GBAPOLSKA

ANALYTICAL LABORATORIES

microbiology - physicochemistry - sensory

A - accredited methodology (AB 1095); reference - if the law so provides (the result can be used to assess compliance in the legally regulated area).

AR - accredited methodology (AB 1095) equivalent to reference (the result can be used to assess compliance in the legally regulated area).

GBA POLSKA Sp. z o.o Member of GBA GROUP

Customer:

Order No.:

area)

Headquarter address: ul. Mochtyńska 65, 03-289 Warsaw, Poland

Ł/0/06/2024/1359

TEST REPORT No.: Ł/0/06/2024/1359/F/2/EN

AE - accredited methodology (AB 1095) of flexible scope - reference if the law so provides / equivalent to reference (the result can be used to assess compliance in the legally regulated

OSTROVIT SPÓŁKA Z OGRANICZONĄ ODPOWIEDZIALNOŚCIĄ 18-300 Zambrów, ul. Sitarska 16

Materia	al/product tested: Fo	ood						
Address (sampling / collection) :		18-300 Zam	ıbrów, ul.Sitarska 16					
Product name: Os		stroVit Beef I	rotein (choco	Date*: 24.05.2024				
Lot nun	production: 1ber:		no data no data 7BEP009 cl	nc; Minimum date durability: 20/				
	collected according to: transported by: Shipping				Sample receiver:	GBA POLSKA	employee no.	: 28
Sample	no. 19728/06/24 S	Sample valuation:	unreservedl	y Analysis start da	te: 11-06-2024 Analy	ysis end date:	12-06-202	4
Lab.	Analyzed paramete	er Ur	it Accred.	Test method	Requirement	Result	MU**	N
	Lead .	mg/	kg A/P	DIN EN 15763, mod., ICP-MS: 2010-04 (Nr Akr. D-PL-14170-01- 00)	no requirements	<0,020		
	Mercury .	mg	kg A/P	DIN EN 15763, mod., ICP-MS: 2010-04 (Nr Akr. D-PL-14170-01- 00)	no requirements	<0,010		
	Cadmium .	mg/	kg A/P	DIN EN 15763, mod., ICP-MS: 2010-04 (Nr Akr. D-PL-14170-01- 00)	no requirements	<0,010		+

Date* - depending on the method of obtaining the sample by GBA Polska, it is the date of: collection (when the sample is collected only by a GBA Polska employee) or collection (when the sample is collected from customer by a GBA Polska employee, is delivered by a courier company or delivered personally by the customer). MU** - expanded measurement uncertainty at the level of confidence app. 95% and the coverage factor k=2, does not take into account the sampling uncertainty, except when indicated in the remarks.

Measurement uncertainty is presented when: it is relevant to the validity or application of the test results, it affects conformity to a specification limit, or a customer's instruction so requires. The test results lower or higher than the measuring ranges of the methods are presented as "<value of the lower limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range " or "> value of the upper limit of the measuring range (or ") value of the upper limit of the measuring range (or ") value of the upper limit of the measuring range (or ") value of the upper limit of respectively. These values provide information about the research results. If expanded uncertainties are given with these test results, they apply to the lower or upper limit of the measuring range of the method. In such a case, if the test results meet the requirements of PCA Communication No. 353 of August 24, 2021, the determination of compliance will be made as part of the opinion and interpretation.

The results relate to the tested samples (sampled or received - as reported in the test report). The underlined information included in the report was provided by the Client. The Laboratory is not responsible for this information. The laboratory is not resposible for the method of sampling and the representativeness of the samples provided by the customer for testing

The test report without the written approval of the Laboratory shall not be reproduced except in full

Customer may file complains within 14 days from receiving the report. The Laboratory does not store the samples after testing, unless otherwise agreed with the customer. Place of performance of the tests ("Lab."): Ł - Łajski, L - Lublin, M - Mysłowice, PS - in situ measurement.

Remarks:

NOTE: The original test reports are issued as PDF file, signed with a qualified electronic signature. Therefore, all prints are copies, unless certified to be true to the original PDF file.

Ł/0/06/2024/1359/F/2/EN

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