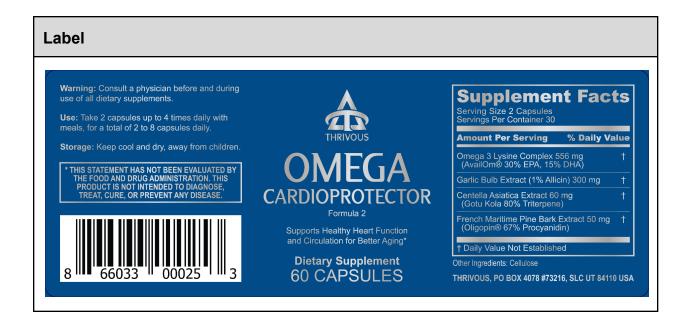


CERTIFICATE OF ANALYSIS AND QUALITY

Product	Omega Cardioprotector
sku	OMEGA
Barcode	866033000253
Formula	2
Date	15 November 2023



Certifications
Letter of Guarantee
Good Manufacturing Practice (GMP) Certificate from Manufacturing
ISO/IEC 17025 Certificate from Third-Party Testing
Certificate of Analysis from Third-Party Testing
Capsule Certificate of Analysis from Supplier
Capsule Certificate of Analysis from Third-Party Testing
Centella Asiatica Certificate of Analysis from Supplier
Centella Asiatica Certificate of Analysis from Third-Party Testing
French Maritime Pine Certificate of Analysis from Supplier
French Maritime Pine Certificate of Analysis from Third-Party Testing
Garlic Certificate of Analysis from Supplier
Garlic Certificate of Analysis from Third-Party Testing
Omega 3 Certificate of Analysis from Supplier
Omega 3 Certificate of Analysis from Third-Party Testing



15 November 2023

RE: Letter of Guarantee for Thrivous Omega Cardioprotector

To whom it may concern,

The undersigned, Lincoln Cannon LLC DBA Thrivous ("Thrivous"), hereby guarantees as follows regarding Omega Cardioprotector ("Product"):

- 1. Product is manufactured according to current Good Manufacturing Practices as indicated in 21 CFR Part 111.
- 2. Product is tested by third party laboratories according to current best practices as indicated in ISO/IEC 17025.
- All ingredients utilized for Product are lawful and safe as defined in section 402(f) of the FD&C Act.
- 4. To the best of Thrivous' knowledge, concentrations of active ingredients, as stated on the label of Product, are safe for consumption.

Thrivous further guarantees that any agent signing on behalf of Thrivous has the authority to bind and obligate Thrivous.

Lincoln Cannon LLC DBA Thrivous

Lincoln Cannon CEO at Thrivous



State of Utah SPENCER J. COX Governor DEIDRE M. HENDERSON Lieutenant Governor

Department of Agriculture and Food

Craig W. Buttars Commissioner Kelly Pehrson Deputy Commissioner Travis Waller Director, Regulatory Services

Certificate No.: REG-2021-12022

GOOD MANUFACTURING PRACTICE CERTIFICATE

We hereby certify that ORIGIN NUTRACEUTICAL INC, located at, 151 E 3450 N, SPANISH FORK, UT 84660 is currently under inspection as a manufacturer of health food or dietary supplements. ORIGIN NUTRACEUTICAL INC has all the facilities to comply with the GOOD MANUFACTURING PRACTICE for food and dietary supplements (Code of Good Manufacturing Practice for food).

We also certify that ORIGIN NUTRACEUTICAL INC, is an inspected facility and the manufacturing plant in which their products are produced are subject to inspections at suitable intervals.

Inspection evaluates and assures compliance with the Utah Wholesome Food Act and Utah Food Protection Rule, which identifies the standard for proper facility construction, good manufacturing practices for food and dietary supplements (GMP), and fulfills requirements of those applicable laws and rules in the State of Utah.

