



TEST REPORT NO 412236/22/GDY

Client SFD SPÓŁKA AKCYJNA GŁOGOWSKA 41 45315 OPOLE		Sample (according to declaration of Client) Sample description: SFD WPC 80 PURE PROTEIN 700 g Batch: 07.2023.PE3.T24 Production date: 31.07.2021 Expiry date: 31.07.2023
Sample reception date:	14.09.2022	Sample status: no objections Sample received from the Client
Start of analysis	19.09.2022	
End of analysis	26.09.2022	
Test report date	26.09.2022	

Test Method	Unit	Result
* Number of yeasts and moulds at 25°C PN-ISO 21527-2:2009 (withdrawn)		
Number of yeasts	cfu/g	<1,0x10 ¹
Number of moulds	cfu/g	<1,0x10 ¹
* 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

Authorized by:
 Ada Okunek, Analysis Expert, Microbiology Laboratory
 Anna Polanin, Manager, Microbiology Laboratory

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

The results refer only to the samples received. When a measurement uncertainty is given, it is an expanded uncertainty estimated for a coverage factor k=2 at 95% confidence level and is not including sampling uncertainty, unless otherwise stated. When the conformity is stated J.S. Hamilton Poland Sp. z o.o. applies the simple acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "result" column of the accredited method contains a record: "<" or ">", it means, that it is the test outcome directly related to the lower or upper limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty relates only to the lower or upper limit of the measuring range of the accredited method respectively. In such a case, the Laboratory presents the opinion and interpretation in the "statement of conformity" column, which is based on the obtained test outcome. This test report may not be copied in part without the prior written permission of J.S. Hamilton Poland Sp. z o.o. The responsibility of J.S. Hamilton Poland Sp. z o.o. is limited solely to the data issued in its original. J.S. Hamilton Poland Sp. z o.o. does not permit the use of the PCA accreditation symbol AB 079 by customers, subcontractors, external service providers and other third parties. For further information please refer to the PCA document - DA-02. The service confirmed by this report is subject to the General Terms and Conditions of Services of J.S. Hamilton Poland Sp. z o.o. published on www.hamilton.com.pl.

* Test method accredited
 # Test performed by external provider

Analytical report AR-22-RE-014189-01

Sample code 122-2022-00012487

Issue date 22.02.2022

Client	SFD S.A. ul. Głogowska 41 45-315 Opole POLSKA
* Type of sample	SFD WPC 80 Pure natural 700 g 005-32419-2846004
* Prescriber	SFD S.A.
* Purchase order date	03.02.2022
* Client Purchase order nr.	1
Transport by	Courier
* Sampling Person	Principal
* Purpose of the testing	fulfillment of legal requirements
* Type of sampling	to guarantee its representativeness
Reception date	07.02.2022
* Batch number	06.2023.PE1.T24
* Best before date	2023-06-30
Sample condition	acceptable
Transport condition	at ambient temp.
Number of tested samples	1
* Client sample code	1
Start analysis	10.02.2022
End Analysis	22.02.2022

Results / Outcomes

Test code	Parameter	Method	Result	Unit	Uncertainty of measurement
DI004	Alanine (A)	AMSUR, IC-UV	4,09	g/100 g	± 14%
	Arginine (A)		2,07	g/100 g	± 14%
	Aspartic acid (A)		8,61	g/100 g	± 14%
	Glutamic acid (A)		14,2	g/100 g	± 14%
	Glycine (A)		1,47	g/100 g	± 14%
	Histidine (A)		1,42	g/100 g	± 14%
	Hydroxyproline (A)		<0,2 (LOQ)	g/100 g	± 14%
	Isoleucine (A)		4,84	g/100 g	± 14%
	Leucine (A)		8,65	g/100 g	± 14%
	Lysine (A)		7,73	g/100 g	± 14%
Ornithine (A)	<0,05 (LOQ)	g/100 g	± 14%		

