



ANALYTICAL LABORATORIES
microbiology - physicochemistry - sensory

GBA POLSKA Sp. z o.o.
Member of GBA GROUP
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AB 1095

TEST REPORT No: B/0/07/2026/518/FM/12/EN

Customer: MZ-STORE SPÓŁKA AKCYJNA 84-240 Reda, ul. ul. Cypriana Kamila Norwida 47

Order No: B/0/07/2026/518

A - accredited methodology (accreditation no. AB 1095); reference – if the law so provides (the result can be used to assess compliance in the legally regulated area).

AE - accredited methodology (accreditation no. 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).

NA - non-accredited methodology, covered by the PN-EN ISO/IEC 17025:2018-02 system

Material/product tested:			Dietary supplements												
Product name:			Omega-3 33/22 TG 1000mg softgel (Fish Oil TG)				Date*: 10 July 2026								
Producer:															
Date of production:															
Lot number:															
Sampling according to:			-			Received by:			GBA POLSKA employee no: 3255						
Samples transported by:			Shipping												
Sample no:		39708/07/26		Sample condition:		correct		Analysis start date:		10-07-2026		Analysis end date:		21-07-2026	
Lab.	Analyzed parameter		Unit	Accred.	Test method		Requirement		Result		U	S/OI			
L	Coliforms count		cfu/g	AE	PN-ISO 4832:2007		no requirements		<1,0 x 10 ³			-			
L	Total microbial count		cfu/g	AE	PN-EN ISO 4833-1:2013-12, PN-EN ISO 4833-1:2013-12/Ap1:2016-11, PN-EN ISO 4833-1:2013-12/A1:2022-06		no requirements		<1,0 x 10 ⁴			-			
L	Presence of presumptive Escherichia coli		1g	AE	PN-ISO 7251:2006		no requirements		absent in 1g			-			
L	Presence of Listeria monocytogenes		25g	AE	PN-EN ISO 11290-1:2017-07		no requirements		not detected in 25g			-			
L	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			-			
L	Count of yeasts and moulds		cfu/g	AE	PN-ISO 7954:1999		no requirements		<1,0 x 10 ³			-			
L	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			-			
L	Mercury		mg/kg	AE	PN-EN 15763:2010		≤ 0.10 ; mg/kg ; Commission Regulation (EU) 2023/915 of 25 April 2023 on maximum levels of certain contaminants in food and repealing Regulation (EC) No 1881/2006 (as amended)		< 0,0010		0.0002	MEET			

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
L	Lead	mg/kg	AE	PN-EN 15763:2010	≤ 3.0 ; mg/kg ; Commission Regulation (EU) 2023/915 of 25 April 2023 on maximum levels of certain contaminants in food and repealing Regulation (EC) No 1881/2006 (as amended)	< 0,010	0.002	MEET
L	Cadmium	mg/kg	AE	PN-EN 15763:2010	≤ 1.0 ; mg/kg ; Commission Regulation (EU) 2023/915 of 25 April 2023 on maximum levels of certain contaminants in food and repealing Regulation (EC) No 1881/2006 (as amended)	< 0,0020	0.0003	MEET
L	Nervonic Acid (C24:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,73	0.35	-
L	Conjugated Linoleic Acid, CLA (C18:2 c9,t11)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,11	0.02	-
L	Cis-8,11,14-Octadecatrienoic Acid (C18:3n4)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,06	0.21	-
L	Octadecatrienoic Acid - Sum Of Trans Isomers (C18:3 trans)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
L	Heneicosanoic Acid (C21:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,05	0.01	-
L	Cis-8,11,14-Eicosatrienoic Acid (C20:3n6)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	2,49	0.50	-
L	Docosanoic Acid (C22:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
L	Cis-7, 10, 13, 16, 19 - Docosapentaenoic Acid, DPA (C22:5n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	5,09	1.02	-
L	Octadecenoic Acid - Sum Of Trans Isomers (C18:1 trans)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,19	0.04	-
L	Octadecadienoic Acid - Sum Of Trans Isomers (C18:2 trans)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,45	0.09	-
L	Heptadecanoic Acid (C17:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,08	0.02	-
L	Stearic Acid (C18:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	2,33	0.47	-

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
Ł	Tetracosanoic Acid (C24:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Tricosanoic Acid (C23:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Cis-11,14,17-Eicosatrienoic Acid (C20:3n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,27	0.05	-
Ł	Cis-13,16-Docosadienoic Acid (C22:2)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Palmitic Acid (C16:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	2,18	0.44	-
Ł	Undecanoic Acid (C11:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Myristic Acid (C14:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,51	0.10	-
Ł	Caprylic Acid (C8:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,13	0.03	-
Ł	Pentadecanoic Acid (C15:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Cis-9,12-Hexadecadienoic Acid (C16:2n4)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,13	0.03	-
Ł	Cis-6,9,12,15-Octadecatetraenoic Acid (C18:4n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	2,57	0.51	-
Ł	Cis-11-Docosic Acid (C22:1n1c)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,29	0.06	-
Ł	Arachidic acid (C20:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,31	0.06	-
Ł	Gamma Linolenic Acid, GLA (C18:3n6)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,12	0.02	-

