

Certificate of Analysis

Characterisation methods are accredited according to

Reference Material

Product name

(2S,3S)-2-Amino-3-methyl-3-sulfino-4-(1H-1,2,3triazol-1-yl)butyric Acid

Product code Lot number MM0989.01-0025 W1008344 **CAS** number **Appearance** not listed light yellow solid Molecular weight Melting point (DSC)

248 26 190 °C

Molecular formula

Long-term storage $C_7H_{12}N_4O_4S$ 2 to 8 °C, dark

99.1 %

Uncertainty² U 0.5 %

Intended Use: Use for identification and quantification, The assay is verified by a second testing method.

25 Oct 2019 Date of shipment:

Producer confirms that this reference material (RM) meets the specification detailed on this Certificate of Analysis for one year from the date of shipment, provided the substance is stored under the recommended conditions unopened in the original container.

Release by: Date of Release:	0	Due doort Delegan
Dr. Sabine Schröder Luckenwalde, 18 Oct 2019	Jarol	Product Release

¹ Calibration and verification were carried out using standards traceable to SI-units. The value is expressed on an "as is" basis.

Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCiPACT $^{\text{TM}}$) Test methods used for characterisation are accredited to ISO/IEC 17025 | DAkkS D-PL-14176-01-00

Producer: LGC GmbH Louis-Pasteur-Str. 30 D-14943 Luckenwalde Germany www.lgcstandards.com Page 1/8

² The uncertainty "U" is the expanded uncertainty of the testing method for the assigned value estimated in accordance with the Guide to the Expression of Uncertainty in Measurement (GUM). It corresponds to a level of confidence of about 95%. Coverage factor k = 2.



Product information

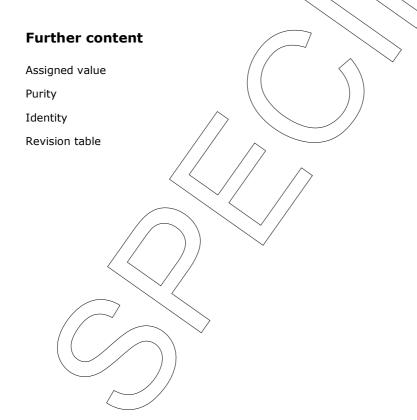
This RM is intended for laboratory use only and is not suitable for human or animal consumption.

This RM conforms to the characteristics of a primary standard as described in the ICH Guidelines. The values quoted in this Certificate of Analysis are the producer's best estimate of the true values within the stated uncertainties and based on the techniques described in this Certificate of Analysis. The characterisation of this material was undertaken in accordance with the requirements of ISO/IEC 17025. The identity is verified by data from international scientific literature.

Storage and handling

Before usage of the RM, it should be allowed to warm to room temperature. No drying is required, as assigned values are already corrected for the content of water and other volatile materials.

Reference Material quality is controlled by regularly performed quality control tests (retests).





Assigned value

Assay "as is":

99.14 %; U = 0.48 %

The assay "as is" is assessed by carbon titration of elemental analysis and is equivalent to the assay based on the not-anhydrous and not-dried substance. The assay is verified by 100% method (mass balance). The verified result lies inside our acceptance criteria, i.e. less than 1.0 % difference to assay assigning technique.

For quantitative applications, use the assay as a calculation value on the as is basis. The uncertainty of the assay can be used for estimation/calculation of measurement uncertainty.

Method 1: Value assigning technique - carbon titration of elemental analysis

Method

percentage carbon found in relation to percentage carbon as calculated for molecular formula

Result (mass fraction, **n = 3**) 99.14 %; U = 0.48 %

Method 2: Value verifying technique - 100% method

100% method (mass balance) with chromatographic purity by HPLC

Result 98. ₹4 %

The calculation of the 100% method follows the formula:

Assay (%) = (100 % - volatile contents (%)) * $\frac{\text{Purity (%)}}{100 \%}$

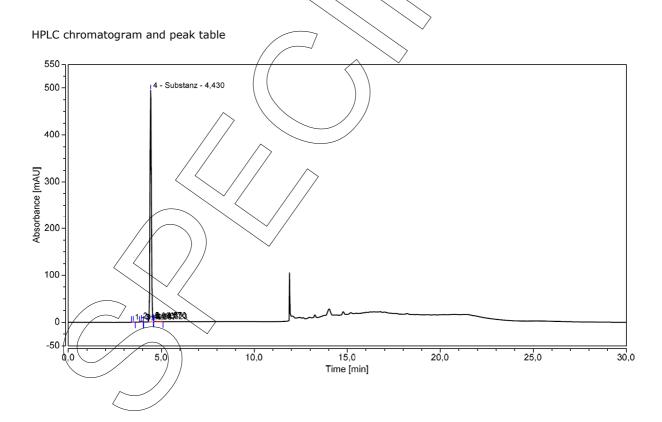
Volatile contents are considered as absolute contributions and purity is considered as relative contribution. Inorganic residues are excluded by additional tests.



