

TEST REPORT NO 602313/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Vitalität Nadarra, 60 capsules Batch: VINAD/22062026 Production date: 22.06.2026 Expiry date: 22.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 602313/26/GDY Sample received from the Client
Start of analysis	27.07.2026	
End of analysis	13.08.2026	
Test report date	13.08.2026	

Test Method	Unit	Result
# Resveratrol ¹⁾ SOP 3.1.139	mg/capsule filling	50 ± 2,5

¹⁾ Capsule filling weight declared by the Client: 660 mg.

Test: Resveratrol was performed in laboratory Certified Laboratories, Inc. Tustin United States of America

Test report authorized by:
ID: 448, Analysis Expert, Authorization Section

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 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 602311/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Vitalität Nadarra, 60 capsules Batch: VINAD/22062026 Production date: 22.06.2026 Expiry date: 22.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 602311/26/GDY Sample received from the Client
Start of analysis	10.07.2026	
End of analysis	07.08.2026	
Test report date	07.08.2026	

Test Method	Unit	Result
# Quercetin ^{1) 2)} LC-MS/MS after extraction		
Quercitin	mg/dose	124,8

1) Capsule weight declared by the Client: 780 mg.

2) Dose declared by the Client: 2 capsules.

Test: Quercetin was performed in laboratory PiCA Prüfinstitut Chemische Analytik GmbH Berlin Germany

Test report authorized by:
 ID: 434, Analysis Expert, 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 602310/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Vitalität Nadarra, 60 capsules Batch: VINAD/22062026 Production date: 22.06.2026 Expiry date: 22.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 602310/26/GDY Sample received from the Client
Start of analysis	24.07.2026	
End of analysis	06.08.2026	
Test report date	06.08.2026	

Test Method	Unit	Result
# Apigenin SOP 3.1.401 (HPLC)		
Apigenin	%	1,74 ± 0,09
	mg/kg	17400 ± 870

Test: Apigenin was performed in laboratory Certified Laboratories, Inc. Tustin United States of America

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.

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 602312/26/GDY

Client NADARRA GmbH Harvestehuder Weg 48 20149 Hamburg		Sample (according to declaration of Client) Sample description: Vitalität Nadarra, 60 capsules Batch: VINAD/22062026 Production date: 22.06.2026 Expiry date: 22.06.2028
Sample reception date	08.07.2026	Sample status: no objections Sample number: 602312/26/GDY Sample received from the Client
Start of analysis	27.07.2026	
End of analysis	29.07.2026	
Test report date	29.07.2026	

Test Method	Unit	Result
* # Coenzyme Q10 ^{1) 2)} SOP O-24	mg/dose	112 ± 22

1) Dose declared by the Client: 2 capsules.

2) Capsule weight declared by the Client: 780 mg.

Test: Coenzyme Q10 was performed in laboratory with an accreditation number L 1162

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.

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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