



TEST REPORT NO 683539/24/GDY

Client OstroVit Sp. z o.o. Sitarska 16 18-300 Zambrów		Sample (according to declaration of Client) Sample description: OstroVit Pharma D3 4000 K2 MK-7 (tablets) Batch: 9PD4K014 Expiry date: 18.10.2027
Sample reception date:	31.10.2024	Sample status: no objections Sample received from the Client
Start of analysis	07.11.2024	
End of analysis	08.11.2024	
Test report date	08.11.2024	

Test Method	Unit	Result	Criteria	Statement of conformity
* Vitamin D3 ^{1) 2) 3)} PN-EN 12821:2009				
Vitamin D3 (cholecalciferol)	IU/dose	4620 ± 690	4000 +50%/-20%	Pass

- 1) Dose 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

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

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