



TEST REPORT NO 726648/24/GDY

Client		Sample (according to declaration of Client)		
NÀDARRA GmbH		Sample description: Vitalität Nadarra, 60 capsules Batch: VN/04092024		
Harvestehuder Weg 48				
20149 Hamburg		Production date: 04.09.2024		
		Expiry date: 04.09.2026		
		Sample No. 565795/24/GDY		
Sample reception date:	13.09.2024	Sample status: no objections		
Start of analysis	23.09.2024			
End of analysis	11.10.2024			
Test report date	21.11.2024	Sample received from the Client		

Test Method	Unit	Result	Criteria	Statement of conformity
# Trans-resveratrol 1) SOP 3.1.58	mg/capsule	35,94 ± 1,80	(#X	n+1
	10.000			

Capsule weight declared by the Client: 800 mg.

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

Authorized by: Subcontracted test results are authorised by persons authorised by the external provider.

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

THE END OF THE REPORT

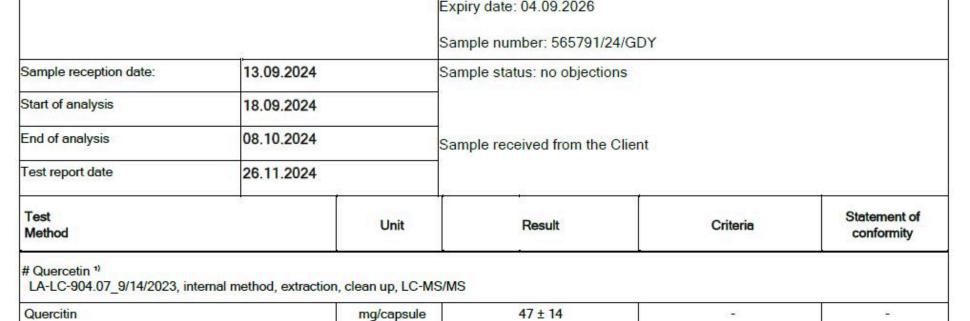
Page 1 / 1

J.S. HAMILTON POLAND Sp. z o.o. **TESTING LABORATORY** Chwaszczyńska 180, 81-571 Gdynia, Poland tel. +48 58 766 99 00

HAMILTON ANALYST

Sample (according to declaration of Client)

Sample description: Vitalität Nadarra, 60 capsules



TEST REPORT NO 722752/24/GDY

Batch: VN/04092024

Production date: 04.09.2024

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. Is limited solely to the data issued in its original. J.S.

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

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.

Authorized by:

Client

NÀDARRA GmbH

20149 Hamburg

Harvestehuder Weg 48

* Test method accredited # Test performed by external provider

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.

Tablet weight declared by the Client: 800 mg.

THE END OF THE REPORT

Client

NÀDARRA GmbH

20149 Hamburg

Test report date

Coenzyme Q10 1)

* Test method accredited

Client

NÀDARRA GmbH

20149 Hamburg

Harvestehuder Weg 48

Sample reception date:

Nicotinamide adenine dinucleotide (NAD+) 1)

Nicotinamide adenine dinucleotide (NAD+)

Start of analysis

End of analysis

Test report date

Test

Method

Authorized by:

* Test method accredited

Test performed by external provider

Test

Method

Harvestehuder Weg 48

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

Page 1 / 1

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

HAMILTON

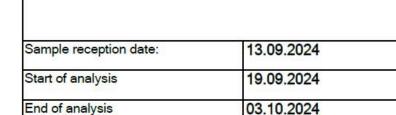
TEST REPORT NO 722754/24/GDY

Batch: VN/04092024

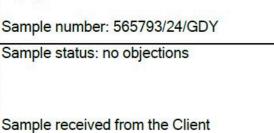
Production date: 04.09.2024 Expiry date: 04.09.2026

Result

 $46,11 \pm 4,6$



26.11.2024



Sample (according to declaration of Client)

Sample description: Vitalität Nadarra, 60 capsules

ANALYST

Criteria

Statement of

conformity

Statement of

conformity

SOP 0-24 Capsule weight declared by the Client: 800 mg. Test: Coenzyme Q10 was performed in laboratory with an accreditation number L 1162 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

THE END OF THE REPORT

Unit

mg/capsule

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 performed by external provider

HAMILTON

TEST REPORT NO 722755/24/GDY

Batch: VN/04092024

Production date: 04.09.2024 Expiry date: 04.09.2026

Sample status: no objections

Sample number: 565794/24/GDY

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

25.10.2024 26.11.2024

13.09.2024

23.09.2024

Sample received from the Client Unit Result

11,75 ± 0,59

Sample (according to declaration of Client)

Sample description: Vitalität Nadarra, 60 capsules

ANALYST

Criteria

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

THE END OF THE REPORT

Capsule weight declared by the Client: 800 mg. Test: Nicotinamide adenine dinucleotide (NAD+) was performed in laboratory Certified Laboratories, Inc. Tustin United States of America

mg/capsule

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 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 "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance decision rule in accordance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported. If the "results of the acceptance with ILAC-G8:09/2019, unless otherwise reported with ILAC-G8:09/2019, unless otherwise reported with ILAC-G8:09/2019, unless otherwise reported with ILAC-G8:09/201 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.

Page 1 / 1