

VitaMoment GmbH

Barnerstraße 14 d
22765 Hamburg



Our Sign : QLT
Date : 17.09.2026

Certificate of analysis 26038138 - 001

Sample name : Leber-Komplex

Marking of sample : Charge: DE2601083; MHD: 07/2028

Customer No. : none

Packaging : Pre-packaging / Plastic container

Sample amount : 3 x 54,4 g

Shipping of sample : Courier Service

Sample entry : 31.07.2026

Entrance temperature : Room temperature

Sample taken : by sender

Begin/end of analysis : 31.07.2026 / 17.09.2026

The test results apply only to the test items described in the report. No responsibility is accepted for the validity of the results if any data or information provided by the customer may affect them. Data provided by the customer are clearly identified. The laboratory assumes no responsibility for the sampling including minimum quantities unless it was carried out by samplers from a company within the GBA Group or on its behalf. In this case, the results apply to the sample as received. The test report may not be published or reproduced, in whole or in part, without the written consent of the issuing company. The general terms and conditions are available at <https://www.gba-group.com/en/general-terms-and-conditions/>.

1 / 4

Dok.-Nr.: ML 510-01 # 2 V1 E, 511, 19.02.2026

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seit 1989

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Test Results

Microbiological Test	Result	Unit
Total Plate Count	$2,0 \cdot 10^1$	cfu/g
Yeasts / moulds		
Yeasts	<10	cfu/g
Moulds	$2,0 \cdot 10^1$	cfu/g
Enterobacteriaceae	<10	cfu/g
E. coli	<10	cfu/g
Salmonella	negative	/ 25 g

Chemical/Physical Test	Result	Unit	Declaration	$\pm$ MU	MU Source	ML
Lead	<0,020	mg/kg			I	3
Cadmium	<0,010	mg/kg			I	1
Mercury	<0,010	mg/kg			I	0,1
Arsenic	0,10	mg/kg		0,02	I	
Cholin, free and bound	81,8	mg/ daily serving	83	16	IV	
PAH						
Benzo(a)anthracene	<1,0	μ g/kg			I	
Chrysen	<1,0	μ g/kg			I	
Benzo(b)fluoranthene	<1,0	μ g/kg			I	
Benzo(a)pyrene	<1,0	μ g/kg			I	10
Sum of PAH	not detectable	μ g/kg			I	50
Weight per dosage form	0,83	g		0,01	VII	
Daily serving	2,0	capsule(s)				
Gluten	<5,0	mg/kg			I	
Iodine	0,4	mg/kg		0,1	IV	

Maximum levels for food supplements according to VO (EU) 2023/915

Assessment:

Results of microbiological analysis are unobjectionable regarding the tests carried out.

Regarding the determined levels of lead, cadmium, mercury and polycyclic aromatic hydrocarbons, the sample complies with the maximum levels for food supplements laid down in Regulation (EU) 2023/915 (Cat. 3.1.28; 3.2.21; 3.3.2; 5.1.16).

Result of Cholin corresponds to the declared value with sufficient accuracy.

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Hamburg, 17.09.2026

This test report is done automatically and is valid without signature.

Methods

Parameter	Method	DR
Total Plate Count	DIN EN ISO 4833-2: 2022-05 ^a ₀	m
Yeasts / moulds	BIOKAR Diagnostics, Symphony-Agar BM20208/BM19108: 2022-11 ^a ; validated according to EN ISO 16140-2 against EN ISO 21527-1/-2 2008-11 ₀	m
Enterobacteriaceae	Biomerieux, Rebecca-Agar AEB520020/AEB150022: 2020-09 ^a ; validated according to EN ISO 16140-2 against ISO 21528-2 2017-07 ₀	m
E. coli	Biomerieux, Rebecca-Agar AEB520020/AEB150022: 2020-09 ^a ; validated according to EN ISO 16140-2 against ISO 16649-2 2001-07 ₀	m
Salmonella	DIN EN ISO 6579-1: 2020-08 ^a ₀	m
Lead	DIN EN 15763, ICP-MS: 2010-04 ^a ₀	y
Cadmium	DIN EN 15763, ICP-MS: 2010-04 ^a ₀	y
Mercury	DIN EN 15763, ICP-MS: 2010-04 ^a ₀	y
Arsenic	DIN EN 15763, ICP-MS: 2010-04 ^a ₀	y
Pressure pulping	§ 64 LFGB L 00.00-19/1: 2015-06 ^a ₀	
Cholin, free and bound	FV-0243 02-003, LC-MS/MS: 2026-04 ^a ₁	z
PAH	HH-MA-M 02-105 # U, HPLC-FLD: 2023-06 ^a ₀	y
Sum of PAH	berechnet α	
Weight per dosage form	HH-MA-M 10-030, gravimetric: 2021-11 ^a ₀	z
Gluten	R-Biopharm AG RIDASCREEN Gliadin R7001, ELISA: 2021-10 ^a ₃	z
Iodine	DIN EN 15111: 2007-06 ^a ₁	z

The methods marked with ^a are accredited methods of the performing laboratory.

Testing laboratory: ₀GBA Hamburg ₁extern α automatically calculated from the system ₃GBA Hameln

MU-Source:

I: According to DIN ISO 11352 as expanded, combined measurement uncertainty with $k = 2$ (95 %), sampling not included

IV: According to subcontractor

VII: According to expert estimation

Decision rules:

m: The conformity assessment of microbiological measured values is performed without considering additional analytical measurands.

y: In conformity assessment, measurement uncertainty is disregarded for measured values below the tolerance limit. For measured values exceeding the tolerance limit, measurement uncertainty is subtracted from the measured value. If no conformity assessment is performed, measurement uncertainty serves as informational data only.

z: In conformity assessment, measurement uncertainty is disregarded and serves as informational data only.

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° According to DIN EN ISO 7218, this result must be stated as detected for statistical reasons.
The value given is not statistically significant.