

## Cannabidiol (CRM)

Certified Reference Material

 Item No.:
 ISO60156

 Batch No.:
 0567962

 CAS Registry No.:
 13956-29-1

 Molecular Formula:
 C<sub>21</sub>H<sub>30</sub>O<sub>2</sub>

 Formula Weight:
 314.50 amu

Expiry Date: 13AUG2023 (valid from date of certification)
Supplied as: A 1 mg/ml (nominal) solution in methand

Volume per Ampule: Not less than 1 ml. Ampules are over aled.

Storage: Unopened at -20°C.

Safety: Refer to Safety Data Sheet

Intended Use: For analytical testing our poses only, not intended for human or animal use.

Instructions for Use: This product is designated for one-time use and should be used immediately after opening.

It is advised that labout ries warm the vial to room temperature prior to opening and use

measured volur es.

### Certified Concents tion 1.000 mg/ml ± 0.021 mg/ml

Concentration is calculated based foroduct mass, solution mass, corrected purity, and density at 20°C. It is traceable to SI units through an unbroken chain of measurements. Unsertainty of concentration is expressed as an expanded uncertainty in accordance with ISO standards for Testing Laboratories and Reference Material Producers at the approximate 95% confidence interval using a coverage factor of k=2 and incorporates uncertainties from the corrected purity, solution preparation, homogeneity, and long- and short-term stability. Concentration was verified by comparison to an independently prepared calibration standard.

### Corrected Purity · 99.70% ± 0.56%

Corrected purity is determined as follows: Corrected Purity = [(100 - % LOD - % ROI)\*Chromatographic Purity/100] or [(100 - % KF - % RS - % ROI)\*Chromatographic Purity/100]. All measurement uncertainties are expressed as expanded uncertainties in accordance with ISO standards for Testing Laboratories and Reference Material Producers at the approximate 95% confidence interval using an appropriate coverage factor. Where applicable, optical rotation, chiral purity, and/or isotopic purity testing are performed to support the identification of the reference material, therefore the uncertainty is considered null.

Approval: Title: ISO Quality Manager Certification Date: 13AUG2019

Cayman Chemical certifies that this standard meets the specifications stated in this certificate and warrants this product to meet the stated acceptance criteria through the expiration date when stored unopened as recommended.



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### **CRM Assay**

Method Parameters		
Cayman Method	TST SD151	
Column	4.6 x150 mm, 5 μm Kinetex Biphenyl	
Mobile Phase	A: 0.1% Trifluoroacetic Acid in Water B: Acetonitrile	
Gradient	Time (min) %B 0-12 50-90% 12-17 90% 17.1-22 50%	
Flow Rate	1 ml/min	
Column Temp	30°C	
Wavelength	UV monitored at 232 nm	

### Homogeneity

A minimum sample size of 2.0  $\mu$ g was used to determine homography. At mographic was determined by HPLC using ampules selected from a random sampling plan from early, mixale, and late  $\ddot{\nu}$  positions.

%RSD	Acceptanc Prite.
1.20%	≤3%

The recommended minimum quantity for use is 2.0 µg. Quant es below this have not been evaluated.

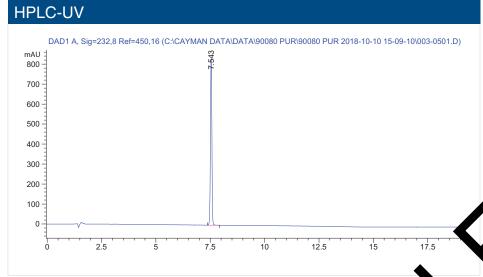
## Neat Material Quality Information (Item To 95-80, Batch No.: 0542338)

