

Test reports

BRAINMAX® ACTIVATED B-COMPLEX



BR	AINMA	X® A	ACTIV	ATED	B-COM	PLEX	<u> </u>	
Quality Declaration – Composi	tion per D	ose (2 C	apsules)		Lab	oratory	Analysis	1
	Quantity of	Measur			Analytical R	esult		
Active Substances	Substances per Dose	ement Unit	*DV Measure Value		Measurement Uncertainty Result F		Page*	Comment
Vitamin B3 (niacin)	100	mg	625%	98	30%		3-4	
Alpha-lipoic acid	90	mg	-					
Vitamin C (liposomal vitamin C)	80	mg	100%	72	15%		5-6	
Betaine (betaine hydrochloride)	60	mg	-					
Vitamin B1 (thiamine hydrochloride)	50	mg	4545%	51	16%		7-8	
Vitamin B2 (riboflavin)	50	mg	3571%	51	16%		9-10	
Vitamin B5 (calcium pantothenate)	50	mg	833%	48	20%		11-12	
Inositol	40	mg	-					
Choline bitartrate	29	mg	-					
Vitamin B6 (pyridoxal-5-phosphate)	25	mg	1786%					
PABA (para-aminobenzoic acid)	25	mg	-					
Coenzyme Q10	20	mg	-	19	20%		13-14	
Vitamin H (biotin)	1000	μg	2000%	1041	30%		15-16	
Vitamin B9 (folic acid)	400	μg	200%	379	20%		17-18	
Vitamin B12 (methylcobalamin)	400	μg	16000%					

^{*}Page in the Test Report





Issue Date · 27-Jun-2025 Work Order : FP2507238-AA

Amendment

Client : ALS Czech Republic, s.r.o. : BRAINMARKET s.r.o. Laboratory

Contact Contact : F&P Client Service : Matěj Veselý

Address : Hladnovská 83/93 Address : Na Harfe 336/9 Prague 9 - Vysocany

> Mualinov 190 00 Czech Republic

712 00 Ostrava

E-mail E-mail : czsupport.food@alsglobal.com

Telephone Telephone : +420 226 226 998

Project Page : Stanovení hladiny účinné látky u : 1 of 2

vitamínů B3 a B5 v produktu Activated

B-Complex

Order number : 20250226 Date Samples : 27-Feb-2025

Received

Quote number : FP2025ROSHO-CZ0001

(CZ-114-25-5010)

Site Date of test : 01-Mar-2025 - 10-Mar-2025

Sampled by QC Level : ALS CR Standard Quality Control

Schedule

General Comments

This report shall not be reproduced except in full, without prior written approval from the laboratory. The laboratory is not responsible for the sample data supplied by the customer and their impact on the validity of the result.

The laboratory declares that the test results relate only to the listed samples. If "ALS" is not included in the test report in the "Sampled by" section, then the results refer to the sample as received.

Amendment 1: protocol was divided and added a comment. This Amendment 1 replaces the original report issued on 10.03.2025.

Responsible for accuracy

Testing Laboratory No. 1163 Accredited by CAI according to CSN EN ISO/IEC 17025:2018

Signatories **Position**

Kateřina Hubáčková Site Manager Food









GMP+ Registered Laboratory-GMP050490

The company is certified according to ČSN EN ISO 14001 (Environmental management systems) and ČSN ISO 45001 (Occupational health and safety management systems)

Sample Information

No. of samples received : 1 Date Samples Received No. of samples analysed : 1 27-Feb-2025 14:00

When sampling date is not provided by the client, the laboratory determines it for procedural reasons, then it is equal to the date of receipt of the sample to the laboratory and is displayed in brackets.

Sub-Matrix: DIETARY SUPPLEMENT

Laboratory sample ID Client sample ID Client sampling date / time

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Issue Date : 27-Jun-2025
Page : 2 of 2

Work Order : FP2507238-AA Amendment 1

Client : BRAINMARKET s.r.o.



