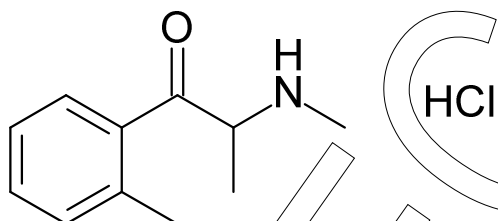




# Certificate of Analysis

## Reference Standard

2-MMC HCl (2-Methylmethcathinone Hydrochloride)



Molecular Formula:  $C_{11}H_{15}NO \cdot HCl$   
Molecular Weight: 213.70  
CAS Number: 1246815-51-9

Catalogue Number: LGCFOR1387.02  
Lot Number: 72240  
Long-term Storage: 2 to 8 °C, dark  
Appearance: white solid  
Melting Point: 167 °C (dec.)  
Assay 'as is': 98.8 %

Date of shipment: **2020-November-30**

This certificate is valid for two years from the date of shipment provided the substance is stored under the recommended conditions unopened in the original container.

LGC Quality

ISO 9001:2008  
DQS 102448 QM08

LGC GmbH, Im Biotechnologiepark, TGZ II, D-14943 Luckenwalde, Germany

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6 Pages

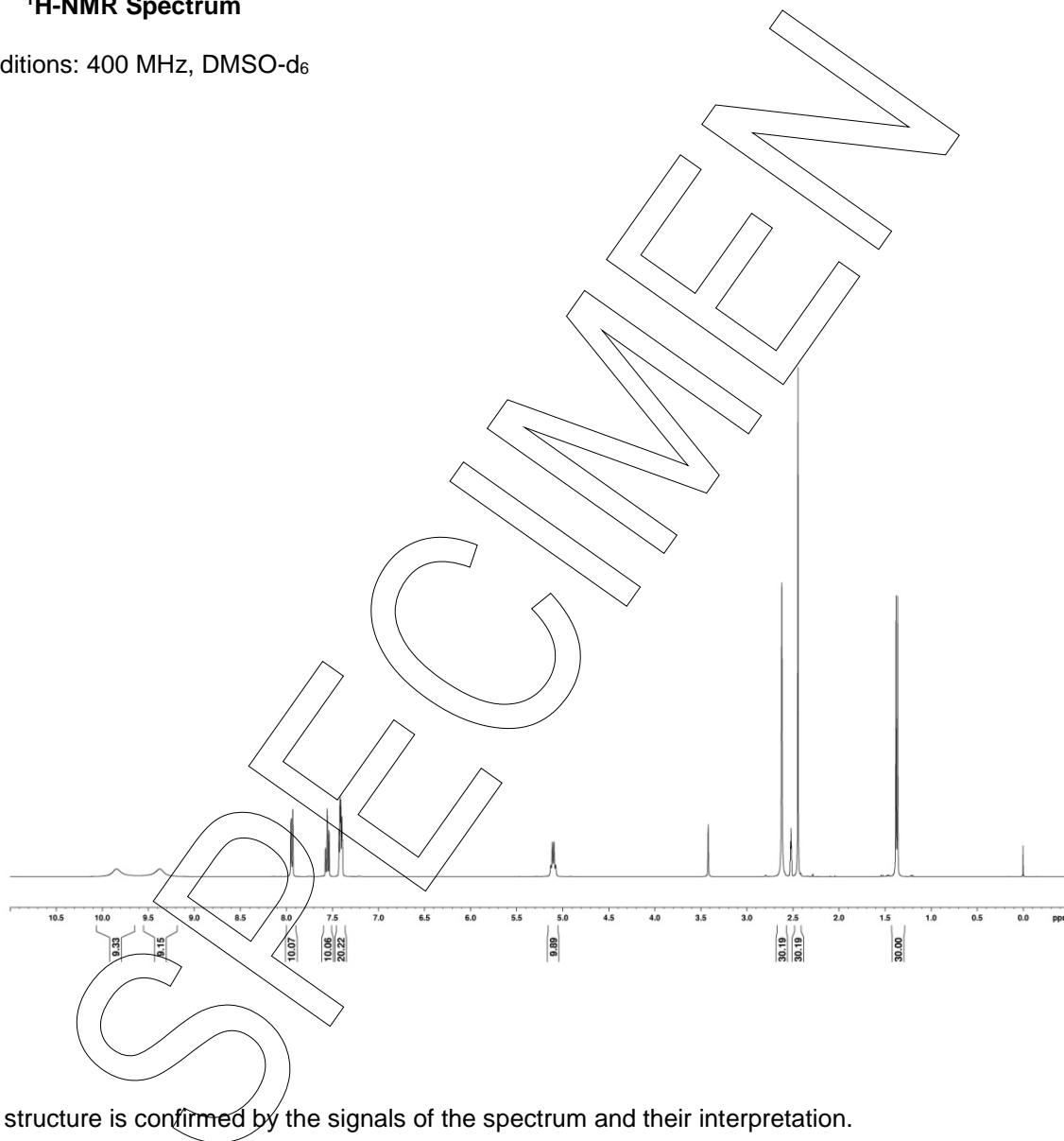


## I. Identity

The identity of the reference substance was established by following analyses.

