

Dronabinol DAC Therismos Kit 1000mg

Dronabinol Kit THC Pure Extract (95%) in 2.25 mL ITC syringe – 1.000mg **Document type:**

Scope:

CoA Number:

F.6010403.001-P.01

04/2025 03/2026

Certificate Of Analysis

(CoA)
Quality Control (QC)

18-25-0158-01

ES-EXTK-001_03

Batch number:
Packaging date:
Retest date:
Specification
reference:

Especificación / Specification Resultado/ Result **Producto / Product** Jeringas de vidrio de borosilicato tipo I (1 mL o 2,25 mL) con tapa de punta integrada y un émbolo con Jeringa tipo I (acondicionamiento primario) tapón. Cumple Complies Borosilicate glass type I syringes (1 mL or 2.25 mL) Type I syringes (primary packaging) with integrated tip cap and a plunger rod with a stopper. Caja de cartón blanco Cartón reciclado de doble capa, blanco. Cumple 28,5 x 75,5 x 30 mm (A x L x A) (acondicionamiento secundario) Complies Double-layer recycled cardboard, white White cartón case 28.5 x 75.5 x 30 mm (W x L x H) (secondary packaging) Cumple **Tube Eppendorf** 1.5 mL Complies **Eppendorf tube** Cumple Aguja hipodérmica 09 x 40 mm (20G) Complies Hypodermic needle Prueba simple de screen de drogas (orina) Cumple Test identificación THC Complies Test ID THC Drug Screen single test (urline) Caja de cartón. 78 x 129 x 41 mm (A x L x A) Cumple Caja kit Complies Kit box Cardboard box 78 x 129 x 41 mm (W x L x H)

Function	Name/Position	Conclusion	Signature/Date
Prepared by:	Alvaro Pascual Quality Control Technician		ANR 12-02-26
Reviewed by:	Joel Prieto Quality Control Manager	COMPLIES	12-02-52 12-02-52
Reviewed by:	Soledad Pérez Quality Assurance Technician	Conplies	14/04/25
Approved by:	Maria de Gracia González Quality Assurance Manager	Complies	12/07/2025





Extracto de THC Puro (95%) en jeringa ITC de 2.25 mL - 1.000mg

THC Pure Extract (95%) in 2.25 mL ITC syringe – 1000mg

Document type: Certificate Of Analysis

(CoA)

Scope: Quality Control (QC)
CoA Number: 18-25-0155-01

Batch number: F.6010403.001-P

Cas nº: 1972-08-3
Manufacturing date: 03/2025
Packaging date: 04/2025
Retest date: 03/2026

Specification ES-EXT-003_4.0 reference:

Ensayo / Test	Método / Method	Especificación / Specification	Resultado/ Result	
	THE THE WEST			
Apariencia Appearance	DAC 2012/2 D 100 Inspección visual DAC 2012/2 D 100 Visual inspection	Líquido de color casi incoloro a amarillo claro, ocasionalmente aceitoso de color púrpura pálido o masa resinosa en la superficie. Almost colourless to light yellow in colour, occasionally pale purple oily liquid or resinous mass on the surface	Conforme Complies	
	In house	Absorbancia a 405 nm < 0,40 uA Absorbance at 405 nm < 0.40 uA	Conforme Complies	
dentificación THC: IR dentificación THC: HPLC dentification THC: IR dentification THC: HPLC	DAC 2012/2 D 100 Farmacopea Europea 2.2.24 Farmacopea Europea 2.2.29 DAC 2012/2 D 100 European Pharmacopoeia 2.2.24 European Pharmacopoeia 2.2.29	Positivo Positive	Conforme Complies Conforme Complies	
Solubilidad Solubility	DAC 2012/2 D 100 Inspección visual DAC 2012/2 D 100 Visual inspection	Prácticamente insoluble en agua; Parcialmente soluble en Etanol 96%, Parcialmente soluble en Glicerol 85%; Soluble en aceites. Practically insoluble in wáter; slightly soluble in Ethanol 96% and Glycerol 85%; Fatty oils soluble	Conforme Complies	
Pureza cromatográfica Cromatography Purity	DAC 2012/2 D 100 Farmacopea Europea 2.2.29 DAC 2012/2 D 100 European Pharmacopoeia 2.2.29	El dronabinol contiene un mínimo de 95,0 y un máximo de 100,0 por ciento (6aR, 10aR)-6a,7,8,10a-tetrahidro-6,6,9-trimetil-3-pentil-6H-dibenzo[b,d]pirano- 1-ol (Δ9-tetrahidrocannabinol) Dronabinol contains a minimum of 95.0 and a maximum of 100.0 percent (6aR, 10aR)-6a,7,8,10a-tetrahydro-6,6,9-trimethyl-3-pentyl-6H-dibenzo[b,d]pyran-1-ol (Δ9-Tetrahydrocannabinol)	98.1%	
		2 ((221)	1 - 25 T W/V	
		Impureza D área corregida (CBN) < 1,0% Impurity D corrected área (CBN) < 1.0%	Imp. D: 0.31 %	
Sustancias Relacionadas	DAC 2012/2 D 100 Farmacopea Europea 2.2.29 DAC 2012/2 D 100 European Pharmacopoeia 2.2.29	Impureza E (Δ ⁸ -THC) < 2,0% Impurity E (Δ ⁸ -THC) < 2.0%	Imp. E: 0.84 %	
Relative Substances		Otras impurezas < 1,0% Other impurities < 1.0%	RRT-0.61: 0.25% RRT-0.78: 0.23% RRT-0.86: 0.18% RRT-1.35: 0.08% Otras < 0.05%	