Qualifier	M thod	Result
Chromatographic Purity, HPLC	Cayman Method TST SD151	>99.90% ± 0.18%
Identity, LC-MS	Cayman Method TST SD13, +ESI	315.2 amu
Identity, GC-MS	Cayman Method TST SD12	Conforms
Identity, FTIR	USP<851> (diamond ATR)	Conforms
% LOD	Cayman Method TST SD24	<0.10% ± 0.48%
% ROI	Cayman Method TST SD06	<0.10% ± 0.21%
Identity, NMR	<sup>1</sup> H NMR	Conforms

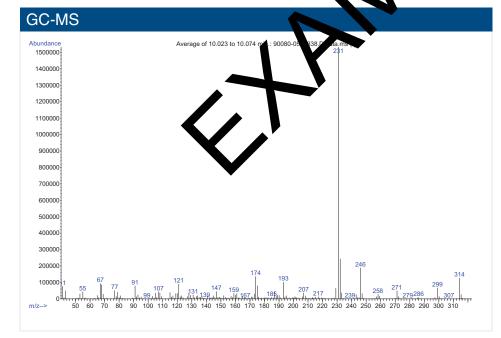
NMR and optical rotation (if applicable) are provided as supplemental information but are not within scope of ISO accreditation. Property values are traceable to SI units through an unbroken chain of measurements.



### Supplemental Data (Neat Material)



Agilent 1100/1200 Series	
4.6 x 150 mm, 5 µm Kinetex Biphenyl	
A: 0.1% Trifluoroacetic Acid in Water B: Acetonitrile	
Time (min) %B 0-12 50-90% 12-17 90% 17.1-22 50%	
1 ml/min	
30°C	
UV monitored at 232 nm	



Conditions		
Instrument	Agilent GC MSD	
Column	30 m x 0.32 mm, 0.5 μm Rtx-5MS	
Carrier Gas	Не	
Flow Rate	2 ml/min	
Inlet Temp	300°C	
Split Ratio	15:1	
Oven Program	50°C hold for 1 min, ramp to 300°C at 30°C per min, hold at 300°C to 15 minutes	
Transfer Line Temp	300°C	
Voltage	70ev EI MS	
Scan Range	40-600 m/z	
Tune File	stune	

Apex spectrum – background (1 min window in front of peak)

