

Certificate of analysis No. 1039/2026

Product:	Cannabis for medical use
Type:	Cannabis Flos
Type of medicinal substance:	dried flower
Strain:	G Wagon
Trade name of the product:	VCC 36/1 GWA
Batch:	LP.159.GWA
Batch amount:	57 000 g
Number and size of packages:	228 x 250 g
Expiration day:	03.2027
Certified by:	Genetia Production s.r.o. IČ: 19452519
Customer:	Lagom Pharmatech s.r.o. IČ: 09514538

Test	Method	Acceptance limits (Specification)	Results	Complies
Pharmacognostic methods				
Macroscopic examination	SOP 5205	Complies with requirements of Ph. Eur. 3028 dark green, yellow or brown, no mould, aromatic odour	complies	yes
Foreign matter	Ph. Eur. 2.8.2	Complies with requirements of Ph. Eur. 3028 ≤ 2 %, no seeds, no leaves larger than 1 cm	complies	yes
Pesticide content -all mentioned pesticides	Ph. Eur. 2.8.13	Complies with requirements of Ph. Eur. 2.8.13 below specified limits	complies	yes
Mycotoxins quality -Aflatoxins B1 -Sum of aflatoxins (B1, B2, G1, G2)	Ph. Eur. 2.8.18	≤ 2 µg/kg ≤ 4 µg/kg	< LOQ (2 µg/kg) < LOQ (2 µg/kg)	yes
Mycotoxins quality -Ochratoxin A	Ph. Eur. 2.8.22	≤ 20 µg/kg	< LOQ (2 µg/kg)	yes
Microscopic examination	Ph. Eur. 2.8.23	Complies with requirements of Ph. Eur. 3028 characteristic glandular/covering trichomes	complies	yes
HPTLC examination	Ph. Eur. 2.8.25	Complies with requirements of Ph. Eur. 3028	complies	yes

This Certificate of Analysis (CoA) has been prepared and issued in accordance with internal SOP No. G SOP 5014 001.

Info@genetia.eu
+420 311 440 038

Genetia production s.r.o.
Inovační 122, 252 41 Zlatníky-Hodkovice
IČ: 194 52 519, DIČ: CZ699007808

		in accordance with reference solution		
Physical and physico-chemical methods				
Assay ¹⁾		Complies with requirements of Ph. Eur. 3028		
-Total THC	Ph. Eur. 2.2.29	32,4 – 39,6 %	34,6 %	yes
-Total CBD		≤ 1,0 %	0,21	
-Total CBN		≤ 1,0 %	0,2 %	
Loss on drying	Ph. Eur. 2.2.32	≤ 12,0 %	7,7 %	yes
Limit methods				
Heavy metal content		Complies with requirements of Ph. Eur. 3028		
-Cadmium	Ph. Eur. 2.4.27	≤ 0,3 ppm	< LOQ (0,002 ppm)	yes
-Mercury and its compounds		≤ 0,1 ppm	< LOQ (0,02 ppm)	
-Lead		≤ 0,5 ppm	< LOQ (0,01 ppm)	
-Arsenic		≤ 0,2 ppm	< LOQ (0,01 ppm)	
Biological methods				
Microbiological quality				
-TAMC	Ph. Eur. 2.6.12	max. 500 000 CFU/g	< 1000 CFU/g	yes
-TYMC		max. 50 000 CFU/g	< 100 CFU/g	
Microbiological quality				
-S. aureus	Ph. Eur. 2.6.13	absence/1 g	absence/1 g	yes
-P. aeruginosa		absence/1 g	absence/1 g	
Microbiological quality				
-BTGN	Ph. Eur. 2.6.31	≤ 10 ⁴ CFU/g	absence/1 g	yes
-E. Coli		absence/1 g	absence/1 g	
-Salmonella sp.		absence/25 g	absence/25 g	

1) Results are reported on a dry weight basis.

Date: 07.07.2026

I hereby confirm that the information provided is true and complete. Testing was performed in accordance with GMP requirements.

Controlled by:



Head of QC

QA Administrator

Qualified Person

This Certificate of Analysis (CoA) has been prepared and issued in accordance with internal SOP No. G SOP 5014 001.

info@genetia.eu
+420 311 440 038

Genetia production s.r.o.
Inovační 122, 252 41 Zlatníky-Hodkovice
IČ: 194 52 519, DIČ: CZ699007808

	Batch Release Certificate	Description:	CLP
		Number:	240/2026

Producer:	Genetia production s.r.o, Inovační 122, Zlatníky – Hodkovice, ID 19452519		
GMP certificate:	sukls225038/2025		
Trade name of the product:	VCC 36/1 GWA		
Product type:	Cannabis flos	Strain:	G Wagon
Destination country and purpose:	Germany – For placement on the market		
Batch number of starting material:	VC.170.080426_2	Country of origin:	Canada
Batch number:	LP.159.GWA	Production date:	01.07.2026
Batch amount and size	227 x 250 g	Expiration date:	03.2027
Retention samples:	2 x 250 g	Active ingredients and their content:	THC: 34,6 % CBD: < 1,0 %

The product was manufactured and packaged according to a GMP-compliant manufacturing process.

Date, stamp, name and signature: 

07 -07- 2026

The product has passed all prescribed checks and complies with the specification given in Certificate of analysis no. 1039/2026 .

Date, stamp, name and signature: 

07 -07- 2026

I hereby certify that all manufacturing phases of this batch of medicinal products have been carried out in full compliance with the requirements of the Czech Republic / EU for Good Manufacturing Practice of Medicinal Products, and is in compliance with the product specification and the marketing authorisation dossier.

Date, stamp, name and signature: 

07 -07- 2026

G SOP 3801 002-P3 Batch Certificate

