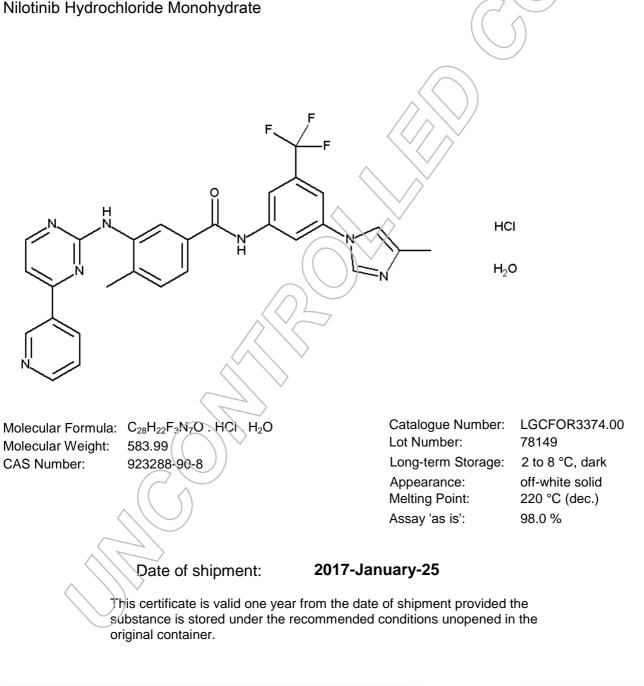


Certificate of Analysis

Reference Standard

Nilotinib Hydrochloride Monohydrate



ISO 9001:2008 DQS 102448 QM08

LGC Quality

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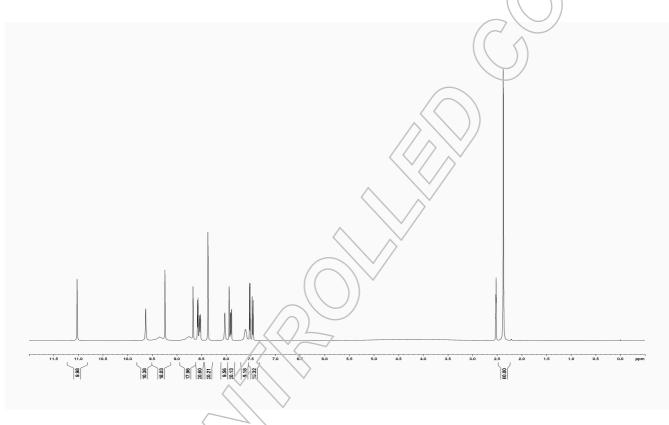


I. Identity

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

¹H-NMR Spectrum la.

Conditions: 400 MHz, DMSO-d₆



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



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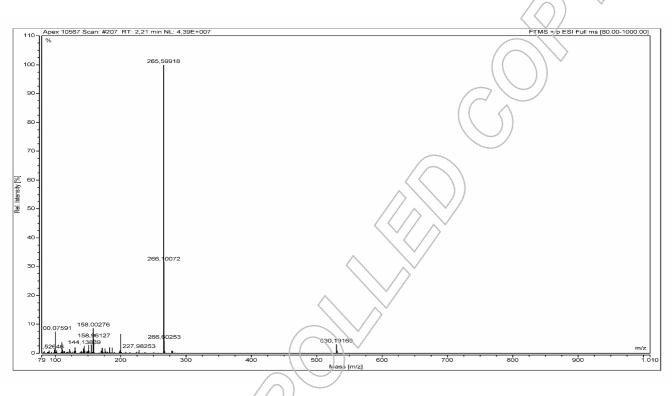
lot number 78149

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Ib. Mass Spectrum

Method: HRMS; 3.5 kV ESI+; capillary temperature: 269 °C



Theoretical value: 530.19107

The signal of the MS spectrum is consistent with the theoretical value and its interpretation is consistent with the structural formula.

