



ANALYTICAL LABORATORIES
microbiology - physicochemistry - sensory

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



AB 1095

TEST REPORT No: B/0/07/2026/389/F/1/EN

Customer: Nadarra GmbH 20149 Hamburg, Harvestehuder Weg 48

Order No: B/0/07/2026/389

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).

Material/product tested:		Dietary supplements						
Product name:		Mama Nadarra, 120 capsules-transparent capsules					Date*: 08 July 2026	
Producer:		NADARRA GmbH						
Date of production:		23/06/2026						
Lot number:		MANAD/23062026						
Sampling according to:		-			Received by:		GBA POLSKA employee no: 3344	
Samples transported by:		Shipping						
Sample no:	35865/07/26	Sample condition:	correct		Analysis start date:	08-07-2026	Analysis end date:	17-07-2026
Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
Ł	Vitamin B12 content (as methylcobalamin)	µg/dose	A	PB-304/LF ed. 2 of 06.03.2025	no requirements	4,2	1.3	-
Ł	Vitamin B12 content (as methylcobalamin)	µg/100g	A	PB-304/LF ed. 2 of 06.03.2025	no requirements	1120,0	336.0	-

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 in italics 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.

The Test Report without the written approval of the Laboratory shall not be reproduced except in full.

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

Place of performance of the tests ("Lab."): Ł - Łajski, ul. Kościelna 2a, 05-119 Legionowo, L - ul. Doświadczalna 50a, 20-280 Lublin, M - ul. Fabryczna 7, 41-404 Mysłowice, P - ul. Jasielska 16a, 60-476 Poznań, W - ul. Żąbkowska 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:

Dose: 375mg capsule - in accordance with the Client's declaration.

Created on: 20-07-2026	Authorized result: GBA POLSKA employee no: 2705	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



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

TEST REPORT No: B/0/07/2026/389/F/NN/2/EN

Customer: Nadarra GmbH 20149 Hamburg, Harvestehuder Weg 48

Order No: B/0/07/2026/389

Material/product tested: Dietary supplements

Product name: Mama Nadarra, 120 capsules-transparent capsules

Date*: 08 July 2026

Producer: NADARRA GmbH

Date of production: 23/06/2026

Lot number: MANAD/23062026

Sampling according to: -

Samples transported by: Shipping

Received by: GBA POLSKA employee no: 3344

Sample no: 35865/07/26

Sample condition: correct

Analysis start date: 08-07-2026

Analysis end date: 17-07-2026

Lab.	Analyzed parameter	Unit	Accred.	Test method	Requirement	Result	U	S/OI
Ł	Content of vitamin B9 (5-Methyltetrahydrofolate Glucosamine)	µg/dose	NN	PB-257/LF ed. 6 of 16.05.2025	no requirements	656	131	-
Ł	Content of vitamin B9 (5-Methyltetrahydrofolate Glucosamine)	µg/100g	NN	PB-257/LF ed. 6 of 16.05.2025	no requirements	175025	35,005	-

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Remarks: Dose: 375mg capsule - in accordance with the Client's declaration.

Created on: 20-07-2026

Authorized result: GBA POLSKA employee no: 2705

Authorized Test report: Food Research Industry Documentation Team Leader

Signed with a qualified electronic signature

GBA POLSKA employee no: 2942



Report prepared in a single copy

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

The end of the Test Report



TEST REPORT NO 619440/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028	
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619440/26/GDY Sample received from the Client	
Start of analysis	14.07.2026		
End of analysis	20.07.2026		
Test report date	20.07.2026		

Test Method	Unit	Result
* Vitamin D3 ¹⁾ PN-EN 12821:2009		
Vitamin D3 (cholecalciferol)	µg/capsule	40,1 ± 6,0
	IU/capsule	1600 ± 240

1) Capsule weight declared by the Client: 800 mg.