Sub-Matrix : DIETARY SUPPLEMENT

Laboratory sample ID	Client sample ID	Client sampling date / time
FP2507238-001	Activated B-Complex 314229	26-Feb-2025 08:10

Analytical Results

Sub-Matrix: DIETARY SUPPLEMENT		Laborate	ory sample ID	FP250723	8-001				
	Client sampling date / time			26-Feb-2025	08:10				
Parameter	Method	LOR	Unit	Result	MU	Result	MU	Result	MU
Vitamins									
Niacin	B-VB3LCF02	7.0	mg/kg	88100	± 30.0%				

Measurement uncertainty is expressed as expanded measurement uncertainty with coverage factor k = 2, representing 95% confidence level. Key: LOR = Limit of reporting; MU = Measurement Uncertainty. The MU does not include sampling uncertainty.

Brief Method Summaries

Analytical Methods	Method Descriptions
Location of test performance	: Na Harfe 336/9 Prague 9 - Vysocany Czech Republic 190 00
B-VB3LCF02	CZ_SOP_D06_09_225 (CSN EN 15652) Determination of niacin by liquid chromatography method with PDA detection.

The symbol "*" for the method indicates a test outside the scope of accreditation of the laboratory or subcontractor. If the UNICO-SUB code is stated in the method table, this only informs that the tests have been performed by a subcontractor and the results are given in an annex to the test report, including information on test accreditation. If the lab used for matrix outside the scope of accreditation or non-standard sample matrix procedure specified in the accredited method and issues non-accredited results, this fact is stated on the title page of this protocol in the section "Notes". If the test report shows the results of subcontracting, the place of performance of the test is outside the laboratories of ALS Czech Republic, s.r.o.

The method for calculating of the summation parameters is available on request in the customer service.



· 27-Jun-2025 Work Order Issue Date : FP2508335-AI

Client : BRAINMARKET s.r.o. Laboratory : ALS Czech Republic, s.r.o.

Contact : Matěj Veselý Contact : F&P Client Service

Address Address : Hladnovská 83/93 : Na Harfe 336/9 Prague 9 - Vysocany

190 00 Czech Republic

Muglinov 712 00 Ostrava

E-mail E-mail : czsupport.food@alsglobal.com

Telephone Telephone : +420 226 226 998

Project Page : Stanovení hladiny účinné látky u : 1 of 2

Liposomálního vitamínu C v

produktech Brainmax

Order number : 20250307 : 10-Mar-2025 **Date Samples**

Received

Quote number : FP2025ROSHO-CZ0001

(CZ-114-25-5010)

Site Date of test : 10-Mar-2025 - 21-Mar-2025 Sampled by QC Level

: ALS CR Standard Quality Control

Schedule

General Comments

This report shall not be reproduced except in full, without prior written approval from the laboratory. The laboratory is not responsible for the sample data supplied by the customer and their impact on the validity of the result.

The laboratory declares that the test results relate only to the listed samples. If "ALS" is not included in the test report in the "Sampled by" section, then the results refer to the sample as received.

Responsible for accuracy

Testing Laboratory No. 1163 Accredited by CAI according to CSN EN ISO/IEC 17025:2018

Signatories Position

Kateřina Hubáčková Site Manager Food









GMP+ Registered Laboratory-GMP050490

The company is certified according to ČSN EN ISO 14001 (Environmental management systems) and ČSN ISO 45001 (Occupational health and safety management systems)

Sample Information

No. of samples received Date Samples Received : 1 No. of samples analysed 10-Mar-2025 09:00 : 1

When sampling date is not provided by the client, the laboratory determines it for procedural reasons, then it is equal to the date of receipt of the sample to the laboratory and is displayed in brackets.

Sub-Matrix: DIETARY SUPPLEMENT

Laboratory sample ID	Client sample ID	Client sampling date / time
FP2508335-009	Brainmax Activated B-Complex 314327	07-Mar-2025 08:10

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 Issue Date
 : 27-Jun-2025

 Page
 : 2 of 2

 Work Order
 : FP2508335-AI

 Client
 : BRAINMARKET s.r.o.



