

## TEST REPORT NO 403337/24/GDY

		Sample (according to declaration of Client) Sample description: Vitamin B Complex, 90 Capsules, Nadara Batch: VBNK/07052024 Production date: 07.05.2024 Expiry date: 07.05.2026
Sample reception date: 25.07.2024		Sample status: no objections
Start of analysis 29.07.2024		
End of analysis 20.08.2024		Sample received from the Client
Test report date 20.08.2024		

Test Method	Unit	Result	Criteria	Statement of conformity
# Methylcobalamin (sum of methyl and hydroxycobalamin) <sup>12</sup> USP 44/NF 39 mod. / LC-CV/DAD	µg/capsule	8.9 ± 1.2	8-15	Pass

1) Packaging label.  
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 Regulation (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.

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 " $\times$ " or " $\times$ ", 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. document. The test report is not valid if it is used for the purposes of the PCA accreditation symbol AB 079 by customers, subcontractors, external service providers and other third parties. A limited access to the data issued in the original J.S. Hamilton Poland Sp. z o.o. document is granted to the PCA. 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

Page 1 / 1

J.S. HAMILTON POLAND Sp. z o.o.  
TESTING LABORATORY

Chwaszczyńska 180, 81-571 Gdynia, Poland tel. +48 58 766 99 00



## TEST REPORT NO 289410/24/GDY

		Sample (according to declaration of Client) Sample description: Vitamin B Komplex Nadara, 90 capsules Batch: VBNK/07052024 Production date: 07.05.2024 Expiry date: 07.05.2026
Sample reception date: 17.05.2024		Sample status: no objections
Start of analysis 21.05.2024		
End of analysis 23.05.2024		Sample received from the Client
Test report date 23.05.2024		

Test Method	Unit	Result	Criteria	Statement of conformity
Vitamin B7 (biotin) <sup>12</sup> 20/22 ed. 2 of 05.09.2022	µg/capsule	62,4 ± 12,5	50 ±50/-20%	Pass

1) Capsule weight declared by the Client: 495 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: 293, 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 " $\times$ " or " $\times$ ", 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. document. The test report is not valid if it is used for the purposes of the PCA accreditation symbol AB 079 by customers, subcontractors, external service providers and other third parties. A limited access to the data issued in the original J.S. Hamilton Poland Sp. z o.o. document is granted to the PCA. 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

Page 1 / 1

J.S. HAMILTON POLAND Sp. z o.o.  
TESTING LABORATORY

Chwaszczyńska 180, 81-571 Gdynia, Poland tel. +48 58 766 99 00



## TEST REPORT NO 289408/24/GDY

		Sample (according to declaration of Client) Sample description: Vitamin B Komplex Nadara, 90 capsules Batch: VBNK/07052024 Production date: 07.05.2024 Expiry date: 07.05.2026
Sample reception date: 17.05.2024		Sample status: no objections
Start of analysis 22.05.2024		
End of analysis 23.05.2024		Sample received from the Client
Test report date 23.05.2024		

Test Method	Unit	Result	Criteria	Statement of conformity
Vitamin B5 (pantothenic acid) <sup>12</sup> 20/22 ed. 2 of 05.09.2022	mg/capsule	5.78 ± 1.16	1.4 ±50/-20%	Pass

1) Capsule weight declared by the Client: 495 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: 293, 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 " $\times$ " or " $\times$ ", 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. document. The test report is not valid if it is used for the purposes of the PCA accreditation symbol AB 079 by customers, subcontractors, external service providers and other third parties. A limited access to the data issued in the original J.S. Hamilton Poland Sp. z o.o. document is granted to the PCA. 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

Page 1 / 1

J.S. HAMILTON POLAND Sp. z o.o.  
TESTING LABORATORY

Chwaszczyńska 180, 81-571 Gdynia, Poland tel. +48 58 766 99 00



## TEST REPORT NO 289412/24/GDY

		Sample (according to declaration of Client) Sample description: Vitamin B Komplex Nadara, 90 capsules Batch: VBNK/07052024 Production date: 07.05.2024 Expiry date: 07.05.2026
Sample reception date: 17.05.2024		Sample status: no objections
Start of analysis 22.05.2024		
End of analysis 23.05.2024		Sample received from the Client
Test report date 23.05.2024		

Test Method	Unit	Result	Criteria	Statement of conformity
Vitamin B1 (riboflavin) <sup>12</sup> 20/22 ed. 2 of 05.09.2024	mg/capsule	0.98 ± 0.197	1.1 ±50/-20%	Pass

1) Capsule weight declared by the Client: 495 mg.  
2) Guidance Document for competent