

REPORT NO 307/10/2022

Customer ¹ :	BLUE TARGET LTD SPÓŁKA KOMANDYTOWA Górska 15, 84-230 Rumia						
Customer order no. ¹ : (if applicable)	no data						
Order no.:	357/09/2022	Order date:	15.09.2022				
Sample type ¹ :	Diet supplement	Sample code:	45/15/09/22				
Laboratory name and address:	Ekolabos sp. z o.o. ul. Duńska 9, 54-427 Wrocław						
Testing purpose ¹ :	Customer's own needs						
Test initiation date:	15.09.2022	Test end date:	06.10.2022				
Reporting date:	06.10.2022						
Sampling							
Sampling person ¹ :	Object taken and delivered by the Customer	Sampling method ¹ :	No data				
Sampling location ¹ :	No data	Sampling point ¹ :	No data				
Sample register date ¹ :	15.09.2022	Sample condition:	Unreservedly				
Sampling protocol:	No data						
Sample description							
Sample name:	APOLLOS HEGEMONY Z-Balance 180 caps						
LOT:	05/2025						
Production date:	No data						
Expiration date:	31.05.2025						
Packaging type:	original						
Material:	No data						
Date of receipt of the sample:	15.09.2022						
Tests done in the laboratory							
No.	Tested parameter	Test method	Result [±uncertainty]	Unit	Authorizing person	The highest allowable value or range	Statement of assesment
1	Detection od Salmonella spp.	PN-EN ISO 6579-1:2017-04+ A1:2020-09	NA	not detected in 25g	-	BD	-
2	Total number of microorganisms	PN-EN ISO 4833-1:2013-12/Ap1:2016-11E	NA	<10	cfu/1 g	BD	-
3	Enumeration of Enterobacteriaceae	PN-ISO 21528-2:2017-08	NA	<10	cfu/1 g	BD	-
4	Enumeration of yeasts and mould	PN-ISO 21527-1:2009	NA	<10	cfu/1 g	BD	-
5	Detection of Staphylococcus aureus	PN-EN ISO 6888-3:2004/AC:2005	NA	not detected in 1g	-	BD	-
6	Detection of Listeria monocytogenes	PN-EN ISO 11290-1:2017-07	NA	not detected in 25g	-	BD	not detected in 25g
7	Enumeration of Escherichia coli B-glucuronidase positive	PN-ISO 16649-2:2004	NA	<10	cfu/1 g	BD	-
8	Mercury	PN-EN 15763:2010	A P1	<0,001	mg/kg	P1	-
9	Lead	PN-EN 15763:2010	A P1	0,030 [+/-0,005]	mg/kg	P1	-



Tests done in the laboratory								
No.	Tested parameter	Test method		Result [±uncertainty]	Unit	Authorizing person	The highest allowable value or range	Statement of assessment
10	Cadmium	PN-EN 15763:2010	A P1	0,068 [+/-0,010]	mg/kg	P1	-	-
11	Arsenic	PN-EN 15763:2010	A P1	0,010 [+/-0,002]	mg/kg	P1	-	-

Legend/Explanations:

Employees authorizing test results:

BD - Bartłomiej Dudek

P1 - testing performed by an external service provider AB 1095

Testing methods marked with a symbol: A - accredited tests, NA - non-accredited tests, S- tests approved by PPIS in Wrocław, decision 5169/21 of 18.10.2021 r. and decision 1824/22 of 22.03.2022, R- the reference method may, if the law so provides, be used to statement of conformity in a regulated area, EP - tests performed by an external service provider.

(W) standard removed from the Polish Standards catalog without replacement.

* Maximum acceptable value or range in order to COMMISSION REGULATION (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs

¹Data declared by the Customer, information obtained from the Customer may affect the validity of the result

In case of sampling and delivery of the sample by the Client, the Laboratory is not responsible for the process of sampling and transportation of the samples for testing.

Results marked with a below (<) sign and an above (>) sign mean that the results lie below and above the range.

The test results apply only to the sample received and tested.

NZ result a statement of nonconformity, ZG result a statement of conformity.

The laboratory provides uncertainty: at the customer's request, when conformity to requirements is evaluated, when it is relevant to the validity and applicability of the test results. The laboratory reports the uncertainty as expanded uncertainty with a confidence level of 95% and a coverage factor of k=2. Where the sample is provided by the customer, the uncertainty does not include the sampling.

Uncertainty of testing and statement of conformity was agreed with the customer at the ordering.

The Laboratory is responsible for all information in the Test Report, except for information provided by the Client.

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The original Test Report is issued in electronic form with the extension*.pdf, signed with a qualified electronic signature. Therefore, all hard copies, unless certified as true copies of the original, are copies.

Complaints must be received no more than 14 days from the date of mailing of the Test Report. Complaints may be received at: biuro@ekolabos.pl.

Reported by:

Elżbieta Świerczek - Pluska

Environmental Project Specialist / Cosmetic Product Research Specialist

Report approved by:

Mateusz Latosiński

Customer Services Manager / Key Account Specialist

---END OF THE REPORT---

