

**TEST REPORT NO 250624/26/GDY/Z1**  
**Replaces test report no. 250624/26/GDY of 27.03.2026**

<b>Client</b> <b>OstroVit Sp. z o.o.</b> Sitarska 16 18-300 Zambrów		<b>Sample (according to declaration of Client)</b> Sample description: OstroVit Vitamin K2 200 Natto MK-7 90 tabs Batch: TAB/0001492 Expiry date: 02.03.2029
Sample reception date	<b>19.03.2026</b>	Sample status: no objections Sample number: 250624/26/GDY  Sample received from the Client
Start of analysis	<b>27.03.2026</b>	
End of analysis	<b>27.03.2026</b>	
Test report date	<b>03.04.2026</b>	

Test Method	Unit	Result	Criteria	Statement of conformity
Total menaquinone-7 content (as sum of cis and trans isomers) <sup>1) 2) 3) 4)</sup> Calculated	µg/dose	222 ± 56	200 (+50%/-20%)	Pass
* Vitamin K2 (all-trans menaquinone-7) <sup>1) 2) 3) 4)</sup> PB-584 ed. 2 of 28.06.2024				
all-trans Menaquinone-7	µg/dose	222 ± 55	200 (+50%/-20%)	Pass
Menaquinone-7 isomeric purity (all-trans isomer content)	%	99,7	-	-

- 1) Dose declared by the Client: 1 tablet.
- 2) Tablet weight declared by the Client: 100 mg.
- 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 requirements.

Vitamin K2 (all-trans menaquinone-7): Identification of the change: addition of the statement of conformity

Test report authorized by:  
 ID: 448, Analysis Expert, Authorization Section  
 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. 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 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](http://www.hamilton.com.pl).

\* Test method accredited  
 # Test performed by external provider

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THE END OF THE REPORT