- A W	
LITAH DEPARTMENT	OF AGRICULTURE AND FOOD
U ARI DEPARTMENT	OF AGRICULTURE AND FOOD

Division of Regulatory Services

State of Utah, County of Salt Lake.
On this date DEC 20 2021 before me, the notary, personally appeared

Travis Waller , proved on the basis of satisfactory

evidence to be person, whose name is subscribed to this document, and acknowledge

that he/she executed the same.

Notary Public





PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

Perry Johnson Laboratory Accreditation, Inc. has assessed the Laboratory of:

Contract Testing Laboratories of America 151 E. 3450 N., Spanish Fork, UT 84660

(Hereinafter called the Organization) and hereby declares that Organization is accredited in accordance with the recognized International Standard:

ISO/IEC 17025:2017

This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory quality management system (as outlined by the joint ISO-ILAC-IAF Communiqué dated April 2017):

Chemical and Microbiological Testing (As detailed in the supplement)

Accreditation claims for such testing and/or calibration services shall only be made from addresses referenced within this certificate. This Accreditation is granted subject to the system rules governing the Accreditation referred to above, and the Organization hereby covenants with the Accreditation body's duty to observe and comply with the said rules.

For PJLA:

President

Initial Accreditation Date:

Issue Date:

Expiration Date:

March 31, 2021

March 24, 2023

June 30, 2025

Accreditation No.:

Certificate No.:

102267

L23-261

Perry Johnson Laboratory Accreditation, Inc. (PJLA) 755 W. Big Beaver, Suite 1325 Troy, Michigan 48084 The validity of this certificate is maintained through ongoing assessments based on a continuous accreditation cycle. The validity of this certificate should be confirmed through the PJLA website: www.pjlabs.com



Issue: 3/2023

Certificate of Accreditation: Supplement

Contract Testing Laboratories of America 151 E. 3450 N., Spanish Fork, UT 84660 Contact Name: Brescia Eppley Phone: 385-477-4999

Accreditation is granted to the facility to perform the following testing:

FIELD OF TEST	ITEMS, MATERIALS OR PRODUCTS TESTED	SPECIFIC TESTS OR PROPERTIES MEASURED	SPECIFICATION, STANDARD METHOD OR TECHNIQUE USED	RANGE (WHERE APPROPRIATE) AND DETECTION LIMIT
Microbiological F	Food, Cosmetic,	Aerobic Plate Count	AOAC 990.12	10 CFU/g
	Supplemental, and Nutraceutical	Escherichia Coli and Total Coliforms	AOAC 991.14	
		Enterobacteriaceae	AOAC 2003.01	
		Yeast and Mold	AOAC 2014.05	
		Escherichia Coli and Total Coliforms	AOAC 2018.13	
		Aerobic Plate Count	FDA BAM Ch. 3	
		Escherichia Coli and Total Coliforms	FDA BAM Ch. 4	
		Salmonella	FDA BAM Ch. 5	Presence or Absence
		Listeria Monocytogenes	FDA BAM Ch. 10	
		Staphylococcus Aureus	FDA BAM Ch. 12	
		Yeast and Mold	FDA BAM Ch. 18	100 CFU/g
		Yeast and Mold, Aerobic Plate Count	USP 2021	
		Escherichia Coli, Staphylococcus Aureus, Salmonella, Listeria Monocytogenes	USP 2022	Presence or Absence
		Aerobic Plate Count, Total Coliform, Yeast and Mold	USP <61>	100 CFU/g
		Escherichia Coli, Staphylococcus Aureus, Salmonella	USP <62>	Presence or Absence





Certificate of Accreditation: Supplement

Contract Testing Laboratories of America

151 E. 3450 N., Spanish Fork, UT 84660 Contact Name: Brescia Eppley Phone: 385-477-4999

Accreditation is granted to the facility to perform the following testing:

FIELD OF TEST	ITEMS, MATERIALS OR PRODUCTS TESTED	S granted to the facility to p SPECIFIC TESTS OR PROPERTIES MEASURED	SPECIFICATION, STANDARD METHOD OR TECHNIQUE USED	RANGE (WHERE APPROPRIATE) AND DETECTION LIMIT
Chemical F	Food, Cosmetic,	Arsenic, Cadmium, Lead,