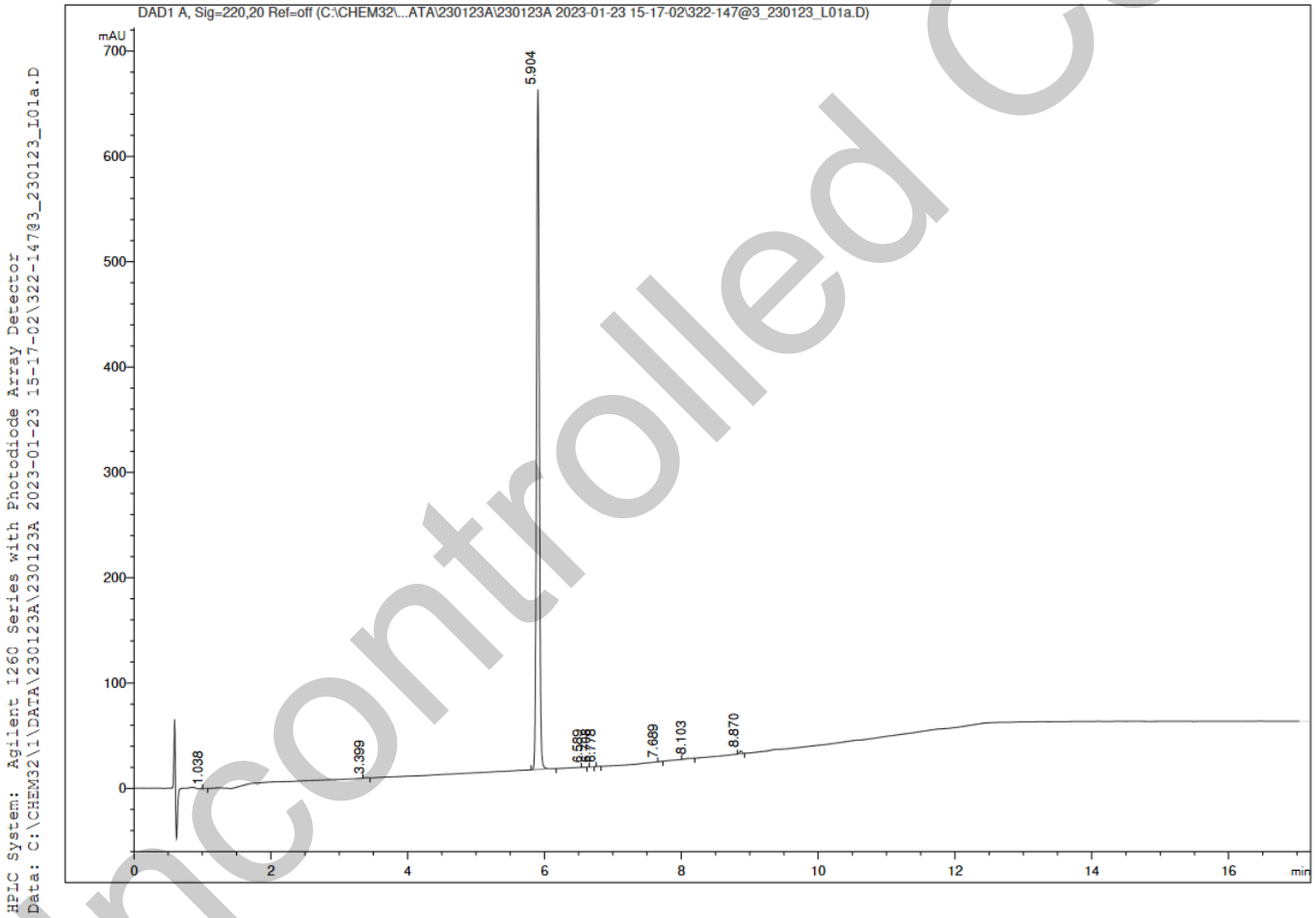
II. Purity

The purity of this material was analysed by high performance liquid chromatography (HPLC) using in-house EM005.WI07.

HPLC Conditions:

Column	Conditions				Detector	Injector
Agilent Poroshell 120 EC-C18 4.6 x 50mm 2.7 micron	25ºC				DAD 220nm	Auto 1.0 µL 0.35 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	85	15	1.0		
	6.00	55	45	1.0		
	11.00	5	95	1.0		
	16.00	5	95	1.0		
	17.00	85	15	1.0		
	20.00	85	15	1.0		



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Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	1.04	0.25	0.01
2	3.40	0.96	0.05
3	5.90	1831.69	99.43
4	6.59	0.31	0.02
5	6.71	0.20	0.01
6	6.78	0.10	0.01
7	7.69	0.28	0.02
8	8.10	2.43	0.13
9	8.87	6.03	0.33
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 99.4% (average of 10 duplicate runs)

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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 2016 Ash (Appendix XI J) as per WS001/28614
Result:
Contains <0.1% ash.

V. Residual Solvents

Method: ^{1}HNMR
Result:
0.4% Toluene detected by ^{1}H NMR analysis.

VI. Final Result

Chromatographic purity (HPLC)	99.4%
Water content	0.1%
Ash content	<0.1%
Residual solvents	0.4%
Purity*	98.9%

This purity is assessed to be 98.9%.

Product Reviewed By:

James Rixson, PhD
Head Reference Standards

Product Released By:

Carol Worth, PhD
Quality Manager
Release Date: 25 January 2023

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The calculation of the purity follows the formula:

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

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