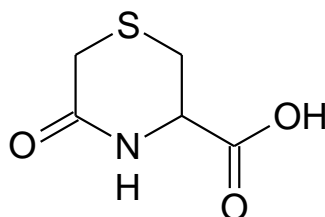


## Reference Material Product Information Sheet

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



<b>Name</b>	(RS)-5-oxothiomorpholine-3-carboxylic acid
<b>Synonym(s)</b>	(RS)-carbocysteine lactam; (RS)-3-oxo-5-carboxyperhydro-1,4-thiazine; (RS)-3-oxo-2,3,5,6-tetrahydro-4H-1,4-thiazine-5-carboxylic acid
<b>Epichem Item #</b>	EPL-AA50 Batch 1
<b>CAS #</b>	14226-97-2
<b>Molecular Formula</b>	C <sub>5</sub> H <sub>7</sub> NO <sub>3</sub> S
<b>Molecular Weight</b>	161.18 g/mol
<b>Appearance</b>	Off-white powder
<b>Melting Point</b>	184.2-190.3°C (decomposition).
<b>Combustion Analysis</b>	Required (%): C:37.3; H:4.4; N:8.7. Found (%): C:37.8; H:4.5; N:8.8.
<b>Purity*</b>	99.3%
<b>Date of Manufacture</b>	24 August 2009
<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)

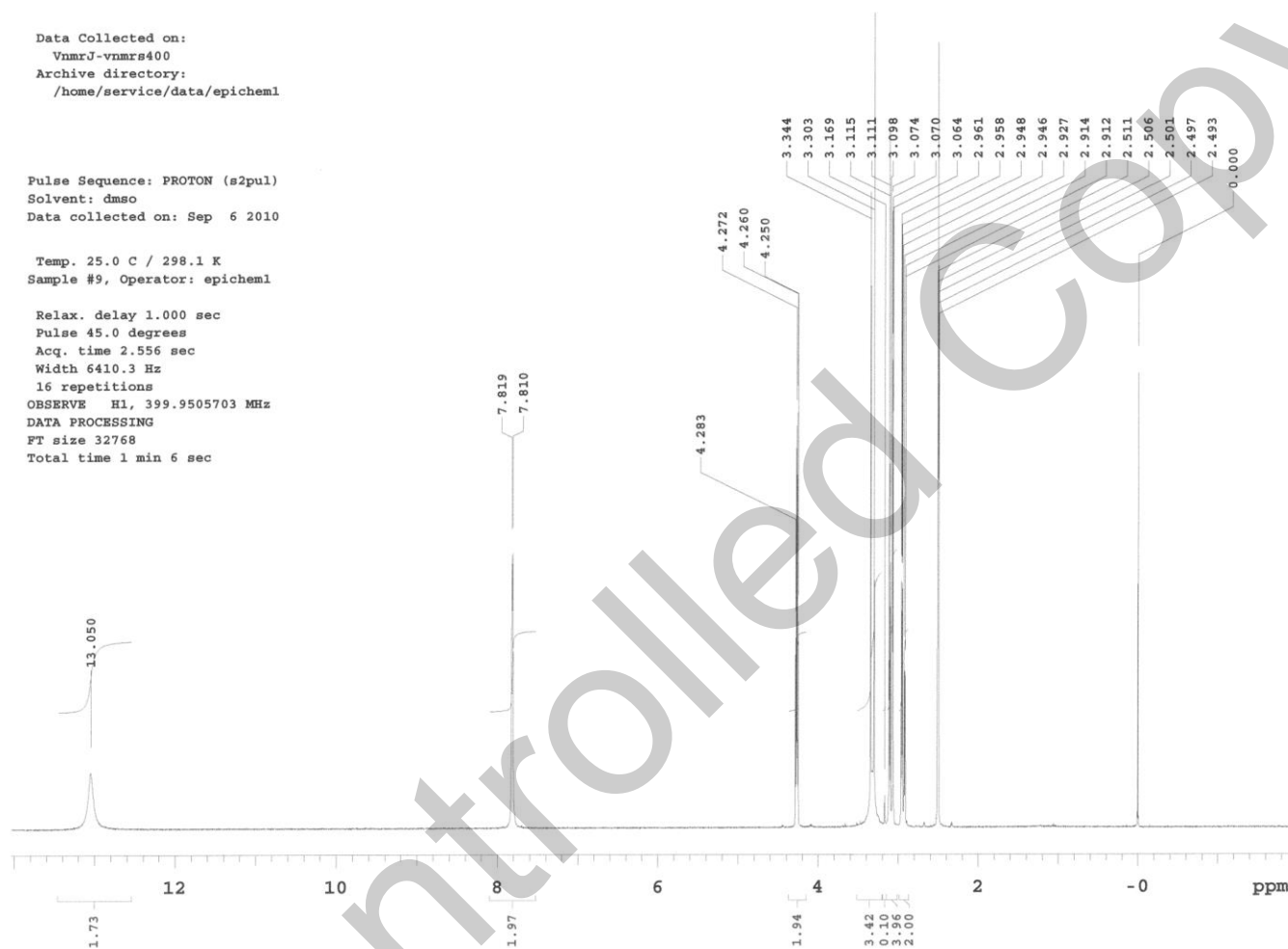
\* NATA accreditation does not cover the performance of this service

