

Certificate of Analysis

ISO 17034

Reference Mate	rial	<	
Product name Thymol			
Product code MM0364.00	Lot number G998469		ОН
CAS number	Appearance	$\sim // >$	
89-83-8	white solid		I
Molecular weight 150.22	Melting point 50 °C		
Molecular formula	Long-term storage	\searrow	
C ₁₀ H ₁₄ O	2 to 8 °C, dark	\sum	
Assay 99.	97 %		rtainty ² U . 6 %
Intended Use: Use for identia Due to the homogeneity studi	fication and quantification. The ass es, the minimum amount of sample	ay is verified by a second test to be used is 10 mg.	ing method.
Date of shipment:	02 Mar 2020		

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:	P	Dreduct Delegas
Dr. Sabine Schröder	Luckenwalde, 25 Feb 2020	Johnok	Product Release

¹ Calibration and verification were carried out using standards traceable to SI-units. The value is expressed on an "as is" basis.

² 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.

Organisation certified to ISO 9001 | DQS 102448 and GMP (EXCiPACTTM) RM Production accredited to ISO 17034 | DAkkS D-RM-14176-01-00 | Test methods used for characterisation are accredited to ISO/IEC 17025 | DAkkS D-PL-14176-01-00



Important 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 production of this RM was undertaken in accordance with the requirements of ISO 17034. 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.

Further content

Assigned value
Purity
Identity
Stability and homogeneity
Revision table

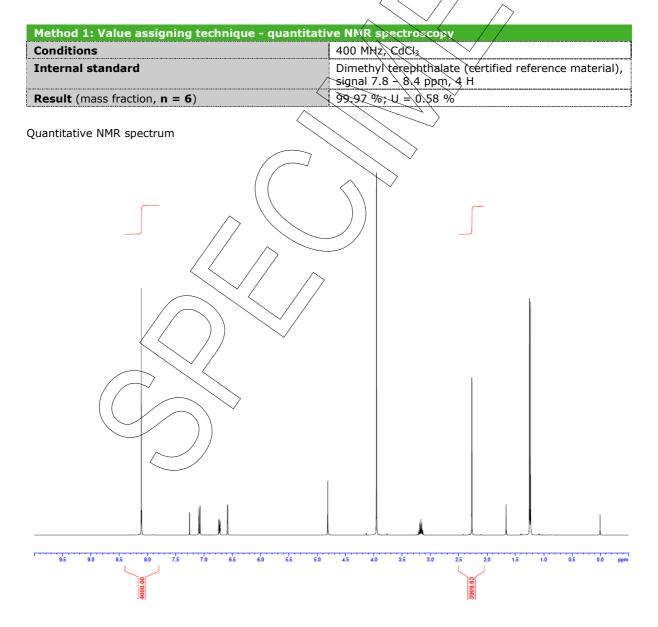


Assigned value

Assay "as is": 99.97 %; U = 0.58 %

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 "as is basis". The uncertainty of 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	99.94 %		

Purity (%) 100 %

The calculation of the 100% method follows the formula:

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

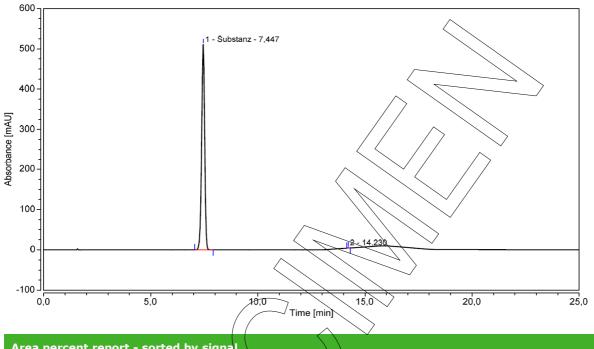
Purity

Purity by High Performance Liquid Chromatography (HPLC)

		litions:	HPLC conditio
	Hypersil Gold C18; 5 µm, 150 x 4.6 mm		Column
	40)°C	mperature	Column tempe
	DAD, 220 nm		Detector
ile 50/50	Auto 5 μl; 0.093 mg/ml in Water/Acetonitrile 5 (v/v)		Injector
	> 1.0 ml/min		Flow rate
	Water, 0.1 % H ₃ PO ₄		Phase A
	Acetonitrile, 0.1 % H₃PO₄		Phase B
	0-10 min A/B 55/45 10-13 min A/B to 20/80 13-15 min A/B to 55/45 15-25 min A/B 55/45 (v/v)	program	Gradient prog
	13-23 IIIIII A/B 33/43 (V/V)		