	Phenylalanine (A)		2,60	g/100 g	± 14%
	Proline (A)		5,05	g/100 g	± 14%
	Serine (A)		4,08	g/100 g	± 14%
	Threonine (A)		5,57	g/100 g	± 14%
	Tyrosine (A)		2,26	g/100 g	± 14%
	Valine (A)		4,67	g/100 g	± 14%
DJ011	Cysteine +Cystine (A)	IC-UV	1,97	g/100 g	± 14%
	Methionine (A)		1,92	g/100 g	± 14%
HU00C	Gliadin (A)	Internal Method FAML-A-32:2021 (RIDASCREEN®G liadin, ELISA)	<2,50	mg/kg	
	Gluten (A)		<5,00	mg/kg	
HU00G	Soy (A)	Internal Method FAML-A-85:2021 (Congen SureFood® A, RT-PCR)	nie wykryto		
KT02C	Mercury (Hg) (A)	LS-PP-CH-30, AAS-AMA	<0,01	mg/kg	
KT0A2	Cadmium (Cd) (A)	LS-PP-CH-85, ICP-MS	<0,1	mg/kg	
KT0A3	Lead (Pb) (A)	LS-PP-CH-85, ICP-MS	<0,3	mg/kg	
ST0E7	Nitrogen - Kjeldahl (A)	PN EN ISO 8968-3: 2008, Calculation	12,31	%	± 0,98
	Protein * 6.38 (A)		78,54	%	± 6,28
ST0YE	Dry matter (A)	PN-A-79011-3:199 8, weight	95,4	%	± 14,3
	Moisture (103 °C) (A)		4,6	%	± 0,7
ST15G	Fructose (A)	PB/CH/36 wyd. 4 z dnia 31.10.2019, LC-RI	<0,10	%	
	Galactose (A)		<0,10	%	
	Glucose (A)		<0,10	%	
	Lactose (A)		3,37	%	± 0,78
	Maltose (A)		<0,10	%	
	Raffinose (A)		<0,10	%	
	Sucrose (A)		<0,10	%	
	Total sugars (A)		3,37	%	± 0,78

A = Method accredited

x = Data provided by the customer

Details of laboratory accreditation:

DI004, DJ011: Eurofins Vitamin Testing Denmark RE00037: DS EN ISO/IEC 17025 DANAK 581

HU00C, HU00G: Eurofins Food Analytica Gyula RE00049: MSZ EN ISO/IEC 17025:2005

ST0E7, ST0YE, ST15G: Eurofins Polska Sp. (Malbork) PS02: AB 1334

KT02C, KT0A2, KT0A3: Eurofins Environment Testing Slovakia Turčianske RE000HB: ISO/IEC 17025:2017 SNAS S-406

+/- Uncertainty of measurement presented as expanded uncertainty of measurement (95%; k=2).

JUDGEMENT

Protein content calculated on dry matter = 82,33 %



Approved by: Alicja Milczarek
Analytical Service Manager

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AB 1334

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 ul. Głogowska 41
 45-315 Opole
 POLSKA

Issue date 11.02.2022

Analytical report AR-22-RE-010474-01



Sample code 122-2022-00012316

x Type of sample	SFD WPC 80 Pure natural 700 g 005-32419-2846004
x Prescriber	SFD S.A.
x Purchase order date	03.02.2022
Reception date	07.02.2022
Transport by	Courier
Sample condition	acceptable
Transport condition	at ambient temp.
x Sampling Person	Principal
x Type of sampling	to guarantee its representativeness
x Purpose of the testing	fulfillment of legal requirements
x Best before date	30.06.2023
x Batch number	06.2023.PE1.T24
Number of tested samples	1
Start analysis	07.02.2022
End Analysis	11.02.2022

Results / Outcomes

UM2PF	The presence of Salmonella spp., Breeding method with biochemical and serological confirmation (A)
Method	PN-EN ISO 6579-1:2017-04+A1:2020-09, D-Cultural technique (non-chromogenic media)
Salmonella	Not Detected /25 g
UMIMW	Coagulase positive Staphylococcus D Abs Pres /1 g ISO 6888-3 (A)
Method	PN-EN ISO 6888-3:2004+AC:2005, D-Cultural technique (MPN tubes)
Coagulase positive Staphylococcus	Not Detected /1 g
UMLS5	The presence of presumptive Escherichia coli, Breeding method with biochemical confirmation (A)
Method	PN-ISO 7251:2006, D-Cultural technique (MPN tubes)
Escherichia Coli	Not Detected /1 g

UMULJ	Enterobacteriaceae 37°C <10 >3000000 /g 4 (1-4) VRBD Agar-P-37 ISO 21528-2 (A)		
Method	PN-EN ISO 21528-2:2017-08, E-Cultural technique (non-chromogenic media)		
Enterobacteriaceae 37°C		< 10	cfu/g
ZM02A	Yeasts & Moulds E [PB/MBK/40] <10 >1 500 000 /g (1-4) PYM-PF 3M™ Petrifilm™ Rapid Yeast and Mold Cou (A)		
Method	3M™ Petrifilm™ Rapid Yeast and Mold Count Plates, E-Cultural technique (media film)		
Yeast & Moulds		< 10	cfu/g

A = Method accredited

x = Data provided by the customer

Authorized by: Anna Broda-Cnota
Analytical Service Manager

Approved by: Justyna Nowak
Analytical Service Manager

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