

Internal Code	NAME		Batch no.	
FIN000237	Remexian 33/1 VCC GWA 250g		2500861	
Manufacture Date	Expiry date	Analysis Date	Analytical Protocol	Specification
14/03/2025	09/2025	24/03/2025	PRT-0000515-1.0	SPE-0001333-ver.1.0

Tests	Methods	Specification	Results
<b>1. Identification</b>			
1.1. Method A: Macroscopy Examination	EP monograph 07/2024:3028	Complies with requirements of monograph	Complies
1.2. Method B: Microscopy Examination	EP monograph 07/2024:3028; Ph.Eur.2.8.23	Complies with requirements of monograph	Complies
1.3. Method C: HPTLC	EP monograph 07/2024:3028; Ph.Eur.2.8.25	Complies with requirements of monograph	Complies
<b>2. Foreign Matter</b>			
	EP monograph 07/2024:3028; Ph.Eur.2.8.2	≤ 2% w/w Cannabis flower does not contain any seeds and the whole herbal drug does not contain any leaves more than 1.0 cm in length	0% Cannabis flower does not contain any seeds and the whole herbal drug does not contain any leaves more than 1.0 cm in length
<b>3. Loss on Drying<sup>a)</sup></b>	EP monograph 07/2024:3028; Ph.Eur.2.2.32	≤ 12.0% w/w	6.8%
<b>4. Assay<sup>a)</sup></b>			
4.1. Total THC	EP monograph 07/2024:3028; Ph.Eur.2.2.29	29.7% - 36.3% w/w	31.5% (31.50%)
4.2. Total CBD		≤ 1.0% w/w	<LOQ (0.2%)
<b>5. Total CBN<sup>a)</sup></b>	EP monograph 07/2024:3028; Ph.Eur.2.2.29	≤ 1.0% w/w	0.1% (0.08%)
<b>6. Heavy Metals<sup>b)</sup></b>			
6.1. Cd		≤ 0.3 ppm	0.0 ppm (0.012 ppm)
6.2. Pb	EP monograph 07/2024:3028; Ph.Eur.2.4.27	≤ 0.5 ppm	< LOQ (0.020 ppm)
6.3. Hg		≤ 0.1 ppm	< LOQ (0.050 ppm)
6.4. As		≤ 0.2 ppm	0.1 ppm (0.096 ppm)

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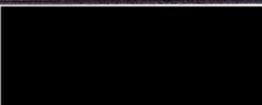
Tests	Methods	Specification	Results
<b>7. Microbiology</b>			
7.1. Total Aerobic Microbial Count (TAMC)	Ph.Eur. (5.1.8C, 2.6.12)	$\leq 10^5$ CFU/g (max: 500 000 CFU/g)	$2.0 \times 10^2$ CFU/g
7.2. Total Yeast and Mold Count (TYMC)		$\leq 10^4$ CFU/g (max: 50 000 CFU/g)	$4.5 \times 10^3$ CFU/g
7.3. Bile-tolerant gram-negative bacteria		$\leq 10^4$ CFU/g	<10 CFU/g
7.4. Escherichia coli		Absence in 1 g	Absent/g
7.5. Salmonella	Ph.Eur. (5.1.8C, 2.6.31)	Absence in 25 g	Absent/25g
7.6. Pseudomonas aeruginosa		Absence in 1 g	Absent/g
7.7 Staphylococcus aureus <sup>c)</sup>		Absence in 1 g	Absent/g
<b>8. Aflatoxins<sup>b)</sup></b>			
8.1. B1	Ph. Eur. 2.8.18	$\leq 2$ µg/kg	< LOD (0.25 µg/kg)
8.2. B1 + B2 + G1 + G2		$\leq 4$ µg/kg	< LOQ (0.5 µg/kg)
9. Ochratoxin A <sup>b)</sup>	Ph.Eur. 2.8.22	$\leq 20$ µg/kg	< LOD (2 µg/kg)
10. Pesticides <sup>b)</sup>	Ph.Eur. 2.8.13	Must comply with requirements of monograph	Complies

Remarks:

- a) From Canapac Certificate of Analysis No: 258/2025, dated on 26/03/2025.
- b) From Canapac Certificate of Analysis No: 192/2025, dated on 11/03/2025.
- c) From LEF Certificate of Analysis No: 25/2048, dated on 24/03/2025.

Legend: LOD - Limit of Detection; LOQ - Limit of Quantitation

Conclusion: Approved - Complies with specification

Prepared by	Approved by
	
Quality Control Manager	Qualified Person
<u>27/03/2025</u>	<u>27/03/2025</u>
Date	Date

<b>Product Name</b>	Remexian 33/1 VCC GWA 250g
<b>Dosage Form</b>	Bags of 250g
<b>Stength/ Potency</b>	33% THC; ≤1% CBD
<b>Batch Number</b>	2500861

<b>Pack Size</b>	250g
<b>Quantity</b>	68 un
<b>Manufacturing Date</b>	03/2025
<b>Expiry Date</b>	09/2025
<b>Country of market place</b>	Germany

<b>Raw Material of Herbal Origin-Cultivator</b>	Otoka Holdings Ltd. D.b.a. Victoria Cannabis Company 344 Mary Street Victoria BC, Canada, V9A 3V9
<b>API-Manufacturer</b>	Canapac, Lda. Rua 3B, Lote 48, Zona Industrial de Cantanhede, 3060-197 Cantanhede, Portugal
<b>Finished-Product Manufacturer</b>	Canapac, Lda. Rua 3B, Lote 48, Zona Industrial de Cantanhede, 3060-197 Cantanhede, Portugal
<b>Finished-Product Quality Control</b>	LEF/ INFOSAÚDE - Instituto de Formação e Inovação em Saúde, S.A. Rua das Ferrarias del Rel, nº6 Urbanização da Fábrica da Pólvora 2730 Barcarena, Portugal  Canapac Lda. Rua 3B, Lote 48, Zona Industrial de Cantanhede, 3060-197 Cantanhede, Portugal

<b>Certificate of Analysis number</b>	277/2025	<b>Issued date</b>	27/03/2025
<b>Specification</b>	SPE-0001333-ver.1.0		

I hereby certify that the above information is authentic and accurate. This batch of product has been manufactured (including packaging and quality control) at the above mentioned sites in full compliance with EU-GMP requirements.

**Deviation(s) Occurred:**     No     Yes    Deviation Nr: \_\_\_\_\_

**OOS(s) Occurred:**         No     Yes        OOS Nr: \_\_\_\_\_

**Comments/ Observations:** Not applicable.

Catarina Paixão  
Qualified Person Name

[Redacted Signature]

Qualified Person Signature

27/03/2025  
Date