## 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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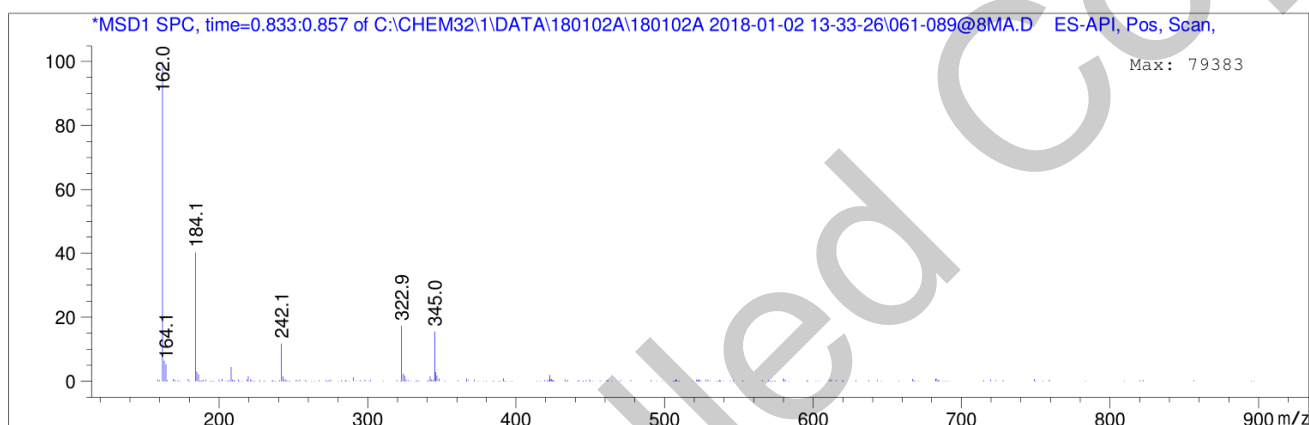
Epichem Pty Ltd, Suite 5, 3 Brodie-Hall Drive, Bentley WA 6102, Australia  
Tel + 61 (0)8 6167 5200 Fax + 61 (0)8 6167 5201 www.epichem.com.au ABN 80 106 769 902

## 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
0.842	859836	345.00 I
		322.90 I
		242.05 I
		184.05 I
		162.00 I

