TEST REPORT NO 722761/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593807/24/GDY

Sample reception date:	24.09.2024
Start of analysis	27.09.2024
End of analysis	30.09.2024
Test report date	27.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Zinc (Zn) [1][2][3] PB-36/ICP ed. 8 of 29.12.2022	mg/capsule	8,51 ± 2,13	10 (+45%/-20%)	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 94, 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

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

THE END OF THE REPORT

TEST REPORT NO 722762/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593808/24/GDY

Sample reception date:	24.09.2024
Start of analysis	27.09.2024
End of analysis	01.10.2024
Test report date	27.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
Selenium (Se) [1][2][3][4] PB-223/ICP ed. 4 of 29.12.2022	µg/capsule	93,4 ± 25,2	100 (+45%/-20%)	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] The result is outside the scope of the accreditation. The scope of accreditation (0,025 - 1,25) mg/kg.

[3] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[4] Client specification.

Authorized by:
ID: 94, 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

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

THE END OF THE REPORT

TEST REPORT NO 722763/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593809/24/GDY

Sample reception date:	24.09.2024
Start of analysis	27.09.2024
End of analysis	30.09.2024
Test report date	27.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Copper (Cu) [1][2][3] PB-36/ICP ed. 8 of 29.12.2022	mg/capsule	0,49 ± 0,24	0,5 (-20%/+45%)	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 295, 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

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

THE END OF THE REPORT

TEST REPORT NO 722764/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593811/24/GDY

Sample reception date:	24.09.2024
Start of analysis	01.10.2024
End of analysis	02.10.2024
Test report date	02.10.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B3 (niacin) [1][2][3] EN 15652:2009	mg/capsule	16,0 ± 3,2	16 +50%/-20%	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 228, Vitamins Testing Laboratory Manager, Vitamin Analysis 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. 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

THE END OF THE REPORT

TEST REPORT NO 722765/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593812/24/GDY

Sample reception date:	24.09.2024
Start of analysis	01.10.2024
End of analysis	02.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B5 (pantothenic acid) [1][2][3][4] PB-325 ed. 2 of 05.09.2022				
Vitamin B$_5$ (pantothenic acid)	mg/capsule	6,20 ± 1,24	6 +50%/-20%	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

[4] Specificity: pantothenic acid, sodium pantothenate, calcium pantothenate. No cross reactivity.

Authorized by:
ID: 758, Analysis Expert, Vitamin Analysis 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. 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

THE END OF THE REPORT

TEST REPORT NO 722766/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593813/24/GDY

Sample reception date:	24.09.2024
Start of analysis	03.10.2024
End of analysis	08.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B6 [1][2][3] PN-EN 14164:2014-08				
Vitamin B$_6$ (pyridoxine hydrochloride)	mg/capsule	3,24 ± 0,65	-	-
Vitamin B6 (pyridoxine)	mg/capsule	2,66 ± 0,53	3 +50%/-20%	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 758, Analysis Expert, Vitamin Analysis 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. 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

THE END OF THE REPORT

TEST REPORT NO 722767/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593814/24/GDY

Sample reception date:	24.09.2024
Start of analysis	01.10.2024
End of analysis	03.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B7 (biotin) [1][2][3][4] PB-326 ed. 2 of 05.09.2022	µg/capsule	50,0 ± 10,0	50 +50%/-20%	Pass

[1] Capsule weight declared by the Client: 375 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

[4] Specificity: D-biotin. No cross reactivity.

Authorized by:
ID: 228, Vitamins Testing Laboratory Manager, Vitamin Analysis 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. 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

THE END OF THE REPORT

TEST REPORT NO 722768/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 593815/24/GDY

Sample reception date:	24.09.2024
Start of analysis	07.10.2024
End of analysis	07.11.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* # 5-methyltetrahydrofolate (5-MTHF) [1][2] USP 41/ NF 36 method 1 mod. / LC-DAD				
Methyltetrahydrofolate (5-MTHF)	µg/capsule	< 37,5	-	-

[1] Capsule weight declared by the Client: 375 mg.

[2] The symbol "<" means below the limit of quantification of the analytical method.

Test: 5-methyltetrahydrofolate (5-MTHF) was performed in laboratory with an accreditation number 581

Authorized by:
Subcontracted test results are authorised by persons authorised by the external provider.
The test report bears the certified electronic seal of J.S. Hamilton Poland Sp. z o.o.

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

THE END OF THE REPORT

TEST REPORT NO 722769/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number:659128/24/GDY

Sample reception date:	24.10.2024
Start of analysis	30.10.2024
End of analysis	12.11.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* # Methylcobalamin (sum of methyl and hydroxycobalamin) [1] USP 44 NF 39 mod. / LC-UV/DAD	µg/capsule	81,75 ± 11,45	-	-

[1] Capsule weight declared by the Client: 375 mg.

Test: Methylcobalamin (sum of methyl and hydroxycobalamin) was performed in laboratory with an accreditation number 581

Authorized by:
Subcontracted test results are authorised by persons authorised by the external provider.
The test report bears the certified electronic seal of J.S. Hamilton Poland Sp. z o.o.