HPLC chromatogram and peak table



Area percent report - sorted by signal				
Pk #	Retention time	Area	Area %	
1	7.447	83.166	99.94	
2	14.233	0.053	0.06	
Totals		83.218	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.

Result (n = 6)

99.94 %; U = 0.19 %



Volatile content

Water content		
Method	Karl Fischer titration	\frown
Result (n = 3)	No significant amounts of residual	l solvents were detected (< 0.05 %).*

*not accredited testing method

Residual solvents			
Method	¹ H-NMR		
Result (n = 1)	No significant amounts of residua	l sol	lvents were detected (< 0.05 %).*

*not accredited testing method

Inorganic residues

Method: Sulphated ash^{*}, EP 8.7 (2.4.14)

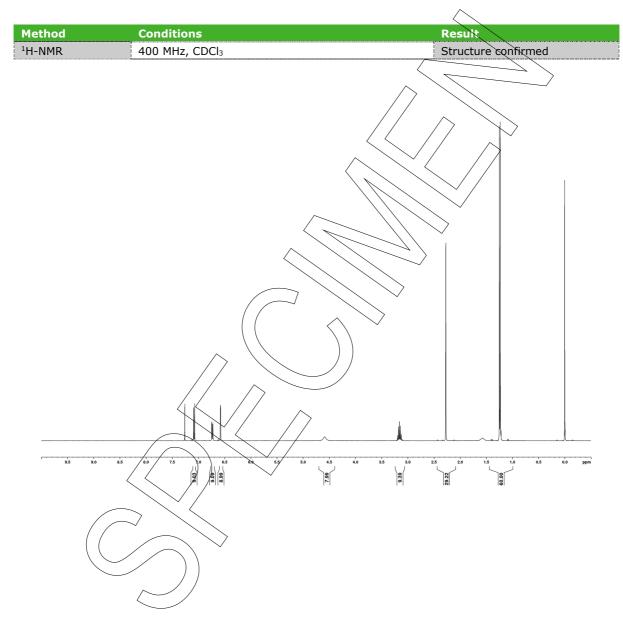
According to the available data, the presence of inorganic impurities in the reference material other than those detectable by sulphated ash is highly unlikely. Inorganic residues can be excluded by results of the sulphated ash. Therefore, no assay correction was performed for inorganic impurities.

*not accredited testing method



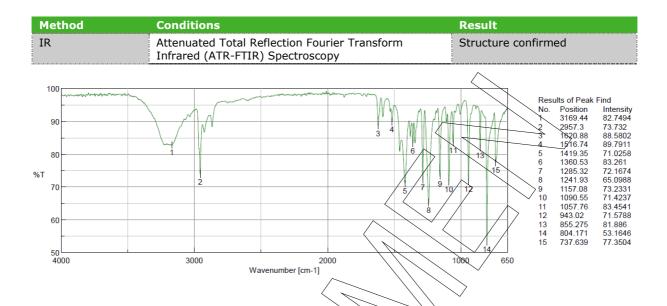
Identity

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









Stability and homogeneity

Accelerated stability studies indicate no significant instability. The given validity period is based on this data. This is backed up by additional stability testing and historical data over the range of several years.

RM quality is controlled by regularly performed quality control tests (re tests). Homogeneity assured by qualified process of preparation and verified by homogeneity testing.

Revision table

Revision	Date	Reason for revision
00	25 Feb 2020	Release of the Certificate of Analysis - initial version

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