



**PARTIAL TEST REPORT NO 623323/26/GDA/2**  
**Translation of the report NO 623323/26/GDA/1 of 13.07.2026**

|                                                                                             |                   |                                                                                                     |
|---------------------------------------------------------------------------------------------|-------------------|-----------------------------------------------------------------------------------------------------|
| Client<br><b>DNL Equity Partners Spółka z.o.o</b><br>ul. Świętojańska 87/3<br>81-389 Gdynia |                   | Sample (according to declaration of Client)<br>Sample description: Alchemy Sérum                    |
| Sample reception date                                                                       | <b>17.10.2024</b> | Sample status: no objections<br>Sample number: 648569/24/GDA<br><br>Sample received from the Client |
| Start of analysis                                                                           | <b>18.10.2024</b> |                                                                                                     |
| End of analysis                                                                             | <b>24.10.2024</b> |                                                                                                     |
| Test report date                                                                            | <b>13.07.2026</b> |                                                                                                     |

| Test Method                                                                                                              | Unit   | Result               |
|--------------------------------------------------------------------------------------------------------------------------|--------|----------------------|
| * Presence of Staphylococcus aureus in 1 g <sup>2)</sup><br>PN-EN ISO 22718:2016-01; PN-EN ISO 22718:2016-01/A1:2023-01  | in 1 g | Absent               |
| * Presence of Escherichia coli in 1 g <sup>1)</sup><br>PN-EN ISO 21150:2016-01; PN-EN ISO 21150:2016-01/A1:2023-03       | in 1 g | Absent               |
| * Presence of Pseudomonas aeruginosa in 1 g <sup>5)</sup><br>PN-EN ISO 22717:2016-01; PN-EN ISO 22717:2016-01/A1:2023-03 | in 1 g | Absent               |
| * Presence of Candida albicans in 1 g <sup>6)</sup><br>PN-EN ISO 18416:2016-01; PN-EN ISO 18416:2016-01/A1:2023-03       | in 1 g | Absent               |
| * Aerobic mesophilic bacteria at 32,5°C <sup>4)</sup><br>PN-EN ISO 21149:2017-07; PN-EN ISO 21149:2017-07/A1:2023-01     | cfu/g  | <1,0x10 <sup>1</sup> |
| * Number of yeasts and moulds at 25°C <sup>3)</sup><br>PN-EN ISO 16212:2017-08; PN-EN ISO 16212:2017-08/A1:2023-01       | cfu/g  | <1,0x10 <sup>1</sup> |

- 1) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
 Culture media: MacConkey Agar  
 Test strains: Escherichia coli ATCC 8739  
 Volume of calibrated suspension: 0.1 ml of the calibrated inoculum Nv - 1.0x10<sup>2</sup> - 5.0x10<sup>2</sup> cfu/g  
 Result of the neutralization: presence of microbial growth in the test mixture with the neutralizer and formulation with 0.1 ml inoculum Nv and absence of microbial growth in the test mixture with the neutralizer and formulation  
 The effectiveness of the neutralization has been confirmed.
- 2) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
 Culture media: Baird Parker Agar  
 Test strains: Staphylococcus aureus ATCC 6538  
 Volume of calibrated suspension: 0.1 ml of the calibrated inoculum Nv - 1.0x10<sup>2</sup> - 5.0x10<sup>2</sup> cfu/g  
 Result of the neutralization: presence of microbial growth in the test mixture with the neutralizer and formulation with 0.1 ml inoculum Nv and absence of microbial growth in the test mixture with the neutralizer and formulation  
 The effectiveness of the neutralization has been confirmed.



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- 3) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
Culture media: SDCA (Sabouraud dextrose Agar with chloramphenicol)  
Test strains: *Candida albicans* ATCC 10231  
Volume of calibrated suspension: 1 ml of the calibrated inoculum  $N_v - 1.0 \times 10^3 - 3.0 \times 10^3$  cfu/g  
Result of the neutralization:  $N_{vw} \geq 0,5 N_{vk}$ , where:  $N_{vw}$  - number of microorganisms present in the test mixture with the neutralizer and formulation and  $N_{vk}$  - number of microorganisms present in the test mixture with the neutralizer in the absence of the formulation  
The effectiveness of the neutralization has been confirmed.
- 4) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
Culture media: TSA (Tryptic Soy Agar)  
Test strains: *Staphylococcus aureus* ATCC 6538, *Pseudomonas aeruginosa* ATCC 9027  
Volume of calibrated suspension: 1 ml of the calibrated inoculum  $N_v - 1.0 \times 10^3 - 3.0 \times 10^3$  cfu/g  
Result of the neutralization:  $N_{vw} \geq 0,5 N_{vk}$ , where:  $N_{vw}$  - number of microorganisms present in the test mixture with the neutralizer and formulation and  $N_{vk}$  - number of microorganisms present in the test mixture with the neutralizer in the absence of the formulation  
The effectiveness of the neutralization has been confirmed.
- 5) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
Culture media: Cetrimide Agar  
Test strains: *Pseudomonas aeruginosa* ATCC 9027  
Volume of calibrated suspension: 0.1 ml of the calibrated inoculum  $N_v - 1.0 \times 10^2 - 5.0 \times 10^2$  cfu/g  
Result of the neutralization: presence of microbial growth in the test mixture with the neutralizer and formulation with 0.1 ml inoculum  $N_v$  and absence of microbial growth in the test mixture with the neutralizer and formulation  
The effectiveness of the neutralization has been confirmed.
- 6) Neutralizer: Eugon LT (1 g of the sample in 9 ml of the neutralizer)  
Culture media: SDCA (Sabouraud dextrose Agar with chloramphenicol)  
Test strains: *Candida albicans* ATCC 10231  
Volume of calibrated suspension: 0.1 ml of the calibrated inoculum  $N_v - 1.0 \times 10^2 - 5.0 \times 10^2$  cfu/g  
Result of the neutralization: presence of microbial growth in the test mixture with the neutralizer and formulation with 0.1 ml inoculum  $N_v$  and absence of microbial growth in the test mixture with the neutralizer and formulation  
The effectiveness of the neutralization has been confirmed.

Identification of the change: sample description

Test report authorized by:

ID: 175, Senior Analysis Specialist, Cosmetics Microbiology Laboratory

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

Laboratory address:

Goździków 1, 43-100 Tychy

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\* Test method accredited

# Test performed by external provider

THE END OF THE REPORT