Test report authorized by:
ID: 91, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

The results refer only to the samples received and tested. The laboratory assumes no responsibility for the method of sample collection, the conditions of their transport, or for the information provided by the Client, including data that may affect the validity of the test results. 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 no criterion for a given test or tests in the analyzed matrix has been specified in the reference document indicated by the Client, confirmation of compliance is not possible. If the "result" column 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 method. If an expanded measurement uncertainty is given for such a test outcome, it relates only to the lower or upper limit of the measuring range of the method, respectively. In the case where the Laboratory base on the obtained test outcome, "statement of conformity" column presents an opinion and interpretation. 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

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TEST REPORT NO 620454/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028	
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620454/26/GDY Sample received from the Client	
Start of analysis	10.07.2026		
End of analysis	20.07.2026		
Test report date	20.07.2026		

Test Method	Unit	Result
* Vitamin B6 ¹⁾ PN-EN 14164:2014-08		
Vitamin B ₆ (pyridoxine hydrochloride)	mg/capsule	3,07 ± 0,61
Vitamin B6 (pyridoxine)	mg/capsule	2,52 ± 0,50

¹⁾ Capsule weight declared by the Client: 375 mg.

Test report authorized by:
ID: 91, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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TEST REPORT NO 620446/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620446/26/GDY Sample received from the Client
Start of analysis	08.07.2026	
End of analysis	16.07.2026	
Test report date	16.07.2026	

Test Method	Unit	Result
* Presence of Escherichia coli in 1 g PN-ISO 7251:2006	in 1 g	Not detected
* 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 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
* Number of Listeria monocytogenes at 37°C PN-EN ISO 11290-2:2017-07	cfu/g	<1,0x10 ¹

Test report authorized by:
ID: 343, Analysis Expert, Microbiology 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

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Test performed by external provider

THE END OF THE REPORT

TEST REPORT NO 619442/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619442/26/GDY Sample received from the Client
Start of analysis	09.07.2026	
End of analysis	15.07.2026	
Test report date	15.07.2026	

Test Method	Unit	Result
* # Iodine ¹⁾ VDLUFA VII, 2.2.2.3; 2011		
Iodine	µg/capsule	215,2 ± 83,1
Iodine calculated as potassium iodate ²⁾	µg/capsule	362,4 ± 139,9

1) Capsule weight declared by the Client: 800 mg.

2) Parameter not included in the scope of accreditation.

Test: Iodine was performed in laboratory with an accreditation number D-PL-14165-01-00

Test report authorized by:
 ID: 548, Authorization Section Manager, Authorization Section

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

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* Test method accredited

Test performed by external provider

THE END OF THE REPORT

TEST REPORT NO 619437/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619437/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	15.07.2026	
Test report date	15.07.2026	

Test Method	Unit	Result
* # Choline ¹⁾ AOAC Vol 91,1 2008 / LC-MS/MS	mg/capsule	208,8 ± 20,9

1) Capsule weight declared by the Client: 800 mg.

Test: Choline was performed in laboratory with an accreditation number 581

Test report authorized by:
ID: 548, Authorization Section Manager, Authorization Section

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* Test method accredited

Test performed by external provider

THE END OF THE REPORT



TEST REPORT NO 619435/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028	
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619435/26/GDY Sample received from the Client	
Start of analysis	07.07.2026		
End of analysis	15.07.2026		
Test report date	15.07.2026		

Test Method	Unit	Result
* Presence of Escherichia coli in 1 g PN-ISO 7251:2006	in 1 g	Not detected
* 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 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
* Number of Listeria monocytogenes at 37°C PN-EN ISO 11290-2:2017-07	cfu/g	<1,0x10 ¹

Test report authorized by:
ID: 343, Analysis Expert, Microbiology Laboratory

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Laboratory address:
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Test performed by external provider

THE END OF THE REPORT



TEST REPORT NO 620451/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620451/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	14.07.2026	
Test report date	14.07.2026	

Test Method	Unit	Result
* Vitamin B1 (thiamine) and B2 (ryboflavin) ¹⁾ PN-EN 14122:2014-07; PN-EN 14152:2014-07		
Vitamin B1 (thiamine)	mg/capsule	0,875 ± 0,175
Vitamin B ₂ (ryboflavin)	mg/capsule	1,50 ± 0,30

¹⁾ Capsule weight declared by the Client: 375 mg.

