



TEST REPORT NO 764622/25/GDY

Client OstroVit Sp. z o.o. Sitarska 16 18-300 Zambrów		Sample (according to declaration of Client) Sample description: OstroVit L-Carnitine 1250 mg Batch: PPK/0000601 Expiry date: 27.09.2027	
Sample reception date:	04.10.2025	Sample status: no objections Sample received from the Client	
Start of analysis	04.10.2025		
End of analysis	10.10.2025		
Test report date	10.10.2025		

Test Method	Unit	Result	Criteria	Statement of conformity
* Aerobic colony count at 30°C ²⁾ PN-EN ISO 4833-1:2013-12; PN-EN ISO 4833-1:2013-12/A1:2022-06				
Aerobic colony count ¹⁾	cfu/g	8,0x10 ¹ [4,0x10 ¹ ; 1,5x10 ²]	≤1,0x10 ⁴	Pass
Aerobic colony count	cfu/g	8,0x10 ¹ [4,0x10 ¹ ; 1,5x10 ²]	-	-
* Number of yeasts and moulds at 25°C PN-ISO 21527-2:2009 (withdrawn)				
Number of yeasts and moulds ¹⁾	cfu/g	<1,0x10 ¹	≤1,0x10 ³	Pass
Number of yeasts and moulds	cfu/g	<1,0x10 ¹	-	-
Number of yeasts	cfu/g	<1,0x10 ¹	-	-
Number of moulds	cfu/g	<1,0x10 ¹	-	-
* Presence of Escherichia coli in 10 g PN-ISO 7251:2006				
Presence of Escherichia coli ¹⁾	in 10 g	Not detected	Absent	Pass
Presence of Escherichia coli	in 10 g	Not detected	-	-
* Presence of coagulase-positive staphylococci (Staphylococcus aureus and other species) in 1 g PN-EN ISO 6888-3:2004; PN-EN ISO 6888-3:2004/AC:2005				
Presence of coagulase-positive staphylococci (Staphylococcus aureus and other species) ¹⁾	in 1 g	Not detected	Absent	Pass
Presence of coagulase-positive staphylococci (Staphylococcus aureus and other species)	in 1 g	Not detected	-	-
* Presence of Listeria monocytogenes in 25 g PN-EN ISO 11290-1:2017-07				
Presence of Listeria monocytogenes ¹⁾	in 25 g	Not detected	Absent	Pass
Presence of Listeria monocytogenes	in 25 g	Not detected	-	-
* Presence of Salmonella spp. in 10 g PN-EN ISO 6579-1:2017-04; PN-EN ISO 6579-1:2017-04/A1:2020-09				
Presence of Salmonella spp. ¹⁾	in 10 g	Not detected	Absent	Pass
Presence of Salmonella spp.	in 10 g	Not detected	-	-



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- 1) Client specification.
- 2) The reported expanded uncertainty of measurement has been estimated according to ISO 19036 and is based on the standard uncertainty multiplied by the coverage factor $k=2$ with a confidence level of 95%. The composite standard uncertainty was taken as equal to the standard deviation of within-laboratory reproducibility.

Authorized by:

ID: 106, Analysis Expert, Microbiology 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

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.

* Test method accredited

Test performed by external provider

THE END OF THE REPORT