

CERTIFICATE OF RELEASE

Certificate number_version:	2025_021_00
-----------------------------	-------------

Product Name	CANNABISBLÜTEN Bathera 33/1
Strength/Potency	33 THC / ≤ 1 CBD
Dosage Form	Herbal drug
Batch Number	PGP2403P1YC
Manufacturing date	11/05/2025
Expiry Date	11/11/2025
API Manufacturer	Batherafarm Unipessoal, Lda Rua 5 de Outubro, Casais 85 e 86, Nucho de Pegões Velhos
Finished Product Manufacturer	2985-155 Santo Isidro de Pegões, Portugal
Finished Product Quality Control	INFOSAÚDE, Instituto de Formação e Inovação em Saúde, S.A. (LEF) Rua das Ferrarias del Rei, nº6, Urbanização da Fábrica da Pólvora 2730-269 Barcarena, Portugal Batherafarm Unipessoal, Lda Rua 5 de Outubro, Casais 85 e 86, Nucho de Pegões Velhos 2985-155 Santo Isidro de Pegões, Portugal
Country of marketplace	Germany
Pack Size (g) / Type	250 / Bag
Quantity (un)	310
Certificate of Analysis number	See remarks
Specification	ES_QA_Finish product 001_DE_rev4

CERTIFICATE OF RELEASE

Parameter		Monograph/ Guideline Reference	Method	Specification	Results
Identification	Method A Macroscopy examination	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Complies with requirements of monograph	Complies ^{b)}
	Method B Microscopy examination				Complies ^{b)}
	Method C HPTLC				Complies ^{b)}
Foreign Matter		- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.8.2	Maximum 2%	0.0 % ^{b)}
Loss on Drying		- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.2.32	Maximum 12.0%	8.8 % ^{b)}
Total CBN		- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.2.29	Maximum 1.0%	0.13 % ^{b)}
Assay	THC + THCA	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.2.29	29.7% - 36.3%	32.8 % ^{b)}
	CBD + CDBA	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.2.29	≤ 1%	0.16 % ^{b)}
Total Ash		- Ph. Eur. Monograph (1433) Herbal drugs	Ph. Eur. 2.4.16	Maximum 20.0%	9.4 % ^{b)}
Aflatoxins	B1	- Ph. Eur. Monograph (1433) Herbal drugs	Ph. Eur. 2.8.18	Not more than 2 µg/kg	<0.25 µg/kg (LOD) ^{b)}
Aflatoxins	Total B1, B2, G1, G2	- Ph. Eur. Monograph (1433) Herbal drugs	Ph. Eur. 2.8.18	Not more than 4 µg/kg for the sum of aflatoxins B1, B2, G1 and G2	<0.25 µg/kg (LOD) ^{b)}
Ochratoxin A		- Ph. Eur. Monograph (1433) Herbal drugs	Ph. Eur. 2.8.22	Not more than 20 µg/kg	<5 µg/kg (LOQ) ^{b)}
Pesticides		- Ph. Eur. Monograph (1433) Herbal drugs	Ph. Eur. 2.8.13	Not more than the limits specified in Ph. Eur. 2.8.13	Complies ^{a)}
Heavy Metals	Cadmium (Cd)	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.4.27	Maximum 0.3 ppm	<0.005 ppm (LOQ) ^{a)}
	Lead (Pb)	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.4.27	Maximum 0.5 ppm	<0.020 ppm (LOQ) ^{a)}
	Mercury (Hg)	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.4.27	Maximum 0.1 ppm	<0.050 ppm (LOQ) ^{a)}
	Arsenic (As)	- Ph. Eur. Monograph (3028) <i>Cannabis flos</i>	Ph. Eur. 2.4.27	Maximum 0.2 ppm	<0.020 ppm (LOQ) ^{a)}
Microbiological Tests	Total aerobic microbial count (TAMC)	- Ph. Eur. Monograph (5.1.8 C) Microbiological quality of herbal medicinal products and extracts	Ph. Eur. 2.6.12	≤ 10 ⁵ CFU/g (max.: 500 000 CFU/g)	1.00x10 ¹ CFU/g ^{b)}
	Total combined yeasts/moulds count (TYMC)			≤ 10 ⁴ CFU/g (max: 50 000 CFU/g)	1.00x10 ¹ CFU/g ^{b)}
	Bile-tolerant gram (-) bacteria		Ph. Eur. 2.6.31	≤ 10 ⁴ CFU/g	<10 CFU/g ^{b)}
	<i>Escherichia coli</i>			Absence / 1 g	Absent/g ^{b)}
	<i>Salmonella spp</i>			Absence / 25 g	Absent/25g ^{b)}
Average weight*		- Internal Method	Internal Method	250g – 252,5g	250 g

Legend: LOD – Limit of detection; LOQ – Limit of quantification

*Obtained from packaging batch record

CERTIFICATE OF RELEASE

Remarks:

- a) From LEF CoA nº 25/2350, issued date 14/04/2025
b) From LEF CoA nº 25/3767, issued date 27/05/2025

NA: Not applicable

Major or critical deviations:

NA

NA: Not applicable

Certification statement:

I hereby certify that all stages of cultivation, manufacturing and quality control, of this batch of finished product have been carried out at the mentioned sites, in full compliance with the GACP and cGMP requirements of the EU and the national/regional requirements of the destination country.

Qualified Person:

Name:

Signature:

Date: 28/05/2025