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
Ł	Cis-11-Eicosenoic Acid (C20:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,58	0.32	-
Ł	Alpha linolenic acid, ALA (C18:3n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,92	0.18	-
Ł	Cis-11,14-Eicosadienoic Acid (C20:2)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,47	0.09	-
Ł	Erucic Acid (C22:1n9)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,53	0.31	-
Ł	Arachidonic acid, ARA (C20:4n6)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,25	0.05	-
Ł	Cis-10-Pentadecenoic Acid (C15:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Palmitoleic Acid (C16:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,95	0.19	-
Ł	Capric acid (C10:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	0,05	0.01	-
Ł	Lauric Acid (C12:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Tridecanoic Acid (C13:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Myristoleic Acid (C14:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Butyric Acid (C4:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Caproic acid (C6:0)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Cis-4,7,10,13,16,19 - Docosahexaenoic Acid, DHA (C22:6n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	24,88	4.98	-

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
Ł	Cis-5,8,11,14,17 - Eicosapentaenoic Acid, EPA (C20:5n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	37,51	7.50	-
Ł	Cis-Vaccenic Acid (C18:1n7c)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	2,08	0.42	-
Ł	Cis-8,11,14,17-Eicosatetraenoic Acid (C20:4n3)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,45	0.29	-
Ł	Cis-10-Heptadecenoic Acid (C17:1)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	< 0,05	0.01	-
Ł	Oleic Acid (C18:1n9c)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	5,74	1.15	-
Ł	Linoleic Acid, LA (C18:2n6c)	% in fat	AE	PB-191/LF ed. 6 of 01.10.2024	no requirements	1,31	0.26	-
Ł	An anisidine number	-	A	PN-EN ISO 6885:2016-04	no requirements	6,8	1.4	-
Ł	Peroxide value	meq O2/kg	A	PB-72/LF ed. 7 of 11.12.2024	no requirements	0,88	0.09	-
Ł	TOTOX indicator	-	NA	PN-EN ISO 6885:2016-04	no requirements	8,6		-

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 receipt (when the sample is collected from the Customer by a GBA POLSKA employee, is delivered by a courier company or delivered personally by the Customer).

U - 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 provided when it is important for the reliability of test results or compliance with requirements/specifications and at the request of the Customer. 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. These values provide information about the research results. If expanded uncertainties are given with these test results, they apply to the lower or upper limit of the measuring range of the method.

S/OI - statements of conformity/opinion and interpretation, where:

S - statements of conformity with the requirements or specifications relating to the results for the parameters indicated in a given row, where CONFORMING means conformity and NON CONFORMING means non-conformity with specification. The decision rules agreed with the Customer and the risks associated with it, as well as the identification of which specifications, standards or parts thereof are met and which are not, are provided in the Remarks. In case of obtaining the "test results", the Statements of Conformity for those "test results" that are meet the requirements of PCA Communication No. 353 of August 24, 2021, it is carried out as part of the opinion and interpretation.

OI - opinion and interpretation of the Laboratory in relation to the qualitative results/results obtained below/above the method range, where MEET means complying with the requirements and NOT MEET means not complying with the requirements.

The results refer only to the tested samples (sampled or received - in accordance with the information presented in the Test Report).

The information included in the Test Report was provided by the Customer. The laboratory is not responsible for this information. The laboratory is not responsible for the method of sampling and the representativeness of the samples provided by the Customer for testing.

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

Place of performance of the tests ("Lab."): Ł - Łąjski, ul. Kościelna 2a, 05-119 Legionowo, Ł - ul. Doświadczalna 50a, 20-280 Lublin, M - ul. Fabryczna 7, 41-404 Mysłowice, P - ul. Jasielska 16a, 60-476 Poznań, W - ul. Ząbkowska 18, 03-735 Warszawa, PS - in situ measurement.

NOTE: Original Test Report is issued in electronic form with the *.pdf extension, signed with a qualified electronic signature. Therefore, all prints, unless certified as true copies, are copies.

Remarks:

The second selective medium for detecting the presence of *Listeria monocytogenes* according to 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. according to PN-EN ISO 6579-1:2017-04, MON-EN ISO 6579-1:2017-04/A1:2020-09 is RVS broth and Brilliance *Salmonella*/Agar. For the detection of staphylococci coagulase-positive Braid Parker RPF/agar medium was used. The temperature used for incubation of coliform bacteria: 37°C±1°C.

Created on: 22-07-2026	Authorized result: GBA POLSKA employee no: 2424 GBA POLSKA employee no: 2486 GBA POLSKA employee no: 2705 GBA POLSKA employee no: 2883 GBA POLSKA employee no: 2978 GBA POLSKA employee no: 3285	Authorized Test report: Food Research Industry Documentation Team Leader GBA POLSKA employee no: 2942 Signed with a qualified electronic signature 
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Report prepared in a single copy

Original of PDF: Customer, copy of PDF to: Laboratory archive

The end of the Test Report