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Thermo Nicolet iS10 FTIR /

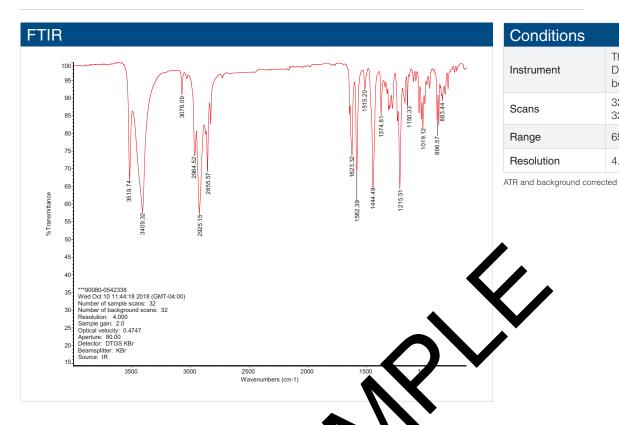
Diamond SmartATR (single

32 background scans

650-4,000 cm-1

bounce)
32 scans /

4.000



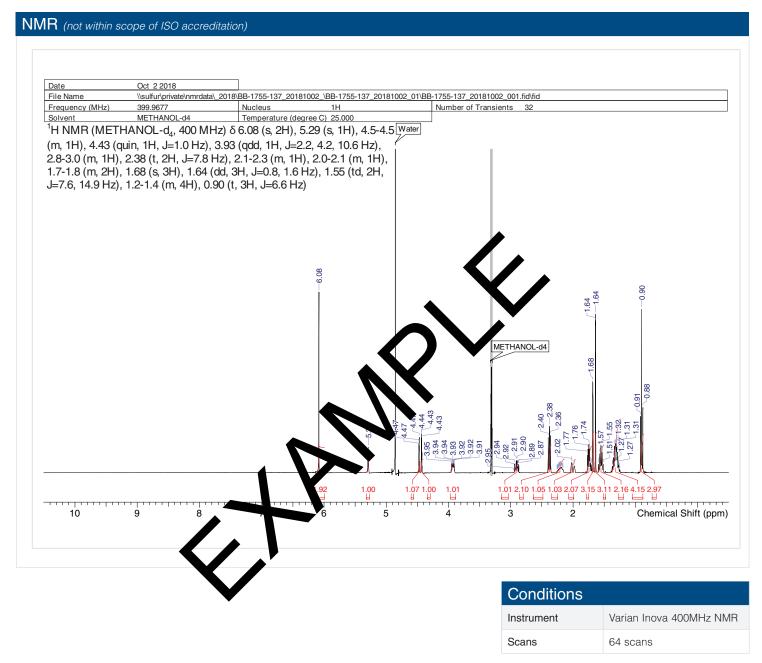
ESI-MS	
*MSD1 SPC, 6me=	273 of CXCAYMANDATAIFIAICAY030.D
80 -	
60 -	
40 -	
20 -	3372
0-	200 300 400 500 600 700 900 900 1000 m/z

Conditions		
Instrument	Agilent 1100 HPLC/MSD	
Mobile Phase	50:50:0.1 methanol/water/acetic acid	
Flow Rate	0.5 ml/min	
Ionization Mode	+ESI	
Mass Range	100-1,000 m/z	
Nebulizer	60 psi	
Desolvation Gas	13 L/min	
Desolvation Temp	350°C	
Electrospray Voltage	4kV	

MS collected across peak width at half height

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### Stability

The effect of the components of stability on the combined standard uncertainty of the CRM property value are considered negligible unless indicated in stability studies.

### Short-Term Stability

A decrease in property value was observed at 60°C during the two-week stability study. No decrease was observed at ambient temperature during the study. This data supports cold shipment of this product.

### Long-Term Stability

Long-term stability data predicts four years stability at the -20°C storage temperature. Long-term stability studies are ongoing and the Certificate of Analysis will be updated upon study completion.



### **Revision History**

Revision No.	Date	Reason for Revision
01	13AUG2019	Initial version

#### Disclaimers

#### Material Safety Data

This material should be considered hazardous until information to the contrary becomes available. Do not ingest, swallow, or inhale. Do not get in eyes, on skin, or on clothing. Wash thoroughly after handling. This information contains some but not all of the information required for the safe and proper use of this material. Before use, review the complete Safety Data Sheet, which has been sent *via* email to your institution.

#### Warranty and Limitation of Remedy

Cayman Chemical Company makes no warranty or guarantee of any kind, whether written or oral, pressed or implied, including without limitation, any warranty of fitness for a particular purpose, suitability and merchantability, which extends beyong the description of the chemicals hereof. Cayman warrants only to the original customer that the material will meet our specifications at the time of Selivary.

Cayman will carry out its delivery obligations with due care and skill. Thus, in no event will Cayman he and obligation or liability, whether in tort (including negligence) or in contract, for any direct, incidental or consequently, lamages, every Cayman is informed about their possible existence.

This limitation of liability does not apply in the case of intentional acts or neg ence of Zayman wis directors or its employees.

Buyer's exclusive remedy and Cayman's sole liability hereunder shall be liked to a sfund of the purchase price, or at Cayman's option, the replacement, at no cost to Buyer, of all material that does not meet our specification.

Said refund or replacement is conditioned on Buyer giving written on to Cryman within thirty (30) days after arrival of the material at its destination. Failure of Buyer to give said notice within thirty (30) days shall constitute a water of Buyer of all claims hereunder with respect to said material.

For further details, please refer to our Warranty and Limitations of speedy located on our website and in our catalog.

This Certificate shall not be reproduced except in full, without written approval from the Cayman Chemical ISO Quality Manager.

ISO CRT SD02 v 4.2

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