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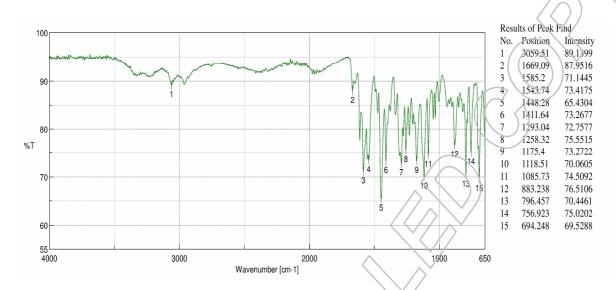
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Ic. IR Spectrum

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



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

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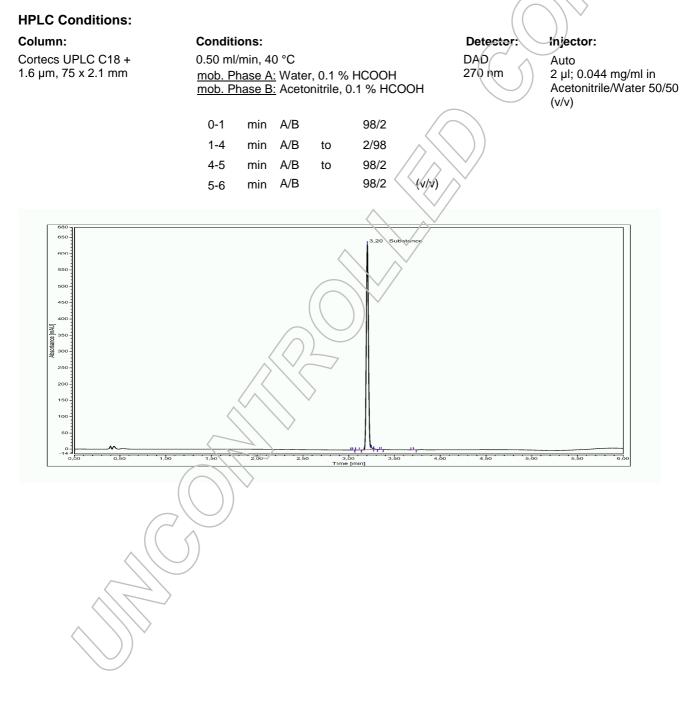
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II. Purity

IIa. High Performance Liquid Chromatography (HPLC)

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



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Cal[®]



Area Percent	Report - Sorte	d by Signal

Pk#	Retention Time	Area	Area %
1	3.043	0.0011	0.01
2	3.117	0.0034	0.02
3	3.203	16.9628	98.89
4	3.247	0.1340	0.78
5	3.273	0.0411	0.24
6	3.357	0.0064	0.04
7	3.707	0.0035	0.02
Totals		17.1523	100.00

For the calculation the system peaks were ignored. The content of the analyte was determined as the 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	0.02 %

IIb. Water Content

Method: Karl Fischer titration

Results:	
Average	4,00 %
Number of results	n=3
Theoretical value	3.08 %
Content of excessive water	0.92 %
Standard deviation	0.11 %
\frown	

IIc. Residual Solvents

Method: ¹H-NMR No significant amounts of residual solvents were detected (< 0.05 %).



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III. Final Result

Chromatographic purity (HPLC) Water content Residual solvents Assay (100 % method)¹

98.88 % 0.92 % No significant amounts of residual solvents were detected (< 0.05 %) 97.97 %

The assay is assessed to be 98.0 % 'as is'

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

Release Date:

Signed:

Luckenwalde, 2015-September-08

Dr. Sabine Schröder Product Release

¹ The calculation of the 100 % method follows the formula: Assay (%) = (100 % - volatile contents) * $\frac{Purity (\%)}{400\%}$

4ssay(%) = (100% - volatile contents) - 100%

Volatile contents are considered as absolute contributions, purity is considered as relative contribution

LGCFOR3374.00 lot number 78149

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