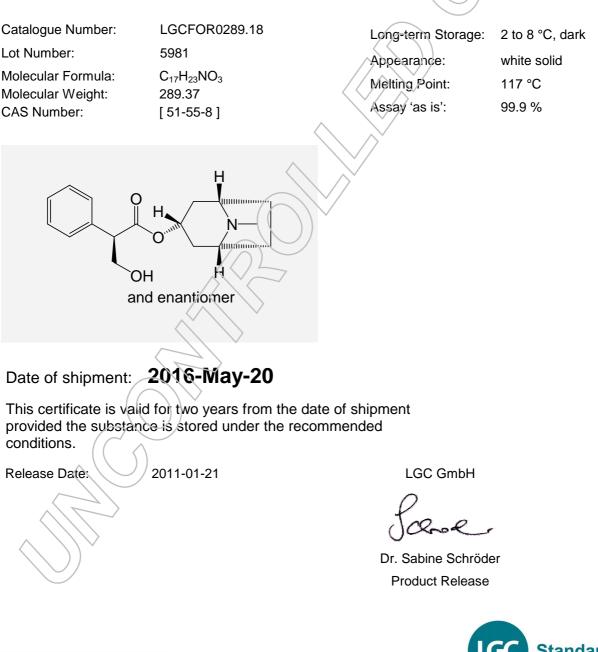


Certificate of Analysis

Reference Substance

Atropine





LGC Quality | ISO 9001:2008 DQS 102448 QM08

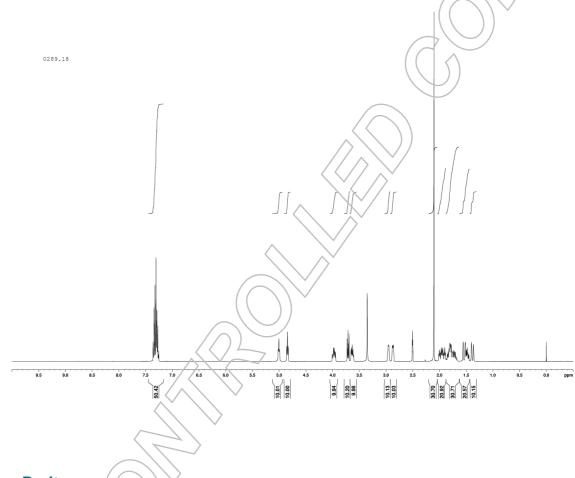
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4 pages



I. Identity

The identity of the reference substance was established by ¹H-NMR spectroscopy. The structure is confirmed with the signals of the spectrum and their interpretation. Conditions: 400 MHz, DMSO-d₆



II. Purity

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

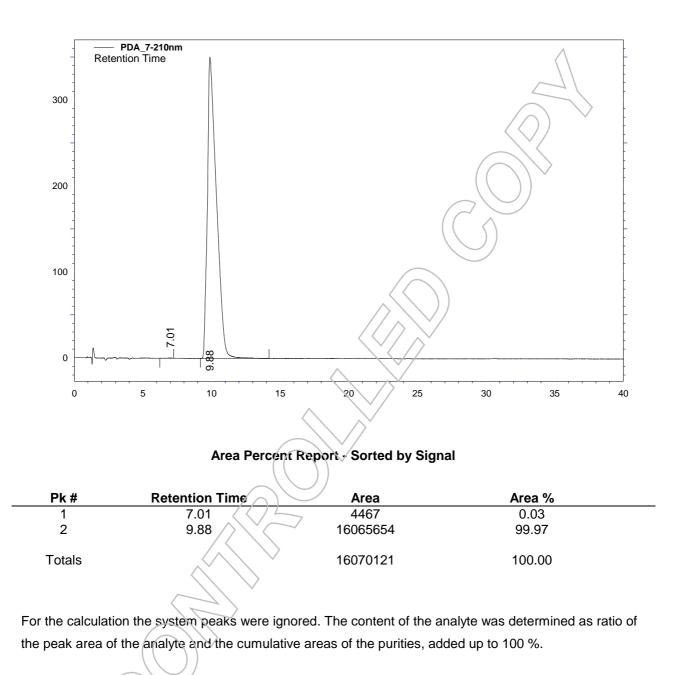
HPLC Conditions:			
Column	Conditions:	Detector:	Injector:
RP 60 Select B	1.0 ml/min, 40 °C	DAD	Auto
5 µm, 125 x 4 mm	Water/Acetonitrile 90/10 (v/v);	210 nm	15 μl; 0.2124 mg/ml in
	0.1 % H ₃ PO ₄		Water/Acetonitrile 50/50 (v/v)



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Results:Average99.96 %Number of resultsn=3Standard deviation0.01 %



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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: ¹H-NMR

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

V. Final Result

Total impurities (HPLC)	0.04 %
Water content	0.05 %
Residual solvents	n. d. (not detected)
Assay (100 % method) ¹	99.91 %

The assay is assessed to be 99.9 % 'as is'

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

¹ 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