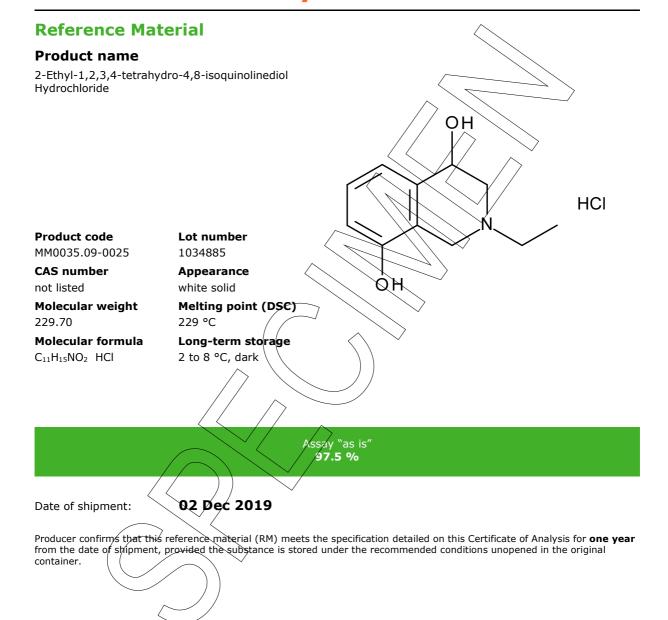


Certificate of Analysis

ISO 9001



Release by:	Date of Release:	P	Draduat Palanca	
Dr. Sabine Schröder	Luckenwalde, 08 Oct 2019	Jarol	Product Release	

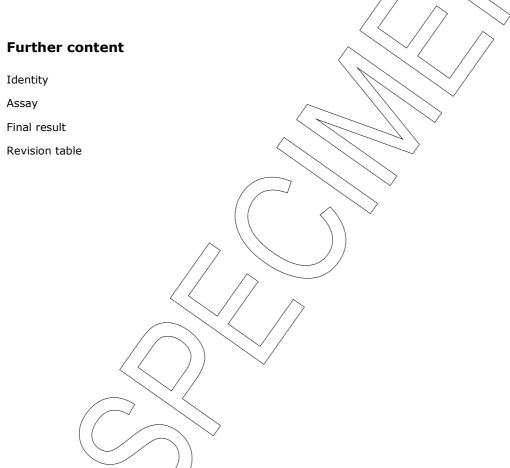


Product information

For laboratory use only. Not suitable for human or animal consumption.

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

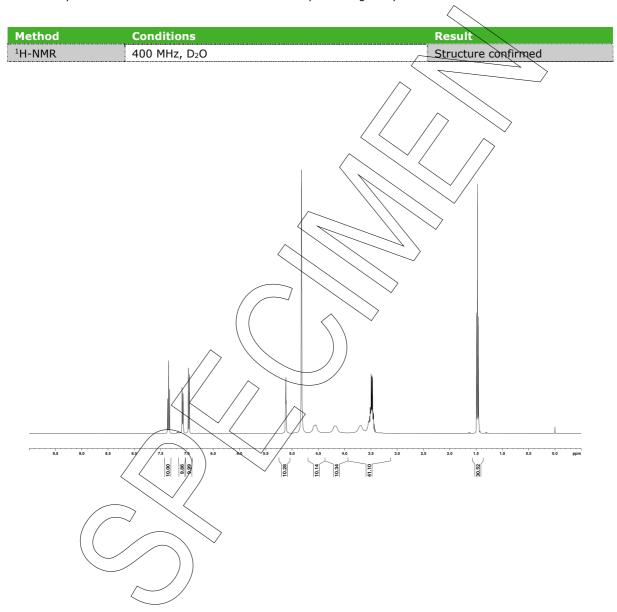
The product quality is controlled by regularly performed quality control tests (retests).