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

THE END OF THE REPORT

TEST REPORT NO 722770/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 659129/24/GDY

Sample reception date:	24.10.2024
Start of analysis	30.10.2024
End of analysis	08.11.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B1 [1][2][3] PN-EN 14122:2014-07				
Vitamin B1 (thiamine)	mg/capsule filling	1,56 ± 0,31	1,1 +50%/-20%	Pass
Vitamin B1 (thiamine hydrochloride)	mg/capsule filling	1,98 ± 0,40	-	-

[1] Capsule filling weight declared by the Client: 300 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 758, Analysis Expert, Vitamin Analysis Laboratory

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Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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TEST REPORT NO 722771/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - transparent capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 659130/24/GDY

Sample reception date:	24.10.2024
Start of analysis	30.10.2024
End of analysis	08.11.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin B2 (ryboflavin) [1][2][3] PN-EN 14152:2014-07				
Vitamin B$_2$ (ryboflavin)	mg/capsule filling	2,49 ± 0,50	1,4 +50%/-20%	Fail

[1] Capsule filling weight declared by the Client: 300 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

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ID: 758, Analysis Expert, Vitamin Analysis Laboratory

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Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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TEST REPORT NO 722774/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - purple capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 594368/24/GDY

Sample reception date:	24.09.2024
Start of analysis	27.09.2024
End of analysis	03.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* # Choline [1] AOAC Vol 91,1 2008 / LC-MS/MS	mg/capsule	210 ± 42	-	-

[1] Capsule weight declared by the Client: 800 mg.

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

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THE END OF THE REPORT

TEST REPORT NO 722775/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - purple capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 594369/24/GDY

Sample reception date:	24.09.2024
Start of analysis	03.10.2024
End of analysis	03.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin A [1][2][3] PB-40/HPLC ed. III of 28.02.2009				
Vitamin A (retinol)	µg/capsule	244 ± 37	300 +50%/-20%	Pass
Vitamin A (retinol) as retinyl acetate	µg/capsule	280 ± 42	-	-

[1] Capsule weight declared by the Client: 800 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

Authorized by:
ID: 127, Analysis Expert, Vitamin Analysis Laboratory

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Laboratory address:
Chwaszczyńska 180, 81-571 Gdynia

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TEST REPORT NO 722776/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - purple capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 594370/24/GDY

Sample reception date:	24.09.2024
Start of analysis	02.10.2024
End of analysis	03.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin E (α-tocopherol) [1][2][3] PB-40/HPLC ed. III of 28.02.2009				
Vitamin E (α-tocopherol)	mg/capsule	12,2 ± 1,7	12 +50%/20%	Pass
Vitamin E (α-tocopherol) as tocopherol acetate	mg/capsule	13,4 ± 1,9	-	-

[1] Capsule weight declared by the Client: 800 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

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Laboratory address:
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TEST REPORT NO 722777/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - purple capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 594371/24/GDY

Sample reception date:	24.09.2024
Start of analysis	02.10.2024
End of analysis	07.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin D3 [1][2][3] PN-EN 12821:2009				
Vitamin D3 (cholecalciferol)	µg/capsule	51,7 ± 7,7	50 +50%/-20%	Pass

[1] Capsule weight declared by the Client: 800 mg.

[2] Guidance Document for competent authorities for the control of compliance with EU legislation on: Regulation (EU) No 1169/2011 of the European Parliament and of the Council of 25 October 2011 on the provision of food information to consumers, amending Regulations (EC) No 1924/2006 and (EC) No 1925/2006 of the European Parliament and of the Council, and repealing Commission Directive 87/250/EEC, Council Directive 90/496/EEC, Commission Directive 1999/10/EC, Directive 2000/13/EC of the European Parliament and of the Council, Commission Directives 2002/67/EC and 2008/5/EC and Commission Regulation (EC) No 608/2004 and Council Directive 90/496/EEC of 24 September 1990 on nutrition labelling of foodstuffs and Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements with regard to the setting of tolerances for nutrient values declared on a label, December 2012. Table 2.

[3] Client specification.

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ID: 127, Analysis Expert, Vitamin Analysis Laboratory

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THE END OF THE REPORT

TEST REPORT NO 722778/24/GDY

Client
NÀDARRA GmbH
Harvestehuder Weg 48
20149 Hamburg

Sample (according to declaration of Client)
Sample description: Mama Nadarra, 120 capsules - purple capsules
Batch: MND/05092024
Production date: 05.09.2024
Expiry date: 05.09.2026

Sample number: 594372/24/GDY

Sample reception date:	24.09.2024
Start of analysis	27.09.2024
End of analysis	16.10.2024
Test report date	28.11.2024

Sample status: no objections

Sample received from the Client

Test Method	Unit	Result	Criteria	Statement of conformity
* # Iodine [1] VDLUFA VII, 2.2.2.3; 2011				
Iodine	µg/capsule	132 ± 35	-	-
Iodine calculated as potassium iodate	µg/capsule	222 ± 59	-	-

[1] The iodine content in the capsules is very inhomogeneously distributed. Individual results: 70,8 mg/kg, 34,8 mg/kg, 57,8 mg/kg, 496 mg/kg.

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

Authorized by:
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