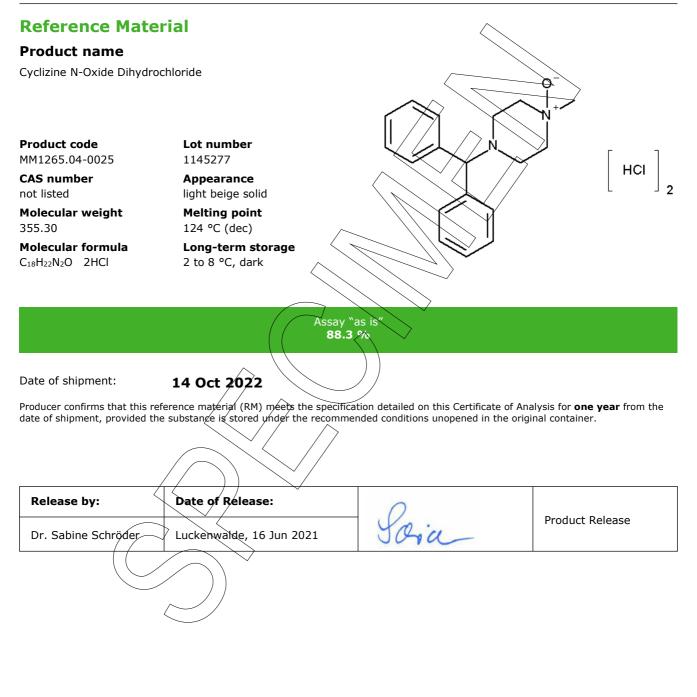


Certificate of Analysis



Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCiPACT $^{\text{TM}})$

Producer: LGC GmbH Louis-Pasteur-Str. 30 D-14943 Luckenwalde Germany www.lgcstandards.com Page 1/7



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)

Further content

Identity Assay Final result

Revision table



Identity

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





Method	Conditions	Result
MS	3.5 kV ESI+; capillary temperature: 269 °C Theoretical value: 283.18049	Structure confirmed
Apex MM1265.04-R	2AW-FA765803-172-0264-1 Scan: #295 RT: 3,07 min NL: 6,76E+007	FTMS + p.ESI Full ms [80.00-1000.00]
100-	283,18003	
90-		
80-		
70-		
60-		
50-		\searrow
40-		
30- 167,0854	43	
20-	284, 18343	
117,10214		
6002 171,992		m/z
80	200 300 400 500 600	700 800 900 1.000
/		





Assay

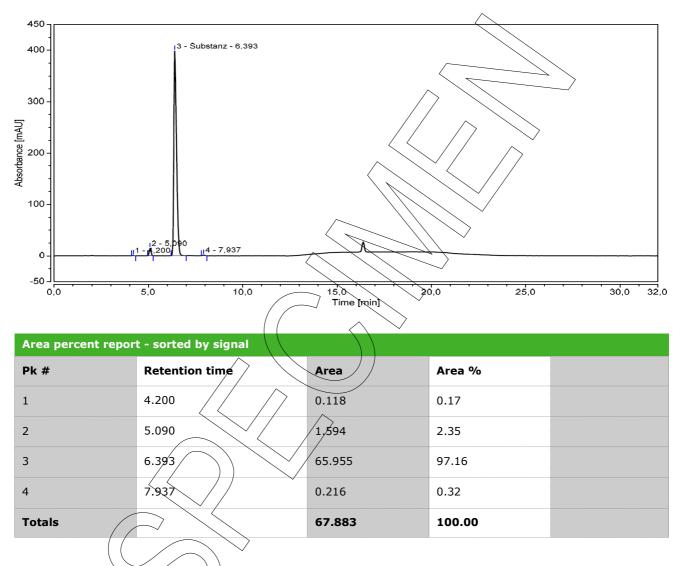
The assay of the reference material was assessed by following analyses.

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, 225 nm
Injector	Auto 2 μl; 0.266 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_3PO_4
Gradient program	0-9 min A/B 70/30
	9-12 min A/B to 20/80
	12-17 min A/B 20/80
	17-20 min A/B to 70/30
	20-32 min A/B 70/30 (v/v)



HPLC chromatogram and peak table



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)

97.16 %; SD < 0.01 %



Volatile content

Water content		~
Method	Karl Fischer titration	
Result (n = 3)	6.62 %; SD = 0.11 %	

Residual solvents		
Method	¹H-NMR	
Result (n = 1)	Sum: 2.49 %	
	2.49 % Ethanol	

Final result

Assay "as is":

88.31 %

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:

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

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



Revision table

Revision	Date	Reason for revision
00	16 Jun 2021	Release of the Certificate of Analysis - initial version

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