

## ANALYTICAL LABORATORIES microbiology - physicochemistry - sensory



GBA POLSKA Sp. z o.o. Member of GBA GROUP ul. Mochtyńska 65, 03-289 Warsaw, Poland

## TEST REPORT No.: B/0/07/2024/856/FM/2/EN

Customer: Order No.: 7FIT s.a. 55-200 Stanowice, ul. Stanowice 28A

B/0/07/2024/856

AE - accredited methodology (accreditation no. AB 1095) of flexible scope - reference if the law so provides / equivalent to reference (the result can be used to assess compliance in the legally regulated area).

Material/product tested: Dietary supplements											
Sample collection address: 55-200 Stanowice , Stanowice 82A											
Produc	Product name: 7NUTRITION ISOTONIC GOLD 1000 g ORANGE Date*: 24 lipca 2024										
Lot nur	production: nber:	15	7NUTRITION 15/07/2024 240715.2								
	collected according to: transported by: Shipping				Sample receiver:	GBA POLSKA er	nployee no.:	2729			
Sample	Sample	ur 1:	nreservedl	y Analysis start da	sis end date: 29-07-2024						
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	S			
Ł	Total microbial count	cfu/g	AE	PN-EN ISO 4833-1:2013-12, PN- EN ISO 4833-1:2013-12/Ap1:2016- 11, PN-EN ISO 4833-1:2013- 12/A1:2022-06	no requirements	<1,0x10 <sup>1</sup>					
Ł	Presence of Salmonella spp.	25g	AE	PN-EN ISO 6579-1:2017-04, PN- EN ISO 6579-1:2017-04/A1:2020- 09	no requirements	not detected in 25g					
Ł	Presence of presumptive Escherichia coli	lg	AE	PN-ISO 7251:2006	no requirements	absent in 1g					
Ł	Presence of coagulase-positive staphylococci (Staphylococcus aureus and other species)	1g	AE	PN-EN ISO 6888-3:2004, PN-EN ISO 6888-3:2004/AC:2005	no requirements	absent in 1g					
Ł	Count of yeasts and moulds	cfu/g	AE	PN-ISO 7954:1999	no requirements	<1,0x10 <sup>1</sup>					
Ł	Lead	mg/kg	AE	PN-EN 15763:2010	≤3.0 ; mg/kg ; Start 2023/915	< 0,010					
Ł	Mercury	mg/kg	AE	PN-EN 15763:2010	≤0.1 ; Start 2023/915	< 0,0010					
Ł	Cadmium	mg/kg	AE	PN-EN 15763:2010	no requirements	0,0027					

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Date\* - depending on the method of obtaining the sample by GBA POLSKA, it is the date of: collection (when the sample is collected only by a GBA POLSKA employee) or receipt (when the sample is collected from the Customer by a GBA POLSKA employee, is delivered by a courier company or delivered personally by the Customer). MU\*\* - expanded measurement uncertainty at the level of confidence app. 95% and the coverage factor k=2, does not take into account the sampling uncertainty, except when indicated in the remarks.

Mos a support interstitution in the request of the clock of the relation of the relation of the clock of the clock of the relation of the clock of t nigher than the measuring ranges of the methods are presented as "cvalue of the lower limit of the measuring range" or "> value of the upper limit of the measuring range, respectively. These values provide information about the research results. If expanded uncertainties are given with these test results, they apply to the lower or upper limit of the measuring range of the method. S – Statements of Conformity with the requirements or specifications relating to the results for the parameters indicated in a given row, where YES means conformity and NO means non-conformity with specification. The decision-making principle agreed with the Customer and the risks associated with it, as well as the identification of which specifications, standards or parts thereof are met and which are not, are provided in the Remarks. In case of obtaining the "test results", the Statements of Conformity for those "test results" that are meet the requirements of PCA Communication No. 353 of

August 24, 2021, it is carried out as part of the opinion and interpretation. The results refer only to the tested samples (sampled or received - in accordance with the information presented in the Test Report).

The information in table is included in the Test Report was provided by the Customer. The laboratory is not responsible for this information. The laboratory is not resposible for the method of sampling and the representativeness of the samples provided by the Customer for testing.

The Test Certificate without the written approval of the Laboratory shall not be reproduced except in full. The Laboratory does not store the samples after testing, unless otherwise agreed with the Customer. Place of performance of the tests ("Lab."): Ł -Łajski, ul. Kościelna 2a, 05-119 Legionowo, L - ul. Doświadczalna 50a, 20-280 Lublin, M - ul. Fabryczna 7, 41-404 Mysłowice, P – ul. Kazimierza Tymienieckiego 34, 60-681 Poznań, PS - in situ measurement.

NOTE: Original Test Report are issued in electronic form with the \*, pdf extension, signed with a qualified electronic signature. Therefore, all prints, unless certified as true copies, are copies.

## **Remarks:**

The second selective medium for detecting the presence of Salmonella spp. in accordance with PN-EN ISO 6579-1:2017-04, Mon-EN ISO 6579-1:2017-04/A1:2020-09 is RVS broth and Brilliance Salmonella/Agar. Coagulase is used to detect staphylococci-positive, Braid Parker RPF/agar was used.

Created on:	Authorized result:	Authorized Test report:	
31-07-2024	GBA POLSKA employee no.: 2642 GBA POLSKA employee no.: 2840 GBA POLSKA employee no.: 2866	Specialist in food and dietary supplements GBA POLSKA employee no: 2793	Signed with a qualified electronic signature

Report prepared in a single copy

The end of the Test Report