Analytical Results

Sub-Matrix: DIETARY SUPPLEMENT		Laboratory sample ID		FP2508335-009					
	Client sampling date / time			07-Mar-2025 08:10					
Parameter	Method	LOR	Unit	Result	MU	Result	MU	Result	MU
Vitamins									
Vitamin C (Ascorbic Acid)	B-VCLCP02	8.0	mg/kg	64400	± 15.0%				

Measurement uncertainty is expressed as expanded measurement uncertainty with coverage factor k = 2, representing 95% confidence level. Key: LOR = Limit of reporting; MU = Measurement Uncertainty. The MU does not include sampling uncertainty.

Brief Method Summaries

Analytical Methods	Method Descriptions
Location of test performance	e: Na Harfe 336/9 Prague 9 - Vysocany Czech Republic 190 00
B-VCLCP02	CZ_SOP_D06_09_207 (CSN EN 14130:2004) Determination of vitamin C (ascorbic acid) by liquid chromatography method
	with PDA detection.

The symbol "*" for the method indicates a test outside the scope of accreditation of the laboratory or subcontractor. If the UNICO-SUB code is stated in the method table, this only informs that the tests have been performed by a subcontractor and the results are given in an annex to the test report, including information on test accreditation. If the lab used for matrix outside the scope of accreditation or non-standard sample matrix procedure specified in the accredited method and issues non-accredited results, this fact is stated on the title page of this protocol in the section "Notes". If the test report shows the results of subcontracting, the place of performance of the test is outside the laboratories of ALS Czech Republic, s.r.o.

The method for calculating of the summation parameters is available on request in the customer service.





AR-25-HD-016907-02



Testing laboratory:

Eurofins Food & Feed Testing Czech Republic s.r.o. Zkušební laboratoř EUROFINS CZ Radiová 1285/7

102 00 Praha 10 - Hostivař

IČO: 27449408

tel.: +420 778 488 111 E-mail: ClientService.cz@ftcee.eurofins.com

Issue date 07.08.2025

Sample code 540-2025-00026225

Sample reception date: 20.05.2025

Date of Testing 20.05.2025 - 23.05.2025

Sample information:

Sample name, extended: 1) BRAINMAX ACTIVATED B-COMPLEX

 Sample description:
 1) 005-32407-226425

 Client Purchase order nr.:
 20250519

 Order date:
 19.05.2025

 Client sample code:
 1) 5

Additional sample description: BRAINMARKET s.r.o.

Physical and chemical tests

Parameter	Unit	Result	Uncertainty of measurement *	Method	Method principle	TZ
Brutto sample weight of supplied sample	kg	0.22	5%	SOP MB.005.PB	Gravimetry	A
Thiamine hydrochloride (vitamin B1-HCl)	mg/100 g	4570	731.200	BS EN 14122-2014.	LC-FLD	SA
Parameter	Unit	Result	Uncertainty of measurement	Method	Method principle	TZ

Testing performed on whole capsule

Yes

Testing on whole capsules SN

Decision rule: If the testing laboratory issues a statement of conformity, the decision-making rule according to ch. 4.2.1 of ILAC document G8:09/2019 Guidelines for the use of decision rules and statement of conformity. In such a case, the measurement uncertainty is not taken into account for the conformity statement. If measurement uncertainty is included the decision, this information is included in the statement of conformity. In such a case, proceed according to chap. 4.2.3 ILAC G8:09/2019.

Notes: SOP, ŠPP - Standard operation procedure TZ - type of test

ND - not detected by given method

A - test within the accreditation scope of EUROFINS CZ

CFU - Colony forming unit

N - test outside of the accreditation scope of EUROFINS CZ

NN - necessary quantity

SA - subcontracted accredited test

SN - subcontracted not accredited test

* - the expanded measurement uncertainty, as determined by the extension coefficient k = 2 (with a 95% probability), does not include sampling uncertainty; if the measurement uncertainty is expressed in %, it is its relative value

 $LOD-limit\ of\ detection,\ LOQ-limit\ of\ quantification,\ result\ between\ LOD\ and\ LOQ=detected$

1) - Information supplied by customer

Unless otherwise stated in the notes, the place of the tests performance is workplace No. 1 - Prague - of EUROFINS CZ testing laboratory.





