

# **Certificate of Analysis**

## **Reference Substance**

## Vinpocetine

Catalogue Number: Lot Number: Molecular Formula: Molecular Weight: CAS Number:	LGCFOR0406.00 36365 C <sub>22</sub> H <sub>26</sub> N <sub>2</sub> O <sub>2</sub> 350.45 [ 42971-09-5 ]	Long-term Storage: Appearance: Melting Point: Assay 'as is':	2 to 8 °C, dark white solid 151 °C 98.4 %		
Date of shipment: 2016-May-20					
	for two years from the e is stored under the re-				
Release Date:	2013-08-22	LGC GmbH			
		Dr. Sabine Schröde Product Release	ЭГ		
$\bigcirc$		Product Release			





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6 pages



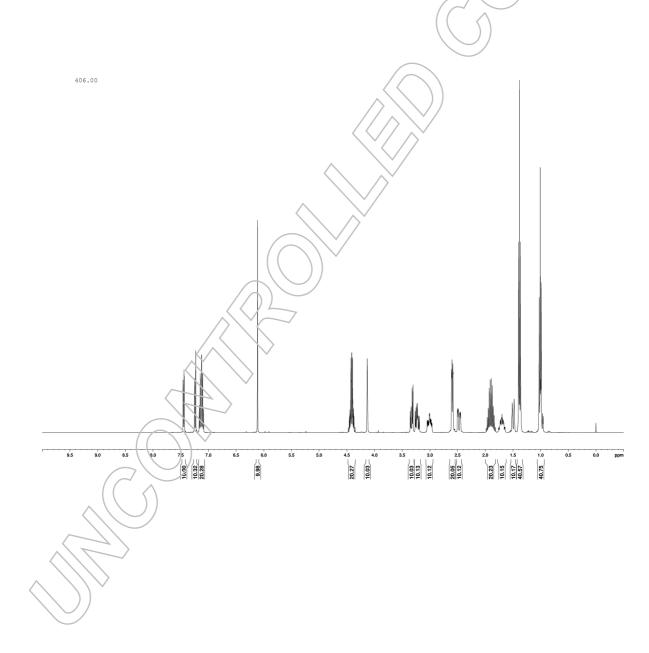
## I. Identity

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

#### Ia. <sup>1</sup>H-NMR Spectrum

Conditions: 400 MHz, CDCl<sub>3</sub>

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



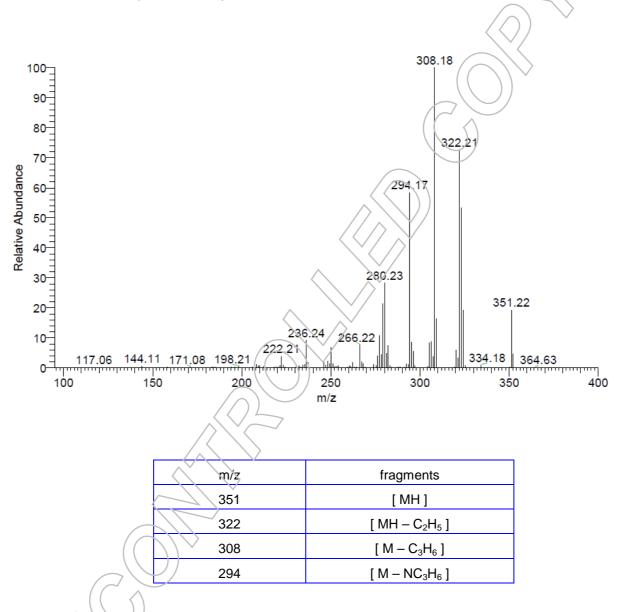


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#### Ib. Mass Spectrum

Method: 4.5 kV ESI; vaporization temperature: 200 °C, direct inlet



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



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#### **IR Spectrum** Ic. Method: Attenuated Total Reflection Fourier Transform Infrared (ATR-FTIR) Spectroscopy 100 Result of Peak Picking No. Position Intensity 1 2929.34 54.0531 2 1715.37 43.78 1604.48 1454.06 72.975 50.1124 80 1257.36 53.5982 1078.98 744.388 54.6228 57.5143 %Т 60 40 4000 3600 3400 3200 3000 2800 2600 2400 2200 2000 1800 1700 1600 1500 1400 1300 1200 1100 1000 900 800 650 Wavenumber [cm-1]

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

## II. Purity

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

#### **HPLC Conditions:**

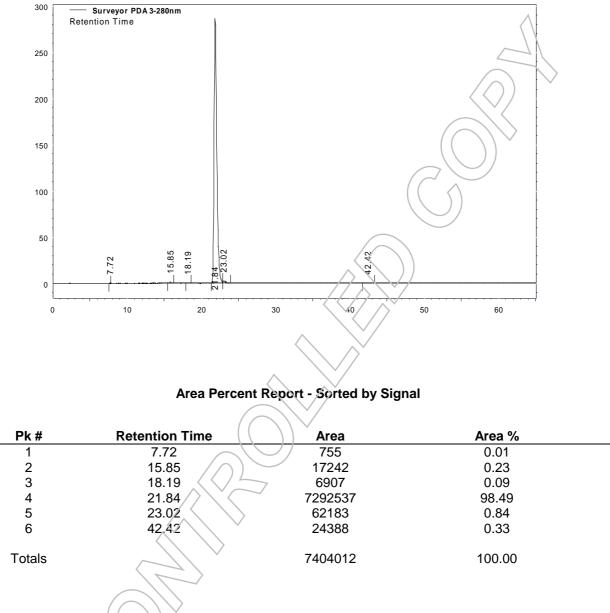
**Column:** Luna C 18 (2) 5 μm, 250 x 4.6 mm Conditions: 1.0 ml/min, 40 °C 15.4 g/l Ammonium acetate/ Acetonitrile 45/55 (v/v)

Detector:	Injector:
DAD	Auto
280 nm	7 µl; 0.14828 mg/ml in Eluent



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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	98.49 %
Number of results	n=15
Standard deviation	0.07 %



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## III. Water Content

Method: Karl Fischer titration

**Results:** 

Average	0.05 %
Number of results	n=3
Standard deviation	0.01 %

## **IV. Residual Solvents**

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

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

## V. Final Result

Total impurities (HPLC)	1.51 %
Water content	0.05 %
Residual solvents	n. d. (not detected)
Assay (100 % method) <sup>1</sup>	98.44 %

3.44 %

The assay is assessed to be 98.4 % 'as is'

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

<sup>1</sup> The calculation of the 100 % method follows the formula:

Assay (%) = (100 % - KF - RES)

Purity HPLC (%) 100 %

Water (KF) and Residual solvents (RES) are considered as absolute contributions, HPLC purity is considered as relative contribution.

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Excellence through measurement