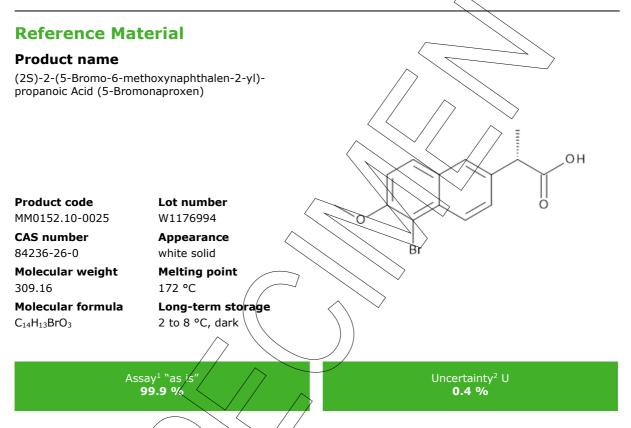




# **Certificate of Analysis**

Characterisation methods are accredited according to

**ISO 17025** 



Intended Use: Use for identification and quantification. The assay is verified by a second testing method.

Date of shipment: **Q8 Nov 2021** 

Producer confirms that this reference material (RM) meets the specification detailed on this Certificate of Analysis for **two years** from the date of shipment, provided the substance is stored under the recommended conditions unopened in the original container.

Release by:	Date of Release:	0	Product Release
Dr. Sabine Schröder	Luckenwalde, 18 Oct 2021	Jarol	Product Release

<sup>&</sup>lt;sup>1</sup> Calibration and verification were carried out using standards traceable to SI-units. The value is expressed on an "as is" basis.

Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCiPACT $^{TM}$ ) Test methods used for characterisation are accredited to ISO/IEC 17025 | DAkkS D-PL-14176-01-00

Producer: LGC GmbH Louis-Pasteur-Str. 30 D-14943 Luckenwalde Germany www.lgcstandards.com Page 1/9

<sup>&</sup>lt;sup>2</sup> The uncertainty "U" is the expanded uncertainty of the testing method for the assigned value estimated in accordance with the Guide to the Expression of Uncertainty in Measurement (GUM). It corresponds to a level of confidence of about 95%. Coverage factor k = 2.



#### **Product information**

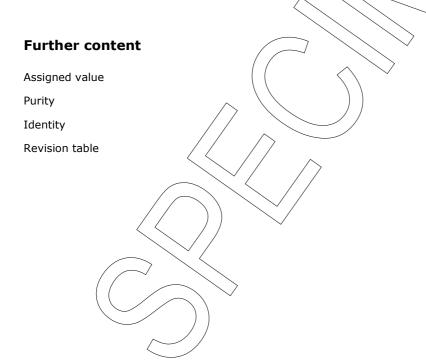
This RM is intended for laboratory use only and is not suitable for human or animal consumption.

This RM conforms to the characteristics of a primary standard as described in the ICH Guidelines. The values quoted in this Certificate of Analysis are the producer's best estimate of the true values within the stated uncertainties and based on the techniques described in this Certificate of Analysis. The characterisation of this material was undertaken in accordance with the requirements of ISO/IEC 17025. The identity is verified by data from international scientific literature.

#### Storage and handling

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

Reference Material quality is controlled by regularly performed quality control tests (retests).



