



## TEST REPORT NO 151096/26/GDY

|                                                                       |            |                                                                                                                                                  |  |
|-----------------------------------------------------------------------|------------|--------------------------------------------------------------------------------------------------------------------------------------------------|--|
| Client<br><b>OstroVit Sp. z o.o.</b><br>Sitarska 16<br>18-300 Zambrów |            | Sample (according to declaration of Client)<br>Sample description: OstroVit MGZB ULTRA 120 tabs<br>Batch: TAB/0000602<br>Expiry date: 09.09.2028 |  |
| Sample reception date                                                 | 21.02.2026 | Sample status: no objections<br>Sample number: 151096/26/GDY<br><br>Sample received from the Client                                              |  |
| Start of analysis                                                     | 26.02.2026 |                                                                                                                                                  |  |
| End of analysis                                                       | 27.02.2026 |                                                                                                                                                  |  |
| Test report date                                                      | 27.02.2026 |                                                                                                                                                  |  |

| Test Method                                                         | Unit    | Result     | Criteria       | Statement of conformity |
|---------------------------------------------------------------------|---------|------------|----------------|-------------------------|
| * Zinc (Zn) <sup>1) 2) 3) 4)</sup><br>PB-36/ICP ed. 9 of 06.10.2025 | mg/dose | 17,3 ± 4,3 | 15 (+45%/-20%) | Pass                    |

- 1) Dose declared by the Client: 3 tablets.
- 2) Tablet weight declared by the Client: 1475 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.

Test report authorized by:  
ID: 371, Senior Analysis Specialist, Spectrometry 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

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\* Test method accredited  
# Test performed by external provider

THE END OF THE REPORT