



Important 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 production of this RM was undertaken in accordance with the requirements of ISO 17034. 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.

Further content

Assigned value

Purity

Identity

Stability and homogeneity

Revision table



Assigned value

Assay "as is": **99.18 %; U = 0.43 %**

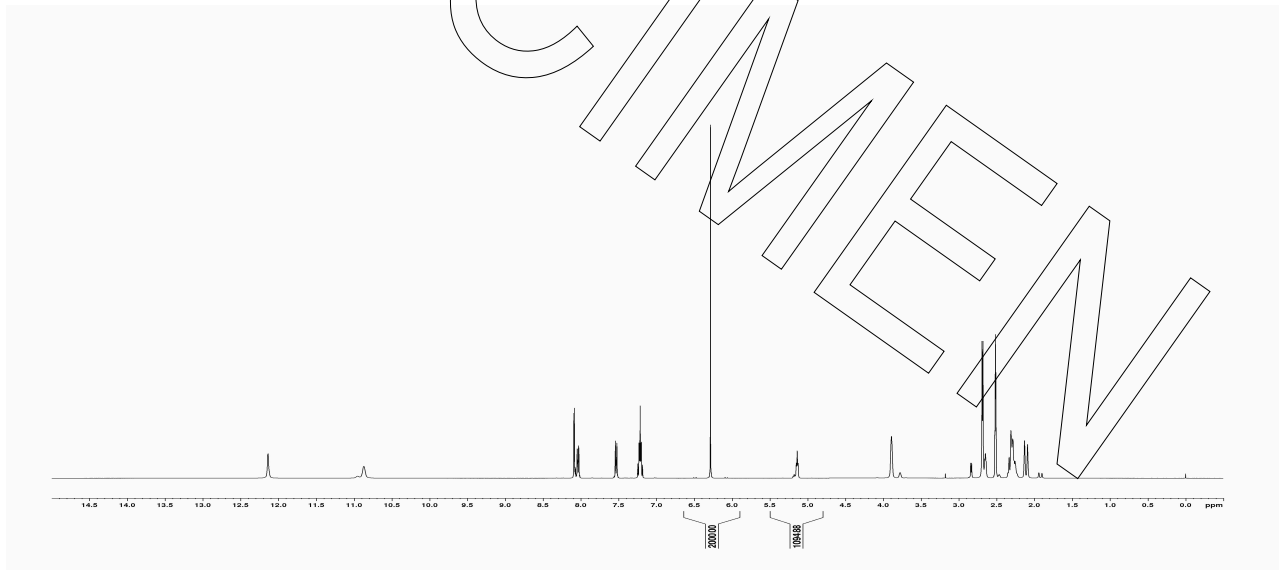
The assay "as is" is assessed by quantitative NMR spectroscopy and is equivalent to the assay based on the not-anhydrous and not-dried substance. The assay is verified by 100% method (mass balance).

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 - quantitative NMR spectroscopy	
Conditions	400 MHz, DMSO-d ₆
Internal Standard	Maleic acid (certified reference material), signal 5.9 - 6.6 ppm, 2 H
Result (mass fraction, n = 6)	99.18 %; U = 0.43 %

Quantitative NMR spectrum





Method 2: Value verifying technique - 100% method

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

Result

99.88 %

The calculation of the 100% method follows the formula:

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

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

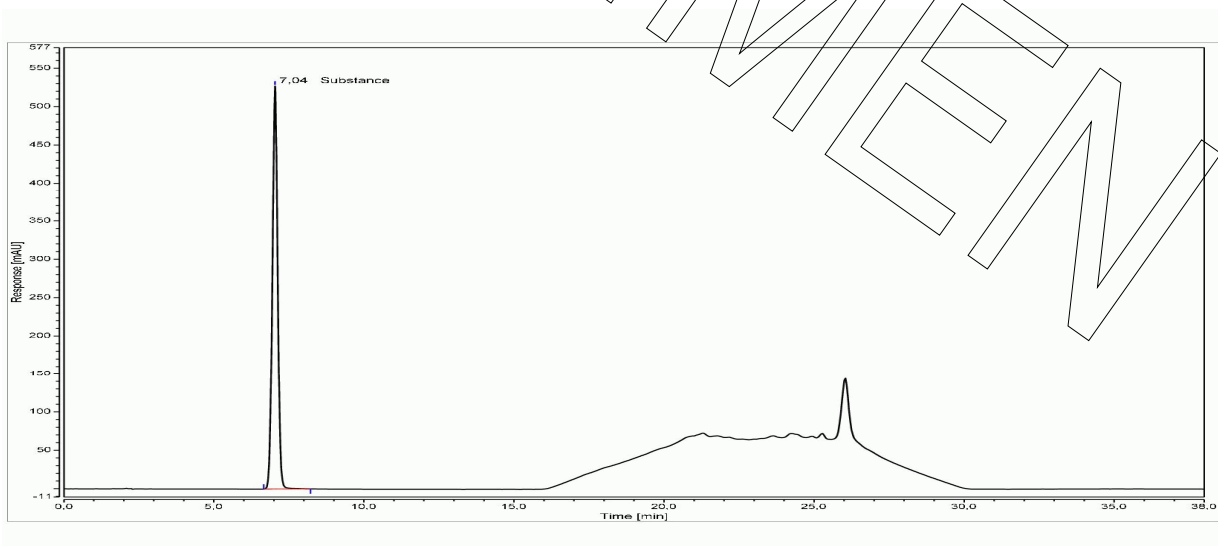


Purity

Purity by high performance liquid chromatography (HPLC)

HPLC Conditions:	
Column	Hypersil Gold C18; 5 µm, 150 x 4.6 mm
Column temperature	40 °C
Detector	DAD, 210 nm
Injector	Auto 2 µl; 0.1006 mg/ml in Acetonitrile/Water 50/50 (v/v)
Flow rate	1.0 ml/min
Phase A	Water, 0.1 % H ₃ PO ₄
Phase B	Acetonitrile, 0.1 % H ₃ PO ₄
Gradient program	0-13 min A/B 82/18 13-18 min A/B to 35/65 18-23 min A/B 35/65 23-27 min A/B to 82/18 27-38 min A/B 82/18 (v/v)

HPLC chromatogram and peak table





Area percent report - sorted by signal

Pk #	Retention time	Area	Area %
1	7.039	107.8217	100.00
Totals		107.8217	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 = 10) 100.00 %; U = 0.18 %

Volatile content

Water content

Method	Karl Fischer titration
Result (n = 3)	0.07 %; U = 0.04 %

Residual solvents

Method	GC headspace
Result (n = 3)	Sum: 0.05 %; U = 0.01 % 0.05 % Methanol

Inorganic residues

Sulphated Ash

Method	EP 10.3, chapter 2.4.14
Result (n = 1)	0.15 %*

*not accredited testing method



Identity

The identity is assessed by ISO/IEC 17025 accredited testing methods.

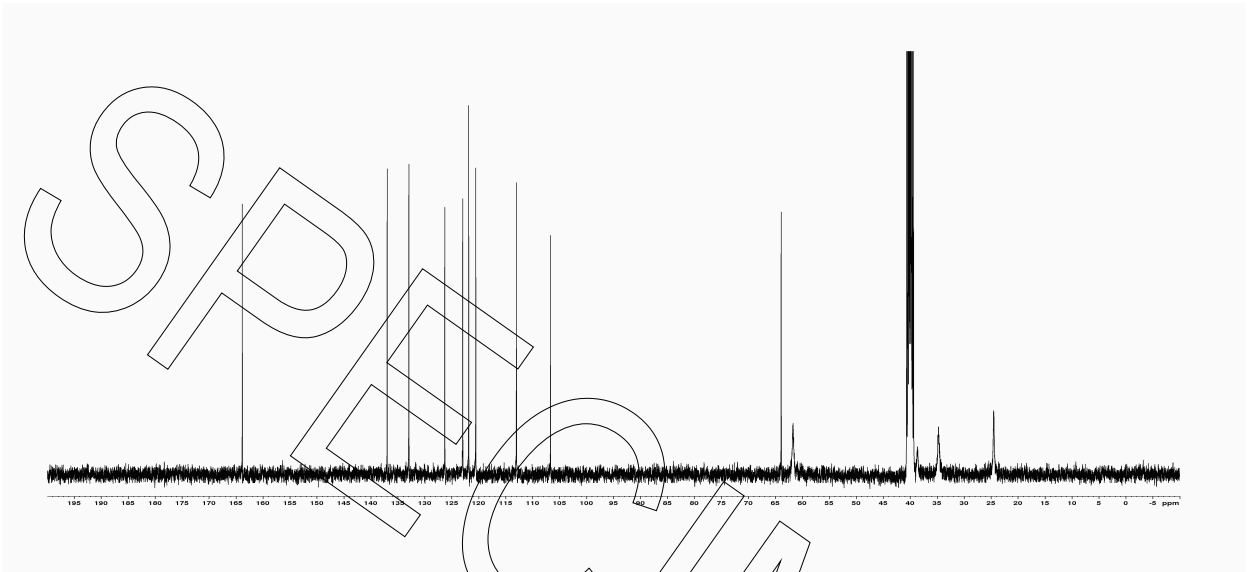
Method	Conditions	Result
¹ H-NMR	400 MHz, DMSO-d ₆	Structure confirmed





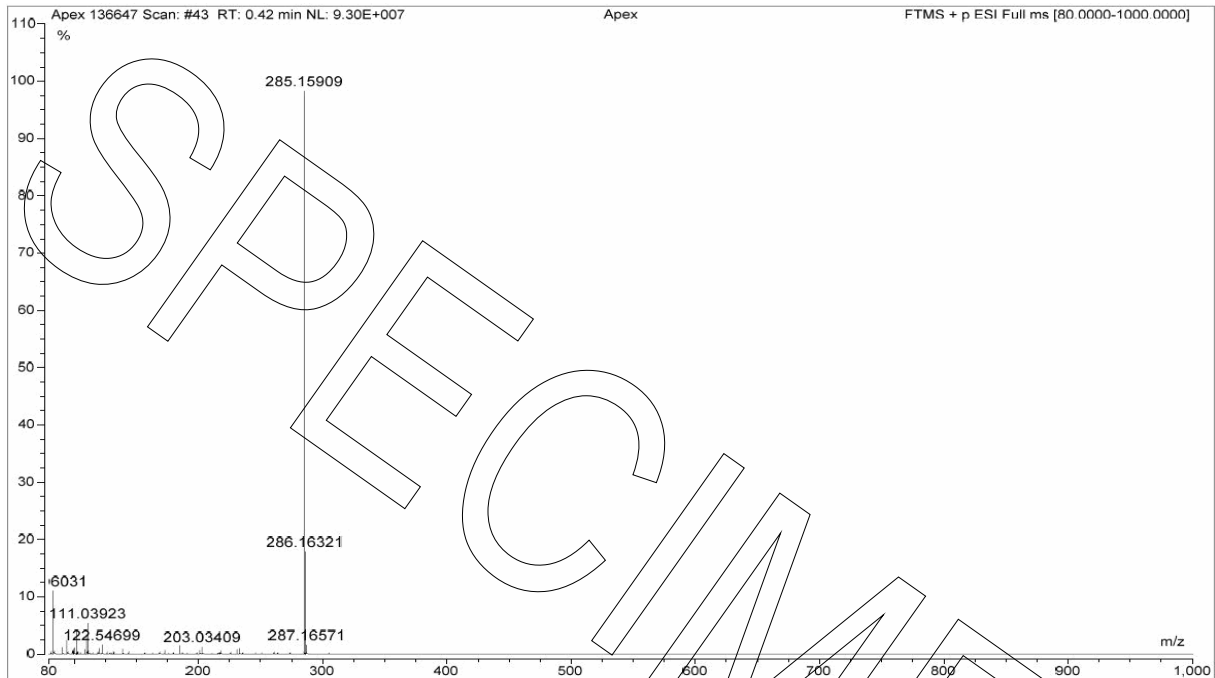
Mikromol™

Method	Conditions	Result
¹³ C-NMR	100 MHz, DMSO-d ₆	Structure confirmed





Method	Conditions	Result
MS	3.5 kV ESI+; capillary temperature: 269 °C Theoretical value: 285.15975	Structure confirmed





Method	Conditions	Result
IR	Attenuated Total Reflection Fourier Transform Infrared (ATR-FTIR) Spectroscopy	Structure confirmed



No.	Position	Intensity
1	3212.83	87.4517
2	3102.9	88.2254
3	2964.05	89.0474
4	2496.4	77.9057
5	1686.44	68.2453
6	1523.49	81.6469
7	1428.03	73.103
8	1311.36	75.7466
9	1242.9	74.8497
10	1175.4	68.6238
11	1123.33	78.3097
12	1081.87	78.2781
13	1032.69	62.3359
14	741.496	67.0408

Stability and Homogeneity

The assessment of stability indicates no significant instability. The given validity period is based on this data. This is backed up by additional stability testing and historical data over the range of several years.

RM quality is controlled by regularly performed quality control tests (re tests). Homogeneity assured by qualified process of preparation and verified by homogeneity testing.

Revision table

Revision	Date	Reason for revision
00	08 Sep 2022	Release of the Certificate of Analysis - initial version

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