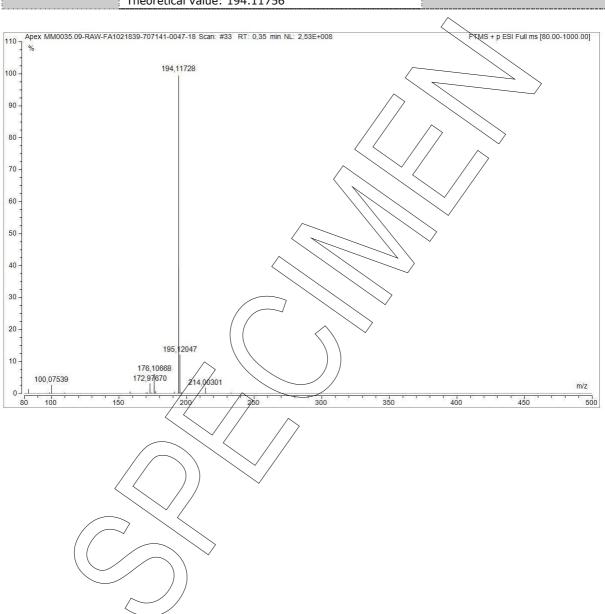
Identity

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

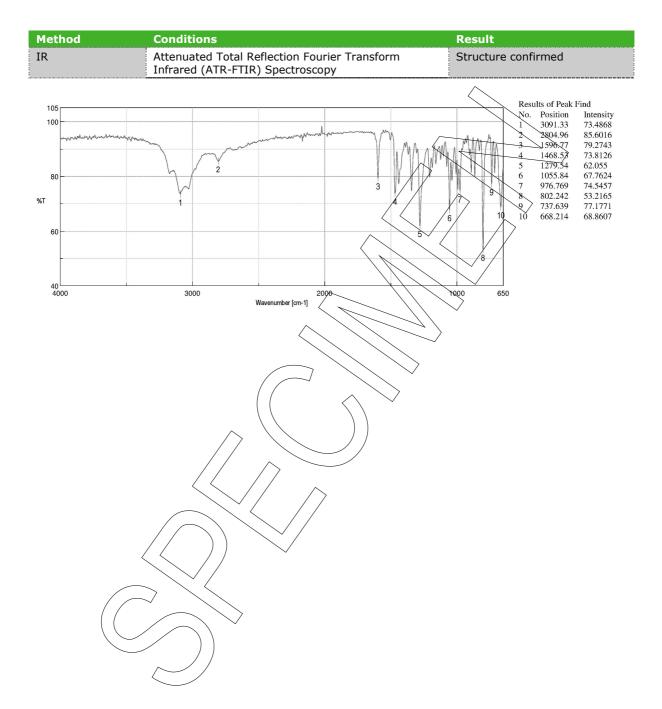




Method	Conditions	Result
MS	3.5 kV ESI+; capillary temperature: 269 °C	Structure confirmed
	Theoretical value: 194.11756	







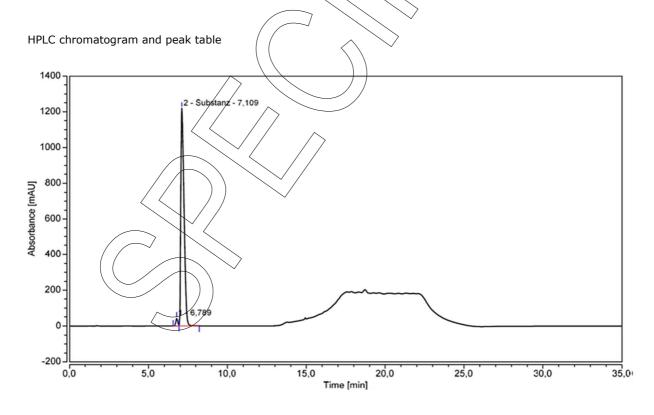


Assay

The assay of the reference material was assessed by following analyses.

Purity by High Performance Liquid Chromatography (HPLC)

HPLC conditions:	
Column	Discovery HS F5; 3 µm, 150 x 4.0 mm
Column temperature	40 °C
Detector DAD, 200/nm	
Injector Auto 2 µl; 0.2108 mg/ml in Water	
Flow rate	1.0 m/kmin
Phase A	Water, Q.1 % H ₃ PQ ₄
Phase B	Acetonitrile, 0.1 % H ₃ RO ₄
Gradient program	0-10 min A/B 99/1 10-15 min A/B to 60/40 15-20 min A/B 60/40 20-23 min A/B to 99/1 23-35 min A/B 99/1 (v/v)





Area percent report - sorted by signal				
Pk #	Retention time	Area	Area %	
1	6.789	6.065	2.05	
2	7.109	289.798	97.95	
Totals		295.863	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.





Volatile content

Water content		
Method	Karl Fischer titration	
Result (n = 3)	0.48 %; SD = 0.03 %	

Residual solvents		^	1			
Method	¹ H-NMR	/				
Result (n = 1)	No significant amounts of residual s			detected (< 0.0	5 %).

Final result

Assay "as is":

97.50 %

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

The calculation of the 100% method follows the formula:

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



Revision	Date	Reason for revision
00	08 Oct 2019	Release of the Certificate of Analysis – initial version

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

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