### Ia. $^1\text{H}$ -NMR Spectrum

Conditions: 400 MHz, DMSO- $d_6$

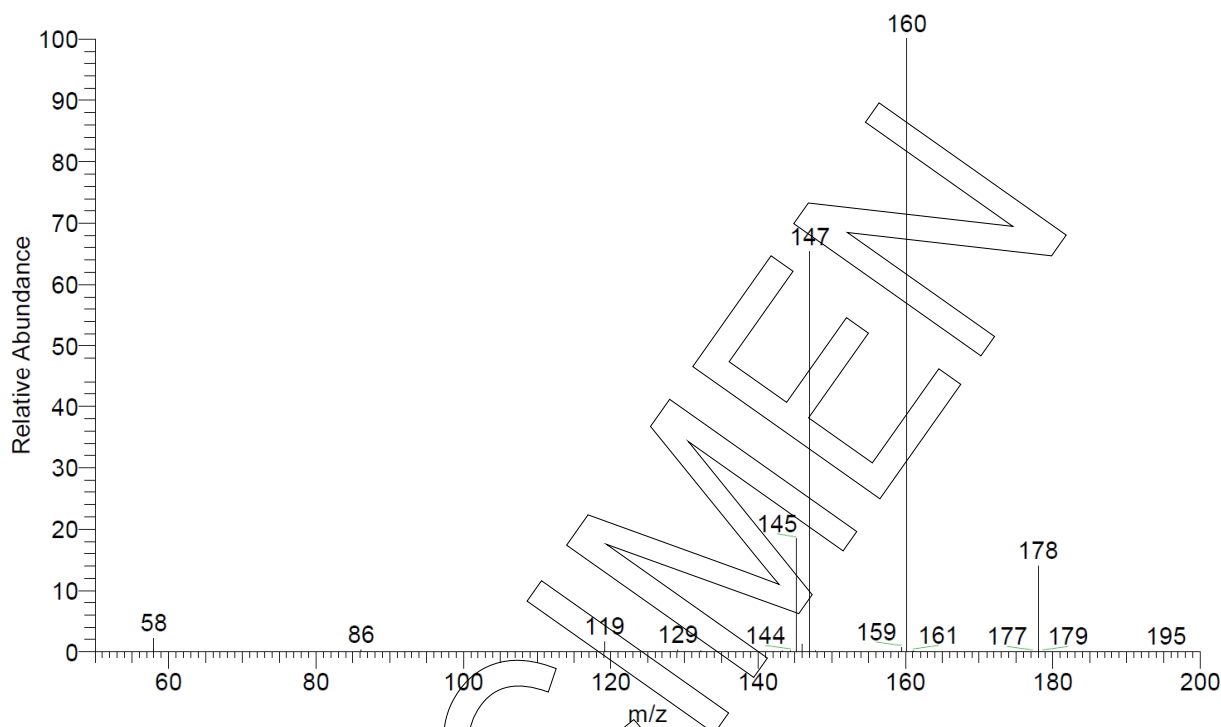


The structure is confirmed by the signals of the spectrum and their interpretation.



## Ib. Mass Spectrum

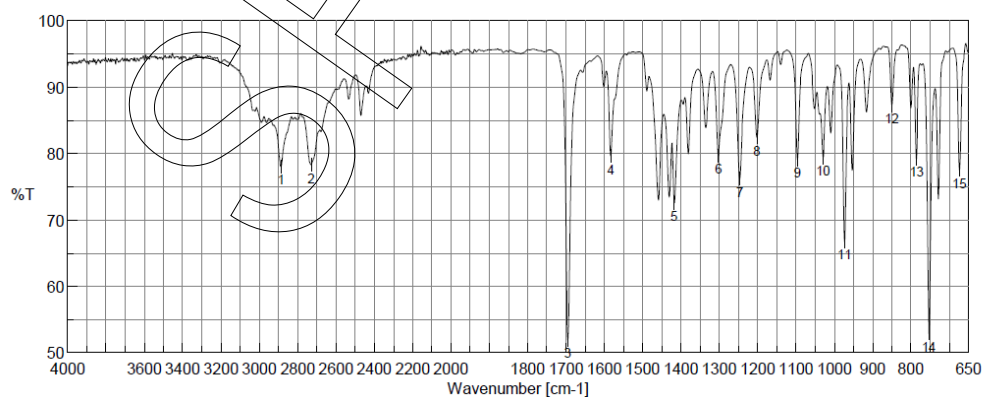
Method: 4.5 kV ESI+; vaporization temperature: 200 °C



The signals of the mass spectrum and their interpretation are consistent with the structural formula.