Purity

Purity by High Performance Liquid Chromatography (HPLC)

HPLC conditions:	
Column	Luna C18(2); 5 μm, 250 x 4.6 mm
Column temperature	40 °C
Detector	DAD, 210 nm //
Injector	Auto 2 μl; 0.168 mg/ml in Water/Acetonitrile 50/50 (v/v)
Flow rate	1.0 ml/min
Phase A	1.32 g/l (NH₄)₂PQ₄, pH 2.5
Phase B	phase A/Acetonitrile 20/80
Gradient program	0-6 min A/B 100/0 6-13 min A/B to 30/70 13-18 min A/B 30/70 18-21 min A/B to 100/0 21-30 min A/B 100/0 (v/v)





Area percent report - sorted by signal				
Pk #	Retention time	Area	Area %	
1	3.490	0.038	0.09	
2	3.933	0.245	0.57	
3	4.037	0.004	0.01	
4	4.430	42.624	98.7/3	\rightarrow
5	4.570	0.058	0.13	
6	4.623	0.203	0.47	
Totals		43.173	100.00	

The content of the analyte was determined as ratio of the peak area of the analyte and the cumulative areas of the purities, added up to 100 %. System peaks were ignored in calculation.

Result (n = 3)

98.74 %; U = 0.18 %

Volatile content

Water content

Method

Result (n = 3)

No significant amounts of water were detected (< 0.05 %).*

*not accredited testing method

Residual solv		//	/
Method	\ /		¹ H-NMR

Result (n = 1) No significant amounts of residual solvents were detected (< 0.05 %).

Inorganic residues

Method: Elementary analysis

Inorganic residues can be excluded by elementary analysis (CHN).

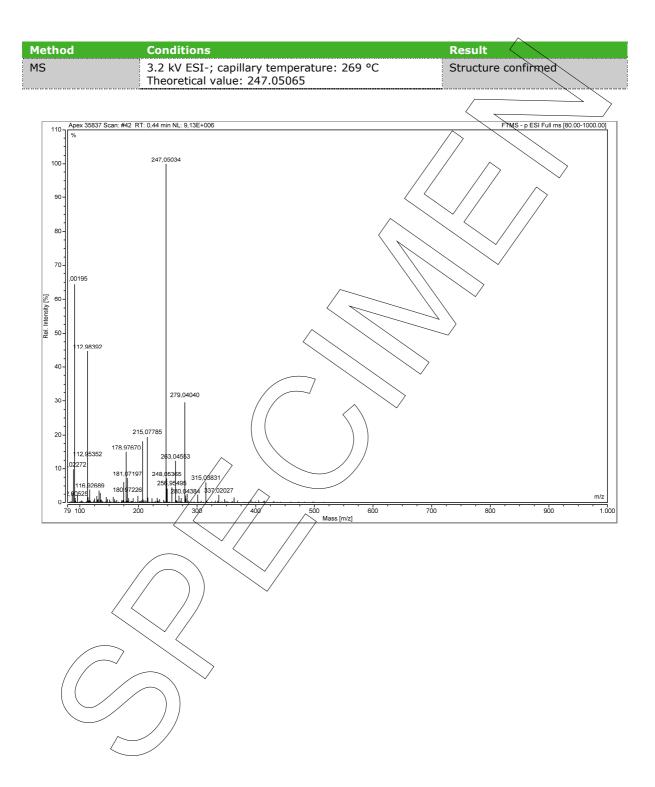
LGC GmbH, Louis-Pasteur-Str. 30, D-14943 Luckenwalde, Germany
MM0989.01-0025
Lot number W1008344

^{*}not accredited testing method

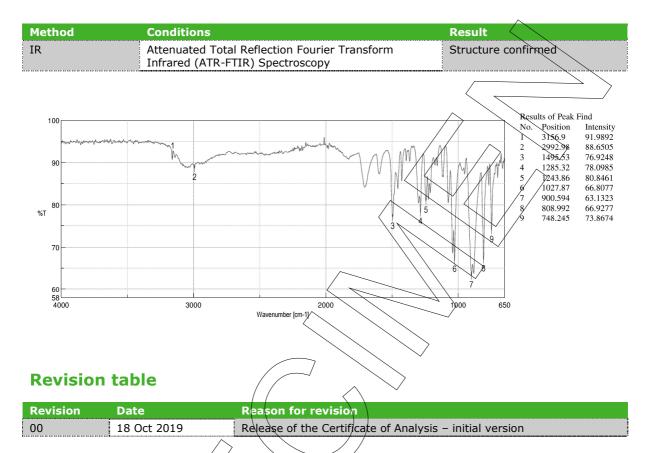


Identity The identity is assessed by ISO/IEC 17025 accredited testing methods. Method Conditions Result ¹H-NMR 400 MHz, D₂O/NaOD, 40 % in D₂O Structure confirmed









Product warranties for the RM are set out in the terms and conditions of purchase.