If the information supplied by the customer could have be to affect the validity of the results, the laboratory disclaims responsibility. For samples supplied by the customer, the results relate to the sample as received and provided by the customer. The measuring devices and gauges used for the test / tests have been calibrated and verified according to valid metrological regulations. The results of the measurements relate only to the subject of the tests have been documents, e.g. of an administrative nature. The result identified as subcontracting in this protocol is the result of subcontractor measurements based on contract, order. The protocol may be reproduced or incorporated into promotional materials only with the written consent of the EUROFINS CZ Testing Laboratory and only to the extent of such approval. Any alteration, reproduction of part of the test report is not permitted and such analytical report automatically becomes invalid. The authenticity and completeness of the report can be verified at the EUROFINS CZ test laboratory stated in the header of analytical report. This Test Report has been issued in accordance with the applicable Conditions of service available on request and accessible at www.eurofins.cz.

Responsible for correctness: Jitka Pinkrová

Worked out by: Júlia Výbostoková No. of document: 20258793342675

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Test Certificate approved by:

Jitka Pinkrová







AR-25-HD-016908-02



Testing laboratory:

Eurofins Food & Feed Testing Czech Republic s.r.o. Zkušební laboratoř EUROFINS CZ Radiová 1285/7

102 00 Praha 10 - Hostivař

IČO: 27449408

tel.: +420 778 488 111 E-mail: ClientService.cz@ftcee.eurofins.com

Issue date 07.08.2025

Sample code 540-2025-00026226

Sample reception date: 20.05.2025

Date of Testing 20.05.2025 - 23.05.2025

Sample information:

Sample name, extended: 1) BRAINMAX ACTIVATED B-COMPLEX

 Sample description:
 1) 005-32407-226452

 Client Purchase order nr.:
 20250519

 Order date:
 19.05.2025

Client sample code:

Additional sample description: BRAINMARKET s.r.o.

Physical and chemical tests

Parameter	Unit	Result	Uncertainty of measurement *	Method	Method principle	TZ
Brutto sample weight of supplied	kg	0.22	5%	SOP MB.005.PB	Gravimetry	A
sample Riboflavin (vitamin B2)	mg/100 g	4610	737.600	EN 14152:2014 mod	LC-FLD	SA
Parameter	Unit	Result	Uncertainty	Method	Method principle	TZ

measurement *	Parameter	Unit	Result	Uncertainty of	Method	Method principle	TZ
				measurement *			

Testing performed on whole capsule

Yes

Testing on whole capsules SN

Decision rule: If the testing laboratory issues a statement of conformity, the decision-making rule according to ch. 4.2.1 of ILAC document G8:09/2019 Guidelines for the use of decision rules and statement of conformity. In such a case, the measurement uncertainty is not taken into account for the conformity statement. If measurement uncertainty is included the decision, this information is included in the statement of conformity. In such a case, proceed according to chap. 4.2.3 ILAC G8:09/2019.

Notes: SOP, ŠPP - Standard operation procedure TZ - type of test

ND - not detected by given method A - test within the accreditation scope of EUROFINS CZ
CFU - Colony forming unit N - test outside of the accreditation scope of EUROFINS CZ
NM - necessary quantity SA - subcontracted accredited test

SN - subcontracted not accredited test

* - the expanded measurement uncertainty, as determined by the extension coefficient k = 2 (with a 95% probability), does not include sampling uncertainty; if the measurement uncertainty is expressed in %, it is its relative value

LOD – limit of detection, LOQ – limit of quantification, result between LOD and LOQ = detected

1) - Information supplied by customer

Unless otherwise stated in the notes, the place of the tests performance is workplace No. 1 - Prague - of EUROFINS CZ testing laboratory.