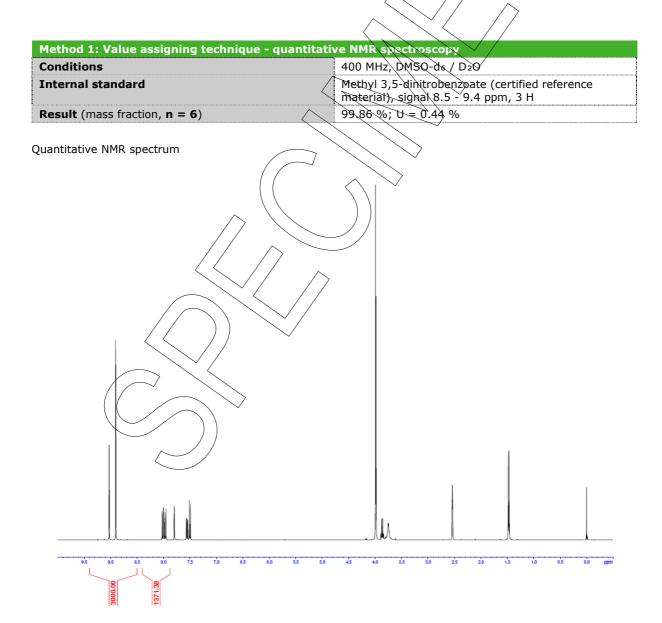


#### **Assigned value**

Assay "as is": 99.86 %; U = 0.44 %

The assay "as is" is assessed by quantitative NMR spectroscopy and is equivalent to the assay based on the not-anhydrous and not-dried substance. The assay is verified by 100% method (mass balance). The verified result lies inside our acceptance criteria, i.e. less than 1.0 % difference to assay assigning technique.

For quantitative applications, use the assay as a calculation value on the assay can be used for estimation/calculation of measurement uncertainty.





# Method 2: Value verifying technique - 100% method 100% method (mass balance) with chromatographic purity by HPLC Result 99.30 %

The calculation of the 100% method follows the formula:

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

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

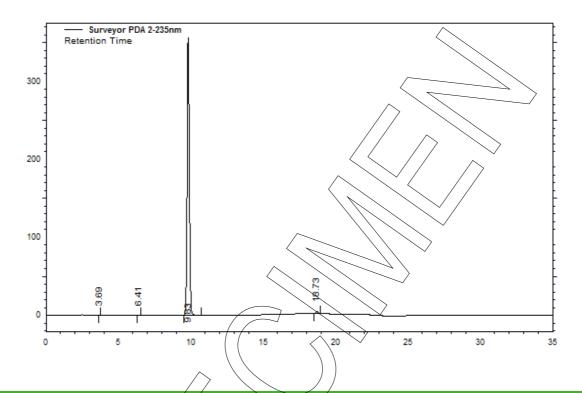
### **Purity**

Purity by High Performance Liquid Chromatography (HPLC)

HPLC conditions:	
Column	Pro C18 RS; 5 μm, 150 x 4.6 mm
Column temperature	40/°¢
Detector	DAD, 235 nm
Injector	Auto 2 μl; 0.032 mg/ml in Water/Acetonitrile 50/50 (v/v)
Flow rate	1.0 ml/min
Phase A	Water, 0.1 % H₃PO₄
Phase B	Acetonitrile, 0.1 % H₃PO₄
Gradient program	0-10 min A/B 55/45 10-15 min A/B to 30/70 15-20 min A/B to 55/45 20-35 min A/B 55/45 (v/v)



#### HPLC chromatogram and peak table



Area percent report - sorted by signal				
Pk #	Retention time	Area	Area %	
1	3.69	7/19	0.02	
2	6.41	1657	0.04	
3	9.83	3988610	99.31	
4	18.73	25312	0.63	
Totals		4016358	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)	No significant amounts of water were de	etected (< 0.05 %).*

<sup>\*</sup>not accredited testing method

Residual solvents			
Method	<sup>1</sup> H-NMR		$\triangle$
Result (n = 1)	No significant amounts of residual	Kşolv	vents were detected (< 0.05 %).*

