



TEST REPORT NO 252540/23/GDY

Client SFD SPÓŁKA AKCYJNA GŁOGOWSKA 41 45315 OPOLE		Sample (according to declaration of Client) Sample description: LOCO ENERGY & SPEED 120 caps Batch: LC230403 Production date: 01.04.2023	
Sample reception date:	17.05.2023	Sample status: no objections	
Start of analysis	17.05.2023	Sample received from the Client	
End of analysis	02.06.2023		
Test report date	02.06.2023		
Test Method		Unit	Result
* Aerobic colony count at 30°C PN-EN ISO 4833-1:2013-12		cfu/g	<1,0x101
* Number of yeasts and moulds at 25 PN-ISO 21527-2:2009 (withdrawn)	5°C	· · · · · ·	
Number of yeasts		cfu/g	<1,0x101
Number of moulds		cfu/g	<1,0x101
* 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		in 1 g	Not detected
* Presence of Escherichia coli in 1 g PN-ISO 7251:2006		in 1 g	Not detected
* Presence of Salmonella spp. in 25 g PN-EN ISO 6579-1:2017-04; PN-EN ISO 6579-1:2017-04/A1:2020-09		in 25 g	Not detected
* Presence of Listeria monocytogenes in 25 g PN-EN ISO 11290-1:2017-07		in 25 g	Not detected
* Pyrrolizidine alkaloids ^{3) 4)} PB-498 ed. I of 23.05.2022			
Echimidine		μg/kg	< 5,0 (5,0 ± 1,8)
Echimidine N-oxide		μg/kg	< 5,0 (5,0 ± 1,8)
Echinatine N-oxide		μg/kg	< 5,0 (5,0 ± 1,8)
Europine		μg/kg	< 5,0 (5,0 ± 1,8)
Europine N-oxide		µg/kg	< 5,0 (5,0 ± 1,8)
Heliosupine		µg/kg	< 5,0 (5,0 ± 1,8)
Heliosupine N-oxide		µg/kg	< 5,0 (5,0 ± 1,8)
Heliotrine		µg/kg	< 5,0 (5,0 ± 1,8)
Heliotrine N-oxide		μg/kg	< 5,0 (5,0 ± 1,8)
Intermedine		μg/kg	< 5,0 (5,0 ± 1,8)
Intermedine-N-oxide (sum of intermedine-N-oxide and indicine-N-oxide as intermedine-N-oxide)		μg/kg	< 5,0 (5,0 ± 1,8)

Page 1/3





TEST REPORT NO 252540/23/GDY

Lasiocarpine	μg/kg	< 5,0 (5,0 ± 1,8)
Lasiocarpine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Lycopsamine (sum of lycopsamine, indicine and echinatine as lycopsamine)	μg/kg	< 5,0 (5,0 ± 1,8)
Lycopamine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Retrorsine (sum of retrorsine and usaramine as retrorsine)	μg/kg	< 5,0 (5,0 ± 1,8)
Retrorsine-N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Rinderine	μg/kg	< 5,0 (5,0 ± 1,8)
Rinderine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Senecionine	μg/kg	< 5,0 (5,0 ± 1,8)
Senecionine-N-oxide (sum of senecionine-N-oxide and integerrimine-N -oxide as senecionine-N-oxide)	μg/kg	< 5,0 (5,0 ± 1,8)
Seneciphylline (sum of seneciphylline and spartioidine as seneciphylline)	μg/kg	< 5,0 (5,0 ± 1,8)
Seneciphylline-N-oxide (sum of seneciphylline-N-oxide and spartioidine N-oxide as seneciphylline-N-oxide)	μg/kg	< 5,0 (5,0 ± 1,8)
Senecivernine (sum of senecivernine and integerrimine as senecivernine)	μg/kg	< 5,0 (5,0 ± 1,8)
Senecivernine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Senkirkine	μg/kg	< 5,0 (5,0 ± 1,8)
Usaramine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)
Sum of pyrrolizidine alkaloids	μg/kg	below quantification limit
Caffeine ¹⁾ PN-ISO 10095:1997 (withdrawn)	mg/dose	218
* Vitamin B12 (cyanocobalamin) ²⁾ PB-328 ed. 2 of 05.09.2022	µg/dose	2,36 ± 0,47
* Vitamin B3 (niacin) ¹⁾ EN 15652:2009	mg/dose	69,7

¹⁾ Dose declared by the Client: 7500 mg (6 capsules).

²⁾ Specificity: cobalamin, cyanocobalamin, hydroxycobalamin, methylocobalamin, adenosylcobalamin. No cross reactivity.

³⁾ The lower limit of the measuring range of the accredited method, which is also the limit of quantification set by the Laboratory.

⁴⁾ Limit of quantification 5,0 (5,0 \pm 1,8) µg/kg.

Authorized by: Ada Okunek, Analysis Expert, Microbiology Laboratory Anna Polanin, Manager, Microbiology Laboratory Ewa Ostrach-Grzybowska, Analysis Expert, Vitamin Analysis Laboratory Kamila Skolmowska, Analysis Expert, Liquid Chromatography Laboratory Marcin Kubiak, Vitamins Testing Laboratory Manager, Vitamin Analysis 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 Ks. Stanisława Kujota 8, 70-605 Szczecin

THE END OF THE REPORT

Page 2/3





TEST REPORT NO 252540/23/GDY

The results refer only to the samples received. 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 of the accredited method 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 accredited method, whereas the given expanded measurement uncertainty relates only to the lower or upper limit of the measuring range of the accredited method respectively. In such a case, the Laboratory presents the opinion and interpretation in the "statement to conformity" column, which is based on the obtained test outcome. This test report may not be copied in part without the prior written permission of 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

Page 3/3