

Certificate of Analysis

Characterisation methods are accredited according to

Reference Material Product name Methenolone Enantate Product code Lot number MM3721.00 W1000940 **CAS** number Appearance 303-42-4 white solid Molecular weight Melting point (DSC) 414.62 71 °C Long-term storage Molecular formula C27H42O3 2 to 8 °C, dark Assay¹ "as is' Uncertainty² U 99.5 % 0.3 %

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

Date of shipment:

02/Sep 2019/

Producer confirms that this reference material (RM) meets the specification detailed on this Certificate of Analysis for **one year** 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	
Dr. Sabine Schröder Luckenwalde, 06 Aug 2019	Joia	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 uncertaint, "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 (EXCiPACT) 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 1/8



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).

Further content

Assigned value Purity Identity Revision table



Assigned value

Assay "as is": 99.51 %; U = 0.29 %

The assay "as is" is assessed by carbon titration of elemental analysis 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 1: Value assigning technique - carbon titration of elemental analysis		
Method	percentage carbon found in relation to percentage carbon as calculated for molecular formula	
Results (mass fraction, n = 3)	99.51 %; U = 0.29 %	
Method 2: Value verifying technique - 100% n	nethod	
100% method (mass balance) with chromatographic purity by GC		
Result	99.62 %	

Purity (%)

The calculation of the 100% method follows the formula:

Assay (%) = (100 % - volatile contents (%)) *

100 %

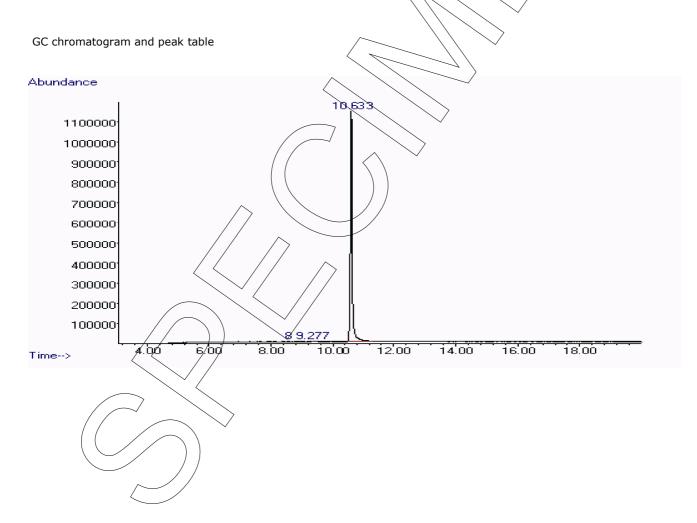
Volatile contents are considered as absolute contributions and purity is considered as relative contribution. Inorganic residues are excluded by additional tests.



Purity

Purity by Gas Chromatography (GC)

GC Conditions:		
Column	HP-5MS 30 m x 0.25 mm x 0.25 μm	
Detector	EI, 70 eV; 35 tø 550 amu; 280 °C	
Injector	Split 20:1, 220 °C	
Flow rate	Helium 1.50 ml/min	
Oven program	Initial Temp.: 220 °C for 3 min Heating Rate: 40 °C/min Final Temp.: 300 °C for 15 min	





Area percent repor	rt - sorted by signal		
Pk #	Retention time	Area	Area %
1	8.900	38374	0.09
2	9.277	109297	0.25
3	10.633	44429846	99.67
Totals		44577517	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 %. Air peaks were ignored in calculation.

Result (n = 3)	99.68 %; U = 0.19 %
Volatile content	
Water content	
Method	Karl Fischer titration
Result (n = 3)	$0.06 \%^{*}; SD = 0.01 \%$
*not accredited testing method	
Residual solvents	\sim $<$ //
Method	1H-NMR
Result (n = 1)	No significant/amounts of residual solvents were detected (< 0.05 %).*
*not accredited testing method	
Inorganic residues Method: Elementary analysis	

Inorganic residues can be excluded by elementary analysis (CHN).



Identity

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