If the information supplied by the customer could have be to affect the validity of the results, the laboratory disclaims responsibility. For samples supplied by the customer, the results relate to the sample as received and provided by the customer. The measuring devices and gauges used for the test / tests have been calibrated and verified according to valid metrological regulations. The results of the measurements relate only to the subject of the tests and do not replace other documents, e.g. of an administrative nature. The result identified as subcontracting in this protocol is the result of subcontractor measurements based on contract, order. The protocol may be reproduced or incorporated into promotional materials only with the written consent of the EUROFINS CZ Testing Laboratory and only to the extent of such approval. Any alteration, reproduction of part of the test report is not permitted and such analytical report automatically becomes invalid. The authenticity and completeness of the report can be verified at the EUROFINS CZ test laboratory stated in the header of analytical report. This Test Report has been issued in accordance with the applicable Conditions of service available on request and accessible at www.eurofins.cz.

Responsible for correctness: Jitka Pinkrová

Worked out by: Júlia Výbostoková No. of document: 20258793451714

Validity check of document

https://www.linktothedocument.com



Test Certificate approved by:

Jitka Pinkrová

Laboratory

Laboratory

Eurofin Furning Food & Feed Testing



Issue Date · 27-Jun-2025 Work Order : FP2507238-AB

Amendment

Client : ALS Czech Republic, s.r.o. : BRAINMARKET s.r.o. Laboratory

Contact Contact : F&P Client Service : Matěj Veselý

Address : Hladnovská 83/93 Address : Na Harfe 336/9 Prague 9 - Vysocany

> Mualinov 190 00 Czech Republic

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Project Page : Stanovení hladiny účinné látky u : 1 of 2

vitamínů B3 a B5 v produktu Activated

B-Complex

Order number : 20250226 Date Samples : 27-Feb-2025

Received

Quote number : FP2025ROSHO-CZ0001

(CZ-114-25-5010)

Site Date of test : 01-Mar-2025 - 10-Mar-2025

Sampled by QC Level : ALS CR Standard Quality Control

Schedule

General Comments

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The laboratory declares that the test results relate only to the listed samples. If "ALS" is not included in the test report in the "Sampled by" section, then the results refer to the sample as received.

Amendment 1: protocol was divided and added a comment. This Amendment 1 replaces the original report issued on 10.03.2025.

Responsible for accuracy

Testing Laboratory No. 1163 Accredited by CAI according to CSN EN ISO/IEC 17025:2018

Signatories **Position**

Kateřina Hubáčková Site Manager Food









GMP+ Registered Laboratory-GMP050490

The company is certified according to ČSN EN ISO 14001 (Environmental management systems) and ČSN ISO 45001 (Occupational health and safety management systems)

Sample Information

No. of samples received : 1 Date Samples Received No. of samples analysed : 1 27-Feb-2025 14:00

When sampling date is not provided by the client, the laboratory determines it for procedural reasons, then it is equal to the date of receipt of the sample to the laboratory and is displayed in brackets.

Sub-Matrix: DIETARY SUPPLEMENT

Laboratory sample ID Client sample ID Client sampling date / time

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Issue Date : 27-Jun-2025
Page : 2 of 2

Work Order : FP2507238-AB Amendment 1

Client : BRAINMARKET s.r.o.



Sub-Matrix: DIETARY SUPPLEMENT

Laboratory sample ID	Client sample ID	Client sampling date / time
FP2507238-001	Activated B-Complex 314229	26-Feb-2025 08:10

Analytical Results

Sub-Matrix: DIETARY SUPPLEMENT			· · · · ·		8-001 5 08:10				
Parameter	Method	LOR	Unit	Result	MU	Result	MU	Result	MU
Vitamins									
Pantothenic acid	B-VB5LCP02	12.5	mg/kg	43500	± 20.0%				
Calcium pantothenate	B-VB5LCP02	12.5	mg/kg	47500	± 20.0%				

Measurement uncertainty is expressed as expanded measurement uncertainty with coverage factor k = 2, representing 95% confidence level. Key: LOR = Limit of reporting; MU = Measurement Uncertainty. The MU does not include sampling uncertainty.

