PO-14/F-01, issue 05 of 07-06-2024



## 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/0/07/2024/664/FM/2/EN

Customer: 7FIT s.a. 55-200 Stanowice, ul. Stanowice 28A

Order No.: B/0/07/2024/664

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 **Product name:** 7NUTRITION DEXTROSE GOLD 1000 g GREEN APPLE Date\*: 24 lipca 2024 Producer: 7NUTRITION Date of production: 28/06/2024 Lot number: 240628.1 Samples collected according to: Sample GBA POLSKA employee no.: 2729 Samples transported by: Shipping receiver Sample Sample no.: 43386/07/24 unreservedly Analysis start date: 24-07-2024 Analysis end date: 29-07-2024 evaluation: MU\*\* S Lab. Analyzed parameter Unit Accred. Test method Requirement Result PN-EN ISO 4833-1:2013-12, PN-Total microbial count cfu/g no requirements <1,0x101 EN ISO 4833-1:2013-12/Ap1:2016 Ł 11, PN-EN ISO 4833-1:2013-12/A1:2022-06 Presence of Salmonella spp. 25g PN-EN ISO 6579-1:2017-04, PNno requirements not detected in EN ISO 6579-1:2017-04/A1:2020-Ł PN-ISO 7251:2006 Presence of presumptive Escherichia 1g ΑE no requirements absent in 1g Ł PN-EN ISO 6888-3:2004, PN-EN Presence of coagulase-positive 1g ΑE no requirements absent in 1g staphylococci (Staphylococcus aureus ISO 6888-3:2004/AC:2005 Ł and other species) Count of yeasts and moulds cfu/g PN-ISO 7954:1999 <1,0x101 AE no requirements Ł PN-EN 15763:2010 < 0.010 +/-0.002 Lead mg/kg AE ≤3.0; mg/kg; Start 2023/915 Ł Mercury mg/kg PN-EN 15763:2010 ≤0.1; Start 2023/915 < 0,0010 +/-0,0002 Ł Cadmium mg/kg PN-EN 15763:2010 no requirements < 0,0020 +/-0,0003 Ł

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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.

Measurement uncertainty is provided when it is important for the reliability of test results or compliance with requirements/specifications and at the request of the Customer. The "test results" lower or higher than the measuring ranges of the methods are presented as "<value of the lower limit of the measuring range" or "> value of the upper limit of the measuring range", respectively. These values

nigher than the measuring ranges of the methods are presented as "cyalue of the lower limit of the measuring range" or "S value of the upper limit of the measuring range or 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 italics 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: 31-07-2024 Authorized result:

GBA POLSKA employee no.: 2642

GBA POLSKA employee no.: 2840 GBA POLSKA employee no.: 2866

**Authorized Test report:** 

Specialist in food and dietary supplements GBA POLSKA employee no: 2793

Signed with a qualified electronic signature

Report prepared in a single copy

Original of PDF: Customer, copy of PDF to: Laboratory archive

The end of the Test Report