Test report authorized by:
ID: 346, Analysis Expert, Authorization Section

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Chwaszczyńska 180, 81-571 Gdynia

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Sample reception date	08.07.2026	Sample status: no objections Sample number: 620453/26/GDY Sample received from the Client
Start of analysis	13.07.2026	
End of analysis	14.07.2026	
Test report date	14.07.2026	

Test Method	Unit	Result
* Vitamin B ₅ (pantothenic acid) ^{1) 2)} PB-325 ed. 3 of 15.04.2025		
Vitamin B ₅ (pantothenic acid)	mg/capsule	5,54 ± 1,11

1) Capsule weight declared by the Client: 375 mg.

2) Specificity: pantothenic acid, sodium pantothenate, calcium pantothenate. No cross reactivity.

Test report authorized by:
ID: 346, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

The results refer only to the samples received and tested. The laboratory assumes no responsibility for the method of sample collection, the conditions of their transport, or for the information provided by the Client, including data that may affect the validity of the test results. 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 no criterion for a given test or tests in the analyzed matrix has been specified in the reference document indicated by the Client, confirmation of compliance is not possible. If the "result" column 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 method. If an expanded measurement uncertainty is given for such a test outcome, it relates only to the lower or upper limit of the measuring range of the method, respectively. In the case where the Laboratory base on the obtained test outcome, "statement of conformity" column presents an opinion and interpretation. 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

THE END OF THE REPORT



TEST REPORT NO 619439/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619439/26/GDY Sample received from the Client
Start of analysis	11.07.2026	
End of analysis	13.07.2026	
Test report date	13.07.2026	

Test Method	Unit	Result
* Vitamin E (α-tocopherol) ¹⁾ PB-40/HPLC ed. III of 28.02.2009		
Vitamin E (α-tocopherol)	mg/capsule	13,4 ± 1,9
Vitamin E (α-tocopherol) as tocopherol acetate	mg/capsule	14,8 ± 2,1

¹⁾ Capsule weight declared by the Client: 800 mg.

Test report authorized by:
ID: 795, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

The results refer only to the samples received and tested. The laboratory assumes no responsibility for the method of sample collection, the conditions of their transport, or for the information provided by the Client, including data that may affect the validity of the test results. 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 no criterion for a given test or tests in the analyzed matrix has been specified in the reference document indicated by the Client, confirmation of compliance is not possible. If the "result" column 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 method. If an expanded measurement uncertainty is given for such a test outcome, it relates only to the lower or upper limit of the measuring range of the method, respectively. In the case where the Laboratory base on the obtained test outcome, "statement of conformity" column presents an opinion and interpretation. 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.

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Test performed by external provider

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TEST REPORT NO 619438/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028	
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619438/26/GDY Sample received from the Client	
Start of analysis	11.07.2026		
End of analysis	13.07.2026		
Test report date	13.07.2026		

Test Method	Unit	Result
* Vitamin A ¹⁾ PB-40/HPLC ed. III of 28.02.2009		
Vitamin A (retinol)	µg/capsule	378 ± 57
Vitamin A (retinol) as retinyl acetate	µg/capsule	434 ± 65

1) Capsule weight declared by the Client: 800 mg.

Test report authorized by:
ID: 795, Analysis Expert, Authorization Section

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Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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Test performed by external provider

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TEST REPORT NO 620455/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620455/26/GDY Sample received from the Client
Start of analysis	09.07.2026	
End of analysis	13.07.2026	
Test report date	13.07.2026	

Test Method	Unit	Result
* Vitamin B7 (biotin) ^{1) 2) 3)} PB-326 ed. 3 of 15.04.2025	µg/capsule	48,4 ± 9,7

1) Masa kapsułki deklarowana przez Zlecniodawcę: 375 mg.

