









TEST REPORT NO 252539/23/GDY/2

Client SFD SPÓŁKA AKCYJNA GŁOGOWSKA 41 45315 OPOLE		Sample (according to declaration of Client) Sample description: LOCO Focus & Stimulus 120 caps Batch: LC230402 Production date: 01.04.2023	
Sample reception date:	17.05.2023	Sample status: no objections	
Start of analysis	17.05.2023		
End of analysis	01.06.2023	Sample received from the Client	
Test report date	15.06.2023		

Test Method	Unit	Result
* Number of yeasts and moulds at 25°C PN-ISO 21527-2:2009 (withdrawn)		
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)
Lasiocarpine	μg/kg	< 5,0 (5,0 ± 1,8)
Lasiocarpine N-oxide	μg/kg	< 5,0 (5,0 ± 1,8)











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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 ²⁾ PN-ISO 10095:1997 (withdrawn)	mg/dose	367
* Vitamin B2 (ryboflavin) ¹⁾ PN-EN 14152:2014-07		
Vitamin B ₂ (ryboflavin)	mg/dose	6,24
* Vitamin B3 (niacin) ¹⁾ EN 15652:2009	mg/dose	26,8

- 1) Dose declared by the Client: 8040 mg (6 capsules).
- 2) Dose declared by the Client: 8040 (6 capsules).
- 3) The lower limit of the measuring range of the accredited method, which is also the limit of quantification set by the Laboratory.
- Limit of quantification 5,0 (5,0 \pm 1,8) μ g/kg.

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

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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, whereas the given expanded measurement uncertainty relates only 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, whereas the given expanded measurement uncertainty relates only 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, whereas the given expanded measurement uncertainty relates only 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, whereas the given expanded measurement uncertainty, unless otherwise reported. If the "result" column 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, whereas the given expanded measurement uncertainty, unless otherwise reported. If the "result" column 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, whereas the given expanded measurement uncertainty, unless of the accredited method, whereas the given expanded measurement uncertainty and the presult of the given expanded measurement uncertainty

- * Test method accredited
- # Test performed by external provider











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