<sup>\*</sup>not accredited testing method

#### **Inorganic residues**

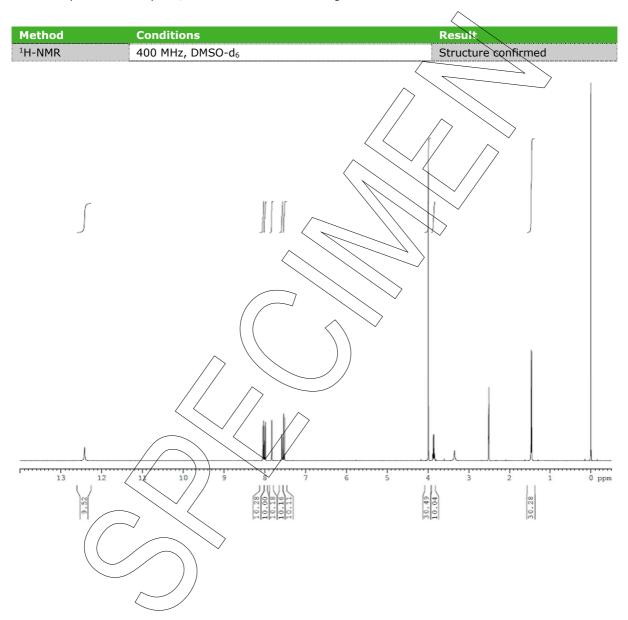
Method: Quantitative NMR spectroscopy

Inorganic residues can be excluded by quantitative NMR spectroscopy.



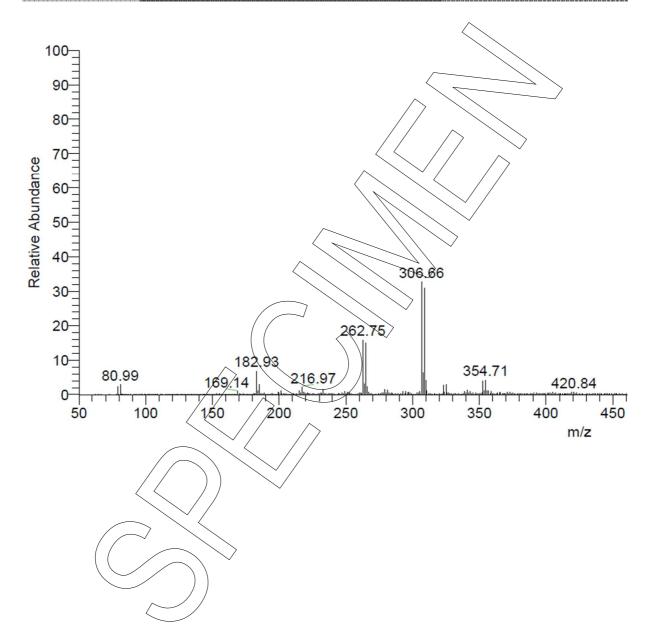
## **Identity**

The identity is assessed by ISO/IEC 17025 accredited testing methods.

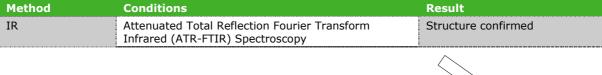


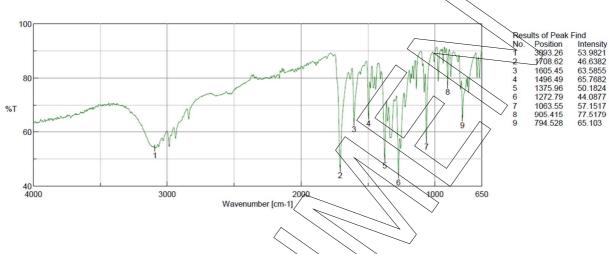


Method	Conditions	Result
MS	4.5 kV ESI; vaporization temperature: 200 °C	Structure confirmed





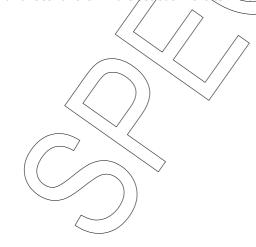




#### **Revision table**

Revision	Date	Reason for revision
00	18 Oct 2021	Release of the Certificate of Analysis – initial version

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



Page 9/9