Brief Method Summaries

Analytical Methods	Method Descriptions						
Location of test performance	Location of test performance: Na Harfe 336/9 Prague 9 - Vysocany Czech Republic 190 00						
B-VB5LCP02	CZ_SOP_D06_09_238 Determination of pantothenic acid (vitamin B5) by liquid chromatography method with PDA detection.						

The symbol "*" for the method indicates a test outside the scope of accreditation of the laboratory or subcontractor. If the UNICO-SUB code is stated in the method table, this only informs that the tests have been performed by a subcontractor and the results are given in an annex to the test report, including information on test accreditation. If the lab used for matrix outside the scope of accreditation or non-standard sample matrix procedure specified in the accredited method and issues non-accredited results, this fact is stated on the title page of this protocol in the section "Notes". If the test report shows the results of subcontracting, the place of performance of the test is outside the laboratories of ALS Czech Republic, s.r.o.

The method for calculating of the summation parameters is available on request in the customer service.





AR-25-HD-017400-02



Testing laboratory:

Eurofins Food & Feed Testing Czech Republic s.r.o. Zkušební laboratoř EUROFINS CZ Radiová 1285/7

102 00 Praha 10 - Hostivař

IČO: 27449408

tel.: +420 778 488 111 E-mail: ClientService.cz@ftcee.eurofins.com

Issue date 07.08.2025

Sample code 540-2025-00026229

20.05.2025 Sample reception date:

Date of Testing 20.05.2025 - 30.05.2025

Sample information:

Sample name, extended: 1) BRAINMAX ACTIVATED B-COMPLEX

1) 005-32407-226456 Sample description: Client Purchase order nr.: 20250519 Order date: 19.05.2025 Client sample code:

BRAINMARKET s.r.o. Additional sample description:

Physical and chemical tests

Parameter	Unit	Result	Uncertainty of measurement *	Method	Method principle	TZ
Brutto sample weight of supplied sample	kg	0.22	5%	SOP MB.005.PB	Gravimetry	A
Coenzyme Q10	mg/Capsule	9.51	20%	0-24	LC-UV/VIS	SA

Decision rule: If the testing laboratory issues a statement of conformity, the decision-making rule according to ch. 4.2.1 of ILAC document G8:09/2019 Guidelines for the use of decision rules and statement of conformity. In such a case, the measurement uncertainty is not taken into account for the conformity statement. If measurement uncertainty is included the decision, this information is included in the statement of conformity. In such a case, proceed according to chap. 4.2.3 ILAC G8:09/2019.

Notes: SOP, ŠPP - Standard operation procedure TZ - type of test

ND - not detected by given method A - test within the accreditation scope of EUROFINS CZ N - test outside of the accreditation scope of EUROFINS CZ CFU - Colony forming unit SA - subcontracted accredited test NM - necessary quantity

SN - subcontracted not accredited test

* - the expanded measurement uncertainty, as determined by the extension coefficient k = 2 (with a 95% probability), does not include sampling uncertainty; if

the measurement uncertainty is expressed in %, it is its relative value

LOD – limit of detection, LOQ – limit of quantification, result between LOD and LOQ = detected

1) - Information supplied by customer

Unless otherwise stated in the notes, the place of the tests performance is workplace No. 1 - Prague - of EUROFINS CZ testing laboratory.

If the information supplied by the customer could have be to affect the validity of the results, the laboratory disclaims responsibility. For samples supplied by the customer, the results relate to the sample as received and provided by the customer. The measuring devices and gauges used for the test / tests have been calibrated and verified according to valid metrological regulations. The results of the measurements relate only to the subject of the tests and do not replace other documents, e.g. of an administrative nature. The result identified as subcontracting in this protocol is the result of subcontractor measurements based on contract, order. The protocol may be reproduced or incorporated into promotional materials only with the written consent of the EUROFINS CZ Testing Laboratory and only to the extent of such approval. Any alteration, reproduction of part of the test report is not permitted and such analytical report automatically becomes invalid. The authenticity and completeness of the report can be verified at the EUROFINS CZ test laboratory stated in the header of analytical report. This Test Report has been issued in accordance with the applicable Conditions of service available on request and accessible at www.eurofins.cz.