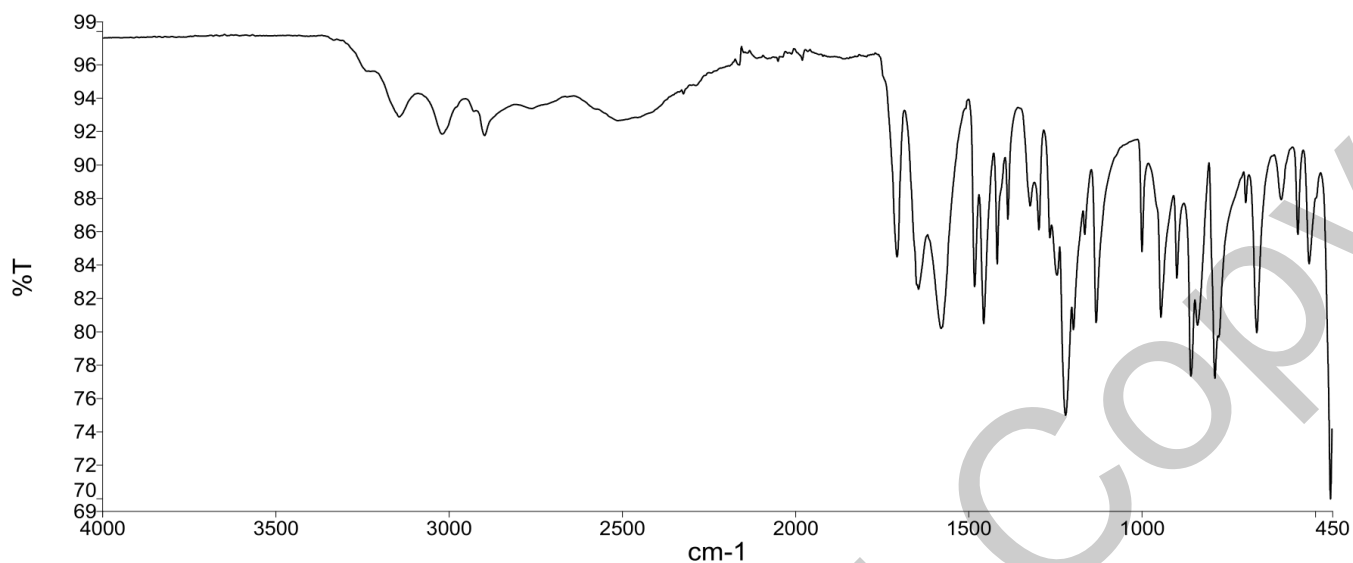


Theoretical value: 162.0 [M+H]<sup>+</sup>.

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.

