AB 079

TEST REPORT NO 125320/23/GDY/2

Client **SFD SPÓŁKA AKCYJNA** GŁOGOWSKA 41 45315 OPOLE	Sample (according to declaration of Client) Sample description: ALLNUTRITION CREATINE MUSCLE MAX 250 g ORANGE Batch: 007223 Expiry date: 31.01.2025

Sample reception date:	**10.03.2023**	Sample status: no objections
Start of analysis	**15.03.2023**	
End of analysis	**22.03.2023**	Sample received from the Client
Test report date	**22.03.2023**	

Test Method	Unit	Result
# Creatine monohydrate Hadorn		
Creatine Monohydrate	g/100 g	76,1
* Number of yeasts and moulds at 25°C PN-ISO 21527-2:2009 (withdrawn)		
Number of yeasts	cfu/g	<1,0x10¹
Number of moulds	cfu/g	<1,0x10¹
* Presence of coagulase-positive staphylococci (Staphylococcus aureus and other species) in 1 g PN-EN ISO 6888-3:2004; PN-EN ISO 6888-3:2004/AC:2005	in 1 g	Not detected
* Presence of Escherichia coli in 1 g PN-ISO 7251:2006	in 1 g	Not detected
* Presence of Salmonella spp. in 25 g PN-EN ISO 6579-1:2017-04; PN-EN ISO 6579-1:2017-04/A1:2020-09	in 25 g	Not detected
* Presence of Listeria monocytogenes in 25 g PN-EN ISO 11290-1:2017-07	in 25 g	Not detected

Test: Creatine monohydrate was performed in laboratory EUROFINS VITAMIN TESTING DENMARK A/S VEJEN Denmark

Authorized by:
Anna Polanin, Manager, Microbiology Laboratory

Subcontracted test results are authorised by persons authorised by the external provider.

The test report bears the certified electronic seal of J.S. Hamilton Poland Sp. z o.o.

Laboratory address:
Ks. Stanisława Kujota 8, 70-605 Szczecin

THE END OF THE REPORT

* Test method accredited

J.S. HAMILTON POLAND Sp. z o.o.
TESTING LABORATORY
Chwaszczyńska 180, 81-571 Gdynia, Poland tel. +48 58 766 99 00

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Test performed by external provider

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TESTING LABORATORY
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GBAPOLSKA

ANALYTICAL LABORATORIES
microbiology - physicochemistry - sensory

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

AB 1095

TEST REPORT No.: B/0/08/2023/202/FM/3/EN

Customer: SFD S.A 45-315 Opole, ul. Głogowska 41

Order No.: B/0/08/2023/202

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:	**Dietary supplements**
Sample collection address:	45-323 Opole, Zielonogórska 4
Product name:	**ALLNUTRITION CREATINE muscle max 500 g passion fruit** **Date*: 21.08.2023**
Producer:	SFD S.A.
Date of production:	June 30, 2023
Lot number:	068223

Samples collected according to:
Samples transported by: Shipping
Sample receiver: GBA POLSKA employee no.: 2729

Sample no.: 26127/08/23 Sample evaluation: unreservedly Analysis start date: 21-08-2023 Analysis end date: 26-08-2023

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	N
Ł	Content of Polycyclic Aromatic Hydrocarbons (PAH) (from calculation)	µg/kg		PB-258/LF ed. 5 dated 02.01.2022	no requirements	< 1,30		
Ł	Benzo(a)pyrene	µg/kg	AE	PB-258/LF ed. 5 dated 02.01.2022	no requirements	< 1,30		
Ł	Mercury	mg/kg	AE	PN-EN 15763:2010	no requirements	< 0,001		
Ł	Lead	mg/kg	AE	PN-EN 15763:2010	no requirements	< 0,010		
Ł	Cadmium	mg/kg	AE	PN-EN 15763:2010	no requirements	< 0,002		
Ł	Arsenic	mg/kg	AE	PN-EN 15763:2010	no requirements	< 0,010		
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		

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	N
Ł	Presence of presumptive Escherichia coli	1g	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	Presence of Listeria monocytogenes	25g	AE	PN-EN ISO 11290-1:2017-07	no requirements	not detected in 25g		
Ł	Count of yeasts and moulds	cfu/g	AE	PN-ISO 7954:1999	no requirements	$<1,0 \times 10^1$		

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 collection (when the sample is collected from customer by a GBA Polska employee, is delivered by a courier company or delivered personally by the customer).
** - 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).
In the case of samples provided by the customer, the information presented in the report regarding these samples is the information provided by the customer. The Laboratory is not responsible for this information or for the method of sampling and 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 detecting the presence of Listeria monocytogenes in accordance with PN-EN ISO 11290-1:2017-07 is Palcam - incubation at 37°C ± 1°C. The second selective medium for detecting the presence 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 Brilliance Salmonella/Agar. Braid Parker RPF/agar was used for the detection of 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:	
29-08-2023	GBA POLSKA employee no.: 2244 GBA POLSKA employee no.: 2337 GBA POLSKA employee no.: 2642	Senior Food Specialist GBA POLSKA employee no.: 2653	**Signed with a qualified electronic signature**