Responsible for correctness: Pavlína Hejduková

Worked out by: Júlia Výbostoková No. of document: 20258793553930 Validity check of document https://www.linktothedocument.com



Test Certificate approved by:





· 27-Jun-2025 Work Order Issue Date : FP2510415

Client : BRAINMARKET s.r.o. Laboratory : ALS Czech Republic, s.r.o.

Contact : Matěj Veselý Contact : F&P Client Service

Address Address : Hladnovská 83/93 : Na Harfe 336/9 Prague 9 - Vysocany

190 00 Czech Republic

712 00 Ostrava

Muglinov

E-mail E-mail : czsupport.food@alsglobal.com

Telephone Telephone : +420 226 226 998

Project Page : Stanovení hladiny účinné latky : 1 of 2

Vitaminu H(Biotin) v produktu Brainmax

Activated B-complex

Order number : 26-Mar-2025 : 20250324 1 **Date Samples**

Received

Quote number : FP2025ROSHO-CZ0001

(CZ-114-25-5010)

Site Date of test : 26-Mar-2025 - 03-Apr-2025 Sampled by QC Level

: ALS CR Standard Quality Control

Schedule

General Comments

This report shall not be reproduced except in full, without prior written approval from the laboratory. The laboratory is not responsible for the sample data supplied by the customer and their impact on the validity of the result.

The laboratory declares that the test results relate only to the listed samples. If "ALS" is not included in the test report in the "Sampled by" section, then the results refer to the sample as received.

The method B-VB7ELI01, LOR of determined parameters may be increased due to character of the matrix/ sample preparation/ dilution of the sample.

Responsible for accuracy

Testing Laboratory No. 1163 Accredited by CAI according to CSN EN ISO/IEC 17025:2018

Signatories Position

Kateřina Hubáčková Site Manager Food









GMP+ Registered Laboratory-GMP050490

The company is certified according to ČSN EN ISO 14001 (Environmental management systems) and ČSN ISO 45001 (Occupational health and safety management systems)

Sample Information

No. of samples received Date Samples Received : 1 26-Mar-2025 09:00 No. of samples analysed : 1

When sampling date is not provided by the client, the laboratory determines it for procedural reasons, then it is equal to the date of receipt of the sample to the laboratory and is displayed in brackets.

Sub-Matrix: DIETARY SUPPLEMENT

Laboratory sample ID	Client sample ID	Client sampling date / time
FP2510415-001	Brainmax Activated B-complex 314639	24-Mar-2025 08:10

 Issue Date
 : 27-Jun-2025

 Page
 : 2 of 2

 Work Order
 : FP2510415

Client : BRAINMARKET s.r.o.



Analytical Results

Sub-Matrix: DIETARY SUPPLEMENT		Laboratory sample ID		FP2510415-001					
	Client sampling date / time			24-Mar-2025 08:10					
Parameter	Method	LOR	Unit	Result	MU	Result	MU	Result	MU
Vitamins									
Vitamin B7 (Biotin)	B-VB7ELI01	10.0	μg/kg	937000	± 30.0%				

Measurement uncertainty is expressed as expanded measurement uncertainty with coverage factor k = 2, representing 95% confidence level. Key: LOR = Limit of reporting; MU = Measurement Uncertainty. The MU does not include sampling uncertainty.

Brief Method Summaries

Analytical Methods	Method Descriptions					
Location of test performance: Na Harfe 336/9 Prague 9 - Vysocany Czech Republic 190 00						
B-VB7ELI01 CZ_SOP_D06_09_220 (Demeditec Manual) Determination of biotin by ELISA method by commercial set						

The symbol "*" for the method indicates a test outside the scope of accreditation of the laboratory or subcontractor. If the UNICO-SUB code is stated in the method table, this only informs that the tests have been performed by a subcontractor and the results are given in an annex to the test report, including information on test accreditation. If the lab used for matrix outside the scope of accreditation or non-standard sample matrix procedure specified in the accredited method and issues non-accredited results, this fact is stated on the title page of this protocol in the section "Notes". If the test report shows the results of subcontracting, the place of performance of the test is outside the laboratories of ALS Czech Republic, s.r.o.