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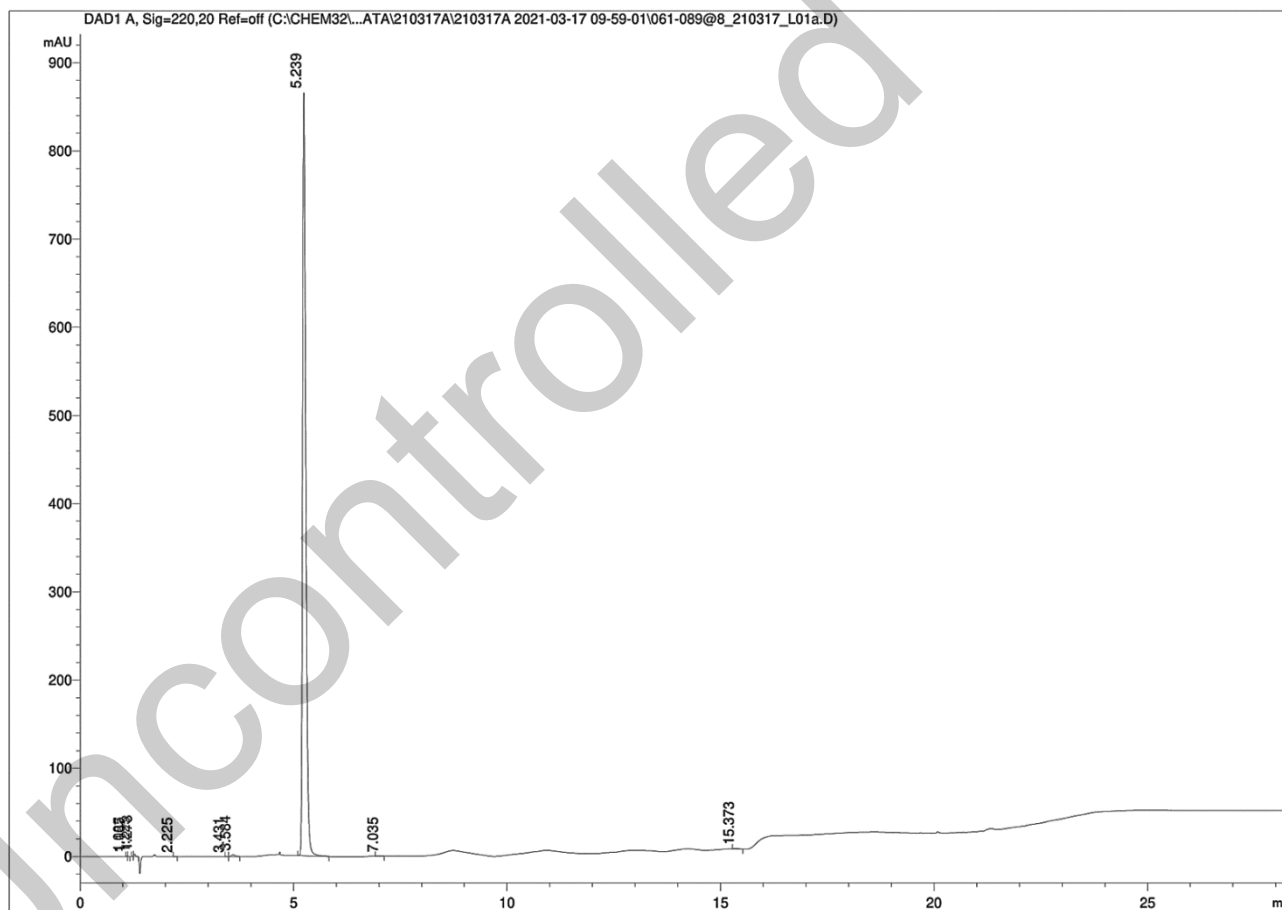
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## II. Purity

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

### HPLC Conditions:

Column	Conditions				Detector	Injector
Fortis H2o 4.6 x 100mm 3 micron	25°C				DAD 220nm	Auto 3.0 µL 3.2 mg/mL in 100% water (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	99.5	0.5	1.0		
	1.00	99.5	0.5	1.0		
	6.00	97	3	1.0		
	11.00	92	8	1.0		
	13.50	87	13	1.0		
	21.70	5	95	1.0		
	26.70	5	95	1.0		
	27.70	99.5	0.5	1.0		
33.70	99.5	0.5	1.0			



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### Area Percent Report – Sorted by Signal

Peak Number	Retention Time (rounded)	Area	Area % (rounded)
1	1.08	0.78	0.02
2	1.11	0.50	0.01
3	1.22	1.02	0.02
4	1.28	6.04	0.14
5	2.23	0.28	0.01
6	3.43	0.14	0.00
7	3.58	10.91	0.24
8	5.24	4444.90	99.46
9	7.03	1.21	0.03
10	15.37	3.24	0.07
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 99.5% (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: Combustion analysis

**Result:**

Contains 0.1% ash.

### V. Residual Solvents

Method: <sup>1</sup>H NMR

**Result:**

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

### VI. Final Result

Chromatographic purity (HPLC)	99.5%
Water content	0.1%
Ash content	0.1%
Residual solvents	<0.1%
Purity*	99.3%

This purity is assessed to be 99.3%.

Product Reviewed By:

James Rixson, PhD  
Head of Production

Product Released By:

Carol Worth, PhD  
Quality Manager  
Release Date: 23 August 2021

*\*NATA accreditation does not cover the performance of this service.*

The calculation of the purity follows the formula:

$$\text{Purity(\%)} = \frac{((\text{Chromatographic purity[HPLC]}) \times (100 - (\text{water content} + \text{ash content} + \text{volatile contents})))}{100}$$

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