

CERTIFICATE OF CONFORMITY

Product Name	Remexian Balanced CNF 250g
Dosage Form	Bags of 250g
Stength/ Potency	9.67% THC; 15.11% CBD
Batch Number	2500131

Pack Size	250g
Quantity	180 un
Manufacturing Date	02/2025
Expiry Date	08/2025
Country of market place	Germany

Raw Material of Herbal Origin-Cultivator	Betralif Corp. 460 CHEMIN INDUSTRIEL L'ANGE GARDIEN, QC, CANADA, J8L 0A9
API-Manufacturer	Canapac, Lda. Rua 3B, Lote 48, Zona Industrial de Cantanhede, 3060-197 Cantanhede, Portugal
Finished-Product Manufacturer	Canapac, Lda. Rua 3B, Lote 48, Zona Industrial de Cantanhede, 3060-197 Cantanhede, Portugal
Finished-Product Quality Control	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

Certificate of Analysis number	160/2025	Issued date	28/02/2025
Specification	SPE-0000931-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.

CERTIFICATE OF ANALYSIS

CoA No.: 160/2025
Rev.: 001
Issued date: 28/02/2025

Internal Code		NAME		Batch no.	
FIN000128		Remexian Balanced CNF 250g		2500131	
Manufacture Date		Expiry date		Analysis Date	
08/02/2025		08/2025		28/02/2025	
				PRT-0000515-1.0 SPE-0000931-ver.2.0	
Analytical Protocol		Specification			
08/02/2025		08/2025		28/02/2025	
				PRT-0000515-1.0 SPE-0000931-ver.2.0	
Tests		Methods		Specification	
				Results	
1. Identification					
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	
2. Foreign Matter		EP monograph 07/2024:3028; Ph.Eur.2.8.2		≤ 2% w/w 0%	
3. Loss on Drying ^{a)}		EP monograph 07/2024:3028; Ph.Eur.2.2.32		≤ 12.0% w/w 8.9%	
4. Assay ^{a)}					
4.1. Total THC		EP monograph 07/2024:3028; Ph.Eur.2.2.29		Declared amount ± 10% 9.67%	
4.2. Total CBD				Declared amount ± 10% 15.11%	
5. Total CBN ^{a)}		EP monograph 07/2024:3028; Ph.Eur.2.2.29		≤ 1.0% w/w 0.1%	
6. Heavy Metals ^{b)}					
6.1. Cd				≤ 0.3 ppm < LOQ (0.005 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 < LOQ (0.020 ppm)	
7. Microbiology ^{b)}					
7.1. Total Aerobic Microbial Count (TAMC)		Ph.Eur. (5.1.8C, 2.6.12)		≤ 10 ⁵ CFU/g (max: 500 000 CFU/g) <1000 CFU/g	
7.2. Total Yeast and Mold Count (TYMC)				≤ 10 ⁴ CFU/g (max: 50 000 CFU/g) <1000 CFU/g	

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08/02/2025	08/2025	28/02/2025	PRT-0000515-1.0	SPE-0000931-ver.2.0
Tests	Methods	Specification	Results	
7.3. Bile-tolerant gram-negative bacteria	Ph.Eur. (5.1.8C, 2.6.31)	$\leq 10^4$ CFU/g	<10 CFU/g	
7.4. Escherichia coli		Absence in 1 g	Absent/g	
7.5. Salmonella		Absence in 25 g	Absent/25g	
7.6. Pseudomonas aeruginosa		Absence in 1 g	Absent/g	
7.7. Staphylococcus aureus		Absence in 1 g	Absent/g	
8. Aflatoxins ^{b)}				
8.1. B1	Ph. Eur. 2.8.18	≤ 2 µg/kg	< LOD (0.25 µg/kg)	
8.2. B1 + B2 + G1 + G2		≤ 4 µg/kg	< LOQ (0.5 µg/kg)	
9. Ochratoxin A ^{b)}	Ph.Eur. 2.8.22	≤ 20 µg/kg	< LOD (2 µg/kg)	
10. Pesticides ^{b)}	Ph.Eur. 2.8.13	Must comply with requirements of monograph	Complies	

Remarks:

- a) From Canapac Certificate of Analysis No: 062/2025, dated on 23/01/2025.
b) From Canapac Certificate of Analysis No: 011/2025, dated on 03/01/2025.

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

Conclusion: Approved - Complies with specification

Prepared by

Approved by