



## **Analytical report**

AR-25-HD-025613-02



## **Testing laboratory:**

Eurofins Food & Feed Testing Czech Republic s.r.o. Zkušební laboratoř EUROFINS CZ Radiová 1285/7 102 00 Praha 10 - Hostivař

IČO: 27449408

tel.: +420 778 488 111 E-mail: ClientService.cz@ftcee.eurofins.com

## **Customer:**

Natios Health, s.r.o. Trocnovská 1088/2a 702 00 OSTRAVA 2, PŘÍVOZ CZECH REPUBLIC

Issue date 29.07.2025

**Sample code** 540-2025-00036479

10.07.2025 Sample reception date:

**Date of Testing** 10.07.2025 - 24.07.2025

**Sample information:** 

Sample name, extended: 1) NATIOS Magtein, Magnesium L-threonate, 90 veganských kapslí

 $^{1)}005\text{-}32407\text{-}2\bar{3}5676$ Sample description: Client Purchase order nr.:

Druhý test rutin/liver/magtein + omega kysel

08.07.2025 Order date: 1) NAT1755 Client sample code: Sampler: Customer Additional sample description: L231380

Microbiological tests

Parameter	Unit	Result	Uncertainty of measurement *	Method	Method principle	TZ
Aerobic Plate Count 30°C	cfu/g	<10		ČSN EN ISO 4833-1	E-Cultural technique (non-chromogenic media) [Total aerobic count 30°C <10>30000000 /g (1-6) PC casting ISO 4833]	A
Escherichia coli	cfu/g	<10		ČSN ISO 16649-2	E-Cultural technique (chromogenic media)	A
Yeast	cfu/g	<10		SOP.MB.014.PB	E-Cultural technique (non-chromogenic media)	A
Moulds	cfu/g	<10		SOP.MB.014.PB	E-Cultural technique (non-chromogenic media) [Plate count:DG18<0.95]	A
Salmonella	/25 g	Not Detected		ČSN EN ISO 6579-1	D-Cultural technique (non-chromogenic media)	A
Staphylococcus aureus	cfu/g	<10		ČSN EN ISO 6888-1	E-Cultural technique (non-chromogenic media)	A

Physical and chemical tests

- Hydrous und differences tools								
Parameter	Unit	Result	Uncertainty of measurement *	Method	Method principle	TZ		
Arsenic (As)	mg/kg	< 0.03		Internal Method LS-PP-CH-85	ICP-MS	SA		
Cadmium (Cd)	mg/kg	< 0.10		Internal Method LS-PP-CH-85	ICP-MS	SA		
Lead (Pb)	mg/kg	< 0.30		Internal Method LS-PP-CH-85	ICP-MS	SA		
Mercury (Hg)	mg/kg	< 0.010		LS-PP-CH-85	ICP-MS	SA		





Decision rule: If the testing laboratory issues a statement of conformity, the decision-making rule according to ch. 4.2.1 of ILAC document G8:09/2019 Guidelines for the use of decision rules and statement of conformity. In such a case, the measurement uncertainty is not taken into account for the conformity statement. If measurement uncertainty is included the decision, this information is included in the statement of conformity. In such a case, proceed according to chap. 4.2.3 ILAC G8:09/2019.

SOP, ŠPP - Standard operation procedure Notes:

ND - not detected by given method CFU - Colony forming unit

SN - subcontracted not accredited test

N - test outside of the accreditation scope of EUROFINS CZ NM - necessary quantity SA - subcontracted accredited test

\* - the expanded measurement uncertainty, as determined by the extension coefficient k = 2 (with a 95% probability), does not include sampling uncertainty; if

Ligora

TZ - type of test

A - test within the accreditation scope of EUROFINS CZ

the measurement uncertainty is expressed in %, it is its relative value

LOD - limit of detection, LOQ - limit of quantification, result between LOD and LOQ = detected

1) - Information supplied by customer

Unless otherwise stated in the notes, the place of the tests performance is workplace No. 1 - Prague - of EUROFINS CZ testing laboratory.

If the information supplied by the customer could have be to affect the validity of the results, the laboratory disclaims responsibility. For samples supplied by the customer, the results relate to the sample as received and provided by the customer. The measuring devices and gauges used for the test / tests have been calibrated and verified according to valid metrological regulations. The results of the measurements relate only to the subject of the tests and do not replace other documents, e.g. of an administrative nature. The result identified as subcontracting in this protocol is the result of subcontractor measurements based on contract, order. The protocol may be reproduced or incorporated into promotional materials only with the written consent of the EUROFINS CZ Testing Laboratory and only to the extent of such approval. Any alteration, reproduction of part of the test report is not permitted and such analytical report automatically becomes invalid. The authenticity and completeness of the report can be verified at the EUROFINS CZ test laboratory stated in the header of analytical report. This Test Report has been issued in accordance with the applicable Conditions of service available on request and accessible at www.eurofins.cz.

Responsible for correctness: Dominika Zigová

Worked out by: Nada Krejcová No. of document: 2025729154717156

Validity check of document

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**Test Certificate approved by:** 

Dominika Zigová Deputy Head of Microbiology