Extracto de THC Puro (95%) en jeringa ITC de 2.25 mL - 1.000mg

THC Pure Extract (95%) in 2.25 mL ITC syringe – 1000mg

Document type:

Certificate Of Analysis

(CoA)

Scope: CoA Number:

Quality Control (QC) 18-25-0155-01

Batch number:

F.6010403.001-P

Cas nº:

1972-08-3

Manufacturing date:

03/2025

Packaging date:

04/2025

Retest date: Specification

03/2026

reference:

ES-EXT-003_4.0

		Impurezas totales excepto impureza E < 3,0% Total Impurities except impurity E < 3.0%	1.05%
		2- Propanol ≤ 500 ppm (0,05%) 2- Propanol ≤ 500 ppm (0.05%)	No detectado Not detected
Disolventes Residuales	Farmacopea Europea 2.9.10 Farmacopea Europea 2.9.11	Heptano ≤ 5000 ppm (0,5%) Heptane ≤ 5000 ppm (0.5%)	No detectado Not detected
Residual Solvents	European Pharmacopoeia 2.9.10 European Pharmacopoeia 2.9.11	Etanol ≤ 5000 ppm (0,5%) Ethanol ≤ 5000 ppm (0.5%)	No detectado Not detected
		Metanol ≤ 3000 ppm (0.3%) Metanol ≤ 3000 ppm (0.3%) Metanol ≤ 3000 ppm (0.3%)	Not detected No detectado Not detected
TAMC (Aerobios totales) TAMC (Total Aerobic Microbial Count)		≤ 10 ⁴ UFC/g ≤ 10 ⁴ UFC/g	610 UFC/g
TYMC (Hongos y levaduras) TYMC (Total Combined Yeast and Moulds Count)		≤ 10² UFC/g ≤ 10² UFC/g	10 UFC/g
Enterobacterias Bile-tolerant Gram (-) bacteria	Farmacopea Europea, 5.1.8 y 5.1.4 European Pharmacopoeia 5.1.8 and 5.1.4	≤ 10 ² UFC/g ≤ 10 ² UFC/g	> 10 UFC/g y < 10 ² UFC/g
Escherichia coli		Ausencia/g Absence/g	Absence/g
Staphyloccoccus aureus		Ausencia/g Absence/g	Absence/g
Pseudomonas aeruginosa		Ausencia/g Absence/g	Absence/g
Salmonella spp.		Ausencia/25g Absence/25g	Absence/25g
Parámetros de acondicionado Packaging parameters Caída de la presión Pressure decay	USP 1207.2	t = 1h, P = 200 mbar	Conforme Complies
Rango de llenado Filling range	In-house	1,0620 g (-3% + 4%) (1,0301 - 1,1044 g)	Conforme Complies

CoAs: F.6010403.001-P

Function	Name/Position	Conclusion	Signature/Date
Prepared by:	Juan Francisco Méndez		
riepaieu by.	Quality Control Technician		1. SEWR 16.07.25
QC Reviewed by:	Joel Prieto	/	
	Quality Control Manager	COMPLIES	16-07-25
QA Reviewed by:	Soledad Pérez		11-
QA Neviewed by:	Quality Assurance Technician	CCRPLIES	- 14/0×/25
Approved by:	Maria de Gracia González	The same of the same	An NICHIC
	Quality Assurance Manager	Complies	12/07/2025



CERTIFICADO DE CONFORMIDAD CERTIFICATE OF COMPLIANCE

Scope: Quality Assurance (QA)

CoC Number:

E25-010

PRODUCTO/PRODUCT	LOTE/BATCH		UND / UND
DRONABINOL KIT – 2.25 mL ITC syringe - 1000 mg	F.6010403.001-P.01		643
FABRICANTE/MANUFACTURER	ENVASADO POR/ PACKAGED BY		LABORATORIO CONTROL CALIDAD/ QUALITY CONTROL LABORATORY
ALCALIBER S.A.	ALLOGA		ALCALIBER S.A.
CLIENTE/CUSTOMER			PAÍS/COUNTRY
Therismos GMBH			Germany

Fecha De Fabricación/Manufacturing Date: Ver certificado de análisis adjunto/See attached certificate of analysis

TO WHOM IT MAY CONCERN

On behalf of Linneo Health S.L., declare:

- This batch of product has been manufactured, including packaging/labelling quality control analysis in full compliance with the GMP Part I / Part II (cross the one which does not apply) requirements of the local Regulatory Authority and with the specification of the importing country or product specification.
- 2. The batch processing and analysis records were reviewed and found to be in compliance with GMP.
- 3. All deviations and OOS have been evaluated and approved based on fixed internal procedures
 - a. No release relevant deviation or OOS.
 - b. Additional quality relevant information for the mentioned batch is enclosed

I hereby certify that the above information is authentic and accurate.

Approved by:

Firma / Signature:

Fecha aprobación/Approval date

18th of July 2025

María de Gracia González QA Manager & QP