

Eurofins Polska Sp. z.o.o. ul. Dubois 118D 93-465 Łódź **POLAND**

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Analytical report AR-22-E8-055513-01

Sample code 297-2022-00078576

Issue date 25.07.2022

Client SFD S.A.

> ul. Głogowska 41 45-315 Opole **POLSKA**

x Type of sample SFD Collagen Premium 400 g

005-32419-2881349

x Prescriber SFD S.A. x Purchase order date 11.07.2022

x Client Purchase order nr. Transport by Courier x Sampling Person principal

x Purpose of the testing fulfillment of legal requirements x Type of sampling to guarantee its representativeness

Reception date 14.07.2022 x Batch number 03.2024.51442 x Best before date 2024-03-31

^x Packaging manufacturer packaging

Sample condition acceptable **Transport condition** at ambient temp.

Number of tested samples x Client sample code

Start analysis 15.07.2022 **End Analysis** 23.07.2022

Results / Outcomes

Test code	Parameter	Method	Result	Unit	Uncertainty of measurement
A7295	Calciferol quantified as cholecalciferol (A)	USP 41/ NF 36 method 1, LC-DAD	192 // 38 ,4	μg/100 g // μg/20 g	± 20%
DJCV3	Vitamin C (A)	ISO 20635:2018, LC-DAD	5740 // 1148	mg/100 g // μg/20 g	± 10%
SZQA3	Calcium (Ca) (A)	LS-PP-CH-2/22, ICP-OES	5630 // 112 ,6	mg/kg // mg/20 g	± 15%
SZQA9	Manganese (Mn) (A)	LS-PP-CH-2/21, ICP-OES	95 ,3 // 1,91	mg/kg // mg/20 g	± 15%

A = Method accredited

Details of laboratory accreditation:

A7295, DJCV3: Eurofins Vitamin Testing Denmark RE00037: DS EN ISO/IEC 17025 DANAK 581

SZQA3, SZQA9: Eurofins Environment Testing Slovakia Turčianske RE000HB: ISO/IEC 17025:2017 SNAS S-406

x = Data provided by the customer

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

Aligo Milmorel

Approved by: Alicja Milczarek Analytical Service Manager

^{1.} The results apply to samples received and analyzed.
2. The test results shall not be reproduced except in full without the written permission of Eurofins Polska Sp. z o.o.
3. Laboratory measurement uncertainty is given when it is relevant to the validity of the test result or the application of the test results; it is agreed with the client; if the uncertainty of measurement affects compliance with the specified limit.
4. The client has the right to submit a complaint within 14 days of receiving the analytical report. May be admitted only complaint in writing, by email reklamacje@eurofins.pl or by mail.
5. Approved analytical results made by subcontractors are authorized by persons authorized in the laboratory of the subcontractor.
6. In case a Customer demands a statement of conformity, or a requirement related to a test and the decision making rule is not included in the documents listed above, the Laboratory appoints a rule to be applied.

applied.
7. The laboratory is not responsible for the data provided by customers. The data provided may affect the validity of the results.











TEST REPORT NO 252534/23/GDY

Client SFD SPÓŁKA AKCYJNA GŁOGOWSKA 41 45315 OPOLE		Sample (according to declaration of Client) Sample description: SFD COLLAGEN PREMIUM RASPBERRY STRAWBERRY 400 g Batch: 03.2025/62033 Production date: 01.03.2023	
Sample reception date:	17.05.2023	Sample status: no objections	
Start of analysis	19.05.2023		
End of analysis	01.06.2023	Sample received from the Client	
Test report date	01.06.2023		

Test Method	Unit	Result	
* Content of elements ²⁾ PN-EN 15763:2010			
Lead (Pb)	mg/kg	< 0,010 (0,010 ± 0,003)	
Cadmium (Cd)	mg/kg	0,0030	
Mercury (Hg)	mg/kg	0,0023	
Collagen 1) Calculated	mg/dose	10300	
Hydroxyproline ¹⁾ PB-53/HPLC ed. II of 30.12.2008	mg/dose	1280	
* Aerobic colony count at 30°C PN-EN ISO 4833-1:2013-12	cfu/g	<1,0x10¹	
* 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	

Dose declared by the Client: 20 g.

²⁾ The lower limit of the measuring range of the accredited method, which is also the limit of quantification set by the Laboratory.











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TEST REPORT NO 252534/23/GDY

Authorized by:
Ada Okunek, Analysis Expert, Microbiology Laboratory
Anna Polanin, Manager, Microbiology Laboratory
Ewa Ostrach-Grzybowska, Analysis Expert, Vitamin Analysis Laboratory
Katarzyna Szpinda, Analysis Expert, Spectrometry Laboratory

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

Chwaszczyńska 180, 81-571 Gdynia 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: "c" 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, whereas the given expanded measurement uncertainty relates only to the lower or upper limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty relates only to the Open limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty relates only to the Open limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty relates only to the Open limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty relates only to the Open limit of the measuring range of the accredited method, whereas the given expanded measurement uncertainty, unless otherwise reported. If the "result" column, which is accredited method contains a record." "C" 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, unless otherwise reported. If the "result" column, which is accredited method contains a recordited method contains a reco

* Test method accredited

Test performed by external provider