The method for calculating of the summation parameters is available on request in the customer service.





AR-25-HD-017462-02



Testing laboratory:

Eurofins Food & Feed Testing Czech Republic s.r.o. Zkušební laboratoř EUROFINS CZ Radiová 1285/7

102 00 Praha 10 - Hostivař

IČO: 27449408

tel.: +420 778 488 111 E-mail: ClientService.cz@ftcee.eurofins.com

Issue date 07.08.2025

Sample code 540-2025-00026227

Sample reception date: 20.05.2025

Date of Testing 20.05.2025 - 30.05.2025

Sample information:

Sample name, extended: 1) BRAINMAX ACTIVATED B-COMPLEX

 Sample description:
 1) 005-32407-226454

 Client Purchase order nr.:
 20250519

 Order date:
 19.05.2025

 Client sample code:
 1) 7

Additional sample description: BRAINMARKET s.r.o.

Physical and chemical tests

Parameter	Unit		Result			Uncertainty of	Method	TZ	
		1	2	3	4	5	measurement*		
Brutto sample weight of supplied sample	kg	0.22					5%	SOP MB.005.PB	A
Folic acid	$\mu g/100~g$	34100					6820.000	AOAC 2013.13	SN
Methyltetrahydrofolate (5-MTHF)	μg/100 g	<5 (LOQ)						AOAC 2013.13	SN

Parameter	Unit	Result	Uncertainty Me	ethod	Method principle	TZ
			measurement *			

Testing performed on whole capsule

Yes

Testing on whole capsules SN

Decision rule: If the testing laboratory issues a statement of conformity, the decision-making rule according to ch. 4.2.1 of ILAC document G8:09/2019 Guidelines for the use of decision rules and statement of conformity. In such a case, the measurement uncertainty is not taken into account for the conformity statement. If measurement uncertainty is included the decision, this information is included in the statement of conformity. In such a case, proceed according to chap. 4.2.3 ILAC G8:09/2019.

Notes: SOP, ŠPP - Standard operation procedure TZ - type of test

ND - not detected by given method

A - test within the accreditation scope of EUROFINS CZ

CFU - Colony forming unit

A - test within the accreditation scope of EUROFINS CZ

N - test outside of the accreditation scope of EUROFINS CZ

NM - necessary quantity SA - subcontracted accredited test SN - subcontracted not accredited test

* - the expanded measurement uncertainty, as determined by the extension coefficient k = 2 (with a 95% probability), does not include sampling uncertainty; if the measurement uncertainty is expressed in %, it is its relative value

LOD – limit of detection, LOQ – limit of quantification, result between LOD and LOQ = detected

1) - Information supplied by customer

Unless otherwise stated in the notes, the place of the tests performance is workplace No. 1 - Prague - of EUROFINS CZ testing laboratory.





If the information supplied by the customer could have be to affect the validity of the results, the laboratory disclaims responsibility. For samples supplied by the customer, the results relate to the sample as received and provided by the customer. The measuring devices and gauges used for the test / tests have been calibrated and verified according to valid metrological regulations. The results of the measurements relate only to the subject of the tests and do not replace other documents, e.g. of an administrative nature. The result identified as subcontracting in this protocol is the result of subcontractor measurements based on contract, order. The protocol may be reproduced or incorporated into promotional materials only with the written consent of the EUROFINS CZ Testing Laboratory and only to the extent of such approval. Any alteration, reproduction of part of the test report is not permitted and such analytical report automatically becomes invalid. The authenticity and completeness of the report can be verified at the EUROFINS CZ test laboratory stated in the header of analytical report. This Test Report has been issued in accordance with the applicable Conditions of service available on request and accessible at www.eurofins.cz.

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Test Certificate approved by:

Jitka Pinkrová

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