

ANALYTICAL LABORATORIES

microbiology - physicochemistry - sensory





GBA POLSKA Sp. z o.o. (formerly: JARS S.A.) Member of GBA GROUP

Headquarter address: Łajski, ul. Kościelna 2a, 05-119 Legionowo, Poland

TEST REPORT No.: Ł/0/05/2022/2268/M/4/EN

Customer: FITNESS TRADING Robert Szulborski 18-300 Zambrów, ul. Sitarska 16

Order No.: Ł/0/05/2022/2268

- A accredited methodology (AB 1095); reference if the law so provides (the result can be used to assess compliance in the legally regulated area).
- AE accredited methodology (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).
- AR accredited methodology (AB 1095) equivalent to reference (the result can be used to assess compliance in the legally regulated area).
- MON methodology accredited in terms of "OiB"
- GMP+ methodology registered in the scope of GMP+ B11 protocol (feed testing)
 - A/P accredited methodology of the subcontractor
 - P non-accredited methodology of the subcontractor

Material/product tested: Food										
Address (sampling / collection):			18-300 Zambrów, ul.Sitarska 16							
Produc	t name: OstroVit	VITAMI	N C SHOT	Γ green apple 2000	Date: 25.05.2022					
Lot nun	production:	Ι	own produc Date min. do WCS001 g	urability: May 21, 2023						
	transported by: Shipping				Sample receiver:	GBA POLSKA er	nployee no.:	2549		
Sample no.: 23031/05/22 Sample evaluation:			nreservedly	y Analysis start da	te: 25-05-2022 Analysi	Analysis end date: 01-06-2022				
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	N		
Ł	Total microbial count	cfu/g	AE	PN-EN ISO 4833-1:2013-12, PN- EN ISO 4833-1:2013-12/Ap1:2016- 11	no requirements	<1,0 x 10 ¹				
Ł	Presence of Listeria monocytogenes	25g	AE	PN-EN ISO 11290-1:2017-07	Absent in 25g; Customer requirements	not detected in 25g				
Ł	Count of yeasts and moulds	cfu/g	AE	PN-ISO 21527-2:2009	no requirements	<1,0 x 10 ¹				
Ł	Presence of presumptive Escherichia coli	10g	AE	PN-ISO 7251:2006	absent in 10g; Principal's requirements	absent in 10g				
Ł	Presence of Salmonella spp.	10g	AE	PN-EN ISO 6579-1:2017-04, PN-EN ISO 6579-1:2017-04/A1:2020-09	absent in 10g; Principal's requirements	not detected in 10g				
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	absent in 1g; Principal's requirements	absent in 1g				

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** - 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 presented when: it is relevant to the validity or application of the test results, it affects conformity to a specification limit, or a customer's instruction so requires. 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. If expanded uncertainties are given with these test results, they apply to the lower or upper limit of the measuring range of the method. Moreover, in the case of these results, the conformity

statement should be treated as an opinion and interpretation. The above-described procedure does not apply to biological tests. The results relate to the tested samples (sampled or received - as reported in the test report).

The Lebstrey does not started the course of samples (sampled or received - as reported in the test report).

In the case of samples provided by the customer, the information presented in the representativeness of the samples provided by the customer for testing.

The test report includes test results of the following number of samples: 1 pc(s) and without the written approval of the Laboratory shall not be reproduced except in full.

Customer may file complains within 14 days from receiving the report.

The Laboratory does not store the samples after testing, unless otherwise agreed with the customer.

Place of performance of the tests (location codes): Ł - Łajski, L - Lublin, M - Mysłowice, PS - in situ measurement.

Remarks:

The second selective medium for the detection of Salmonella spp. In accordance with PN-EN ISO 6579-1: 2017-04, PN-EN ISO 6579-1: 2017-04 / A1: 2020-09 is RVS broth and RAPID Salmonella / Agar, and for detecting the presence of Listeria monocytogenes in accordance with PN-EN ISO 11290-1: 2017-07 is Palcam. Braid Parker RPF / agar was used to detect coagulase positive staphylococci.

NOTE: The original test reports are issued as PDF file, signed with a qualified electronic signature. Therefore, all prints are copies, unless certified to be true to the original PDF file.

Report prepared in a single copy	The end of	the Report	Original of PDF: Customer, copy of PDF to: Laboratory archive		
Created on:	Authorized by:	Approved by:			
10-06-2022	GBA POLSKA employee no.: 2244	Junior food specialist	t Signed with a qualified electronic signature		
		GBA POLSKA employe no.: 2550	ee		

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