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

TEST REPORT No.: Ł/0/03/2023/1297/M/6/EN

Customer:

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

Ł/0/03/2023/1297

Order No.:

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

Materi	al/product tested: Dietary s	upplemen	ts							
Sample	collection address:	4	5-315 Opo	le, ul.Głogowska 41						
Produc	t name: ALLNUT	RITION	FITKING	DELICIOUS PROTEIN CHI	PS 60g cheese	Date*: 09.0.	3.2023			
Produce			own production							
Lot nur	production: nber:		ebruary 16 6022023	, 2023						
-	collected according to: transported by: Shipping				Sample receiver:	GBA POLSKA e	nployee no.:	: 2684		
Sample	no.: 15311/03/23 Sample evaluation	n: u	nreservedly	y Analysis start da	te: 13-03-2023 Anal	ysis end date:	17-03-202	3		
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	Ν		
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 presumptive Escherichia coli	1g	AE	PN-ISO 7251:2006	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	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				

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 $37^{\circ}C \pm 1^{\circ}C$.

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	f the Report	Original of PDF: Customer, copy of PDF to: Laboratory archive			
Created on:	Authorized by:	Approved by:				
17-03-2023	GBA POLSKA employee no.: 2282	Senior Food Speciali	Signed with a qualified electronic signature			
		GBA POLSKA employ no.: 2653				

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

TEST REPORT No.: Ł/0/03/2023/1297/F/1/EN

Customer:

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

Ł/0/03/2023/1297

Order No.:

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

Materia	al/product tested: Dietary s	uppleme	nts						
Sample	collection address:	4	45-315 Оро	le, ul.Głogowska 41					
Produc	t name: ALLNU	FRITION	FITKING	DELICIOUS PROTEIN CHI	PS 60g cheese	Date*: 09.03.2023			
Produce Date of Lot nun	production:]	own production February 16, 2023 16022023						
	collected according to: transported by: Shipping				Sample receiver:	GBA POLSKA	employee no.:	2684	
Sample	no.: 15313/03/23 Sample evaluation	on:	unreservedly	y Analysis start da	ate: 10-03-2023 Analy	sis end date:	13-03-2023	3	
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	MU**	Ν	
Ł	Sucrose	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	0,80	+/-0,08		
Ł	Maltose	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	0,31	+/-0,04		
Ł	Total sugars (from calculations)	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	1,1	+/-0,1		
Ł	Lactose	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	< 0,05			
Ł	Glucose	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	< 0,05			
Ł	Fructose	g/100g	AE	PB-22/LF ed. 4 dated 02.01.2022	no requirements	< 0,05			

Date* - d	lepending on the	method of ob	taining the sa	mple by GBA I	Polska, it is the	date of: collecti	on (when the s	sample is collected	d only by a	GBA Polsk	a employee) or colle	ection (v	vhen the	sample is
collected	from customer b	y a GBA Pols	ka employee	, is delivered b	y a courier com	pany or deliver	ed personally l	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 "="" limit="" lower="" measuring="" of="" or="" range="" the=""> value of the upper limit of the measuring range",</value>
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 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:

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 o	f the Report	Driginal of PDF: Customer, copy of PDF to: Laboratory archive		
Created on:	Authorized by:	Approved by:			
13-03-2023	GBA POLSKA employee no.: 2337	Senior Food Specialist	Signed with a qualified electronic signature		