2) Capsule weight declared by the Client: 375 mg.

3) Specificity: D-biotin. No cross reactivity.

Test report authorized by:
ID: 346, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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* Test method accredited

Test performed by external provider

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TEST REPORT NO 620448/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620448/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	13.07.2026	
Test report date	13.07.2026	

Test Method	Unit	Result
* Zinc (Zn) ¹⁾ PB-36/ICP ed. 9 of 06.10.2025	mg/capsule	11,2 ± 2,8

1) Capsule weight declared by the Client: 375 mg.

Test report authorized by:
ID: 379, Senior Analysis Specialist, 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

The results refer only to the samples received and tested. The laboratory assumes no responsibility for the method of sample collection, the conditions of their transport, or for the information provided by the Client, including data that may affect the validity of the test results. 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 no criterion for a given test or tests in the analyzed matrix has been specified in the reference document indicated by the Client, confirmation of compliance is not possible. If the "result" column 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 method. If an expanded measurement uncertainty is given for such a test outcome, it relates only to the lower or upper limit of the measuring range of the method, respectively. In the case where the Laboratory base on the obtained test outcome, "statement of conformity" column presents an opinion and interpretation. 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

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TEST REPORT NO 620450/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620450/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	13.07.2026	
Test report date	13.07.2026	

Test Method	Unit	Result
* Copper (Cu) ¹⁾ PB-36/ICP ed. 9 of 06.10.2025	mg/capsule	0,48 ± 0,12

1) Capsule weight declared by the Client: 375 mg.

Test report authorized by:
ID: 379, Senior Analysis Specialist, 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

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Test performed by external provider

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TEST REPORT NO 620449/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620449/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	13.07.2026	
Test report date	13.07.2026	

Test Method	Unit	Result
Selenium (Se) ^{1) 2)} PB-223/ICP ed. 5 of 12.08.2025	µg/capsule	103 ± 28

1) Capsule weight declared by the Client: 375 mg.

2) Result outside the scope of accreditation.

Test report authorized by:
ID: 379, Senior Analysis Specialist, 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

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* Test method accredited

Test performed by external provider

THE END OF THE REPORT



TEST REPORT NO 620452/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- transparent capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 620452/26/GDY Sample received from the Client
Start of analysis	09.07.2026	
End of analysis	10.07.2026	
Test report date	10.07.2026	

Test Method	Unit	Result
* Vitamin B3 (niacin) ¹⁾ EN 15652:2009	mg/capsule	17,1 ± 3,4 3,4

1) Capsule weight declared by the Client: 375 mg.

Test report authorized by:
ID: 91, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

The results refer only to the samples received and tested. The laboratory assumes no responsibility for the method of sample collection, the conditions of their transport, or for the information provided by the Client, including data that may affect the validity of the test results. 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 no criterion for a given test or tests in the analyzed matrix has been specified in the reference document indicated by the Client, confirmation of compliance is not possible. If the "result" column 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 method. If an expanded measurement uncertainty is given for such a test outcome, it relates only to the lower or upper limit of the measuring range of the method, respectively. In the case where the Laboratory base on the obtained test outcome, "statement of conformity" column presents an opinion and interpretation. 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

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TEST REPORT NO 619441/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Mama Nadarra, 120 capsules- purple capsules Batch: MANAD/23062026 Production date: 23.06.2026 Expiry date: 23.06.2028
Sample reception date	07.07.2026	Sample status: no objections Sample number: 619441/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	10.07.2026	
Test report date	10.07.2026	

Test Method	Unit	Result
* Vitamin K2 (MK-4 and MK-7) ^{1) 2) 3)} PB-584 ed. 2 of 28.06.2024		
Menaquinone 4	µg/dose	< 68,0
Menaquinone 7	µg/dose	45,3 ± 11,3

- 1) Dose declared by the Client: 1 capsule.
2) Capsule filling weight declared by the Client: 680 mg.
3) The test was performed in capsule fill.

Test report authorized by:
ID: 91, Analysis Expert, Authorization Section

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

Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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