## Ic. IR Spectrum

Method: Attenuated Total Reflection Fourier Transform Infrared (ATR-FTIR) Spectroscopy



Result of Peak Picking		
No.	Position	Intensity
1	2886.92	78.0374
2	2729.74	78.2779
3	1695.12	51.9172
4	1583.27	79.5876
5	1417.42	72.5508
6	1302.68	79.6489
7	1246.75	76.1766
8	1201.43	82.4088
9	1096.33	79.0211
10	1029.8	79.4161
11	973.876	66.7361
12	850.454	87.2821
13	785.85	79.154
14	753.066	52.8824
15	673.999	77.5201

The signals of the IR spectrum and their interpretation are consistent with the structural formula.



## II. Purity

### IIa. High Performance Liquid Chromatography (HPLC)

The purity of the reference substance was analysed by high performance liquid chromatography (HPLC).

#### HPLC Conditions:

##### Column:

Hypersil Gold C18  
5  $\mu$ m, 150 x 4.6 mm

##### Conditions:

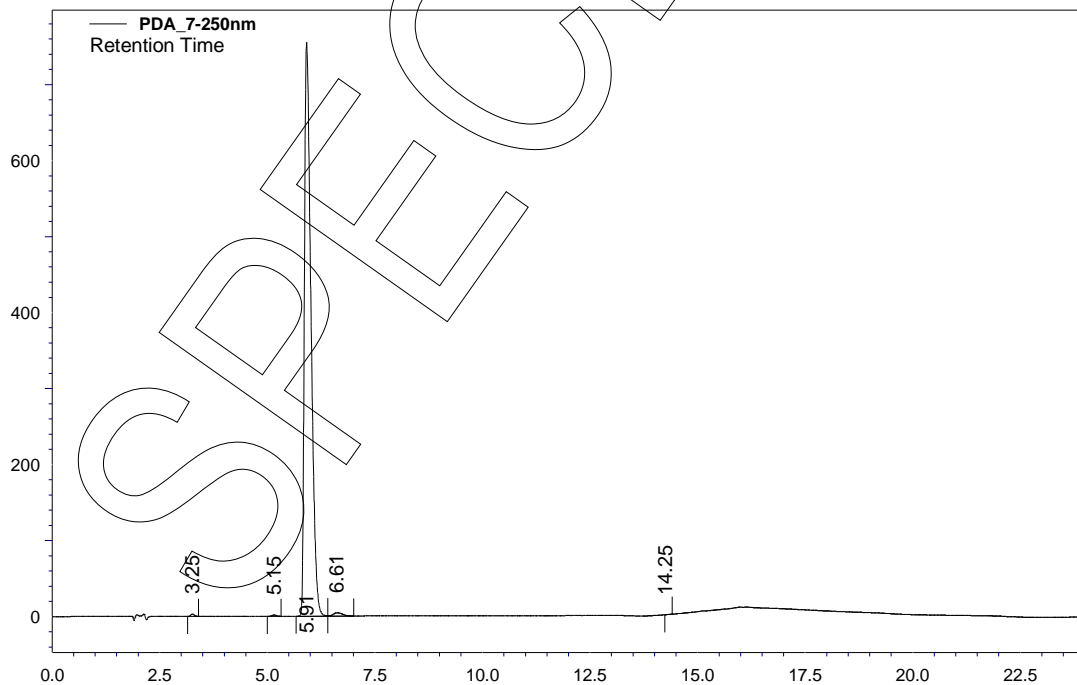
1.0 ml/min, 40 °C  
0 – 10 min Water/Acetonitrile 90/10  
10 – 13 min Water/Acetonitrile to 50/50  
13 – 16 min Water/Acetonitrile to 90/10  
16 – 24 min Water/Acetonitrile 90/10 (v/v);  
0.1 % H<sub>3</sub>PO<sub>4</sub>

##### Detector:

DAD  
250 nm

##### Injector:

Auto  
2  $\mu$ l; 0.256 mg/ml in  
Methanol





### Area Percent Report - Sorted by Signal

Pk #	Retention Time	Area	Area %
1	3.25	15959	0.20
2	5.15	11410	0.14
3	5.91	8014489	98.88
4	6.61	62847	0.78
5	14.25	763	0.01
Totals		8105468	100.00

For the calculation the system peaks were ignored. 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 %.

#### Results:

Average 98.88 %  
Number of results n=3  
Standard deviation < 0.01 %

#### IIb. Water Content

Method: Karl Fischer titration

#### Results:

Average 0.07 %  
Number of results n=3  
Standard deviation 0.01 %

#### IIc. Residual Solvents

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

No significant amounts of residual solvents were detected (< 0.05 %).



### III. Final Result

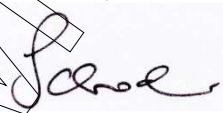
<b>Chromatographic purity (HPLC)</b>	98.88 %
<b>Water content</b>	0.07 %
<b>Residual solvents</b>	No significant amounts of residual solvents were detected (< 0.05 %).
<b>Assay (100 % method)<sup>1</sup></b>	98.81 %

The assay is assessed to be 98.8 % 'as is'

The assay 'as is' is equivalent to the assay based on the not anhydrous and not dried substance respectively.

Release Date:

Luckenwalde, 2015-03-05



Dr. Sabine Schröder  
Product Release

<sup>1</sup> 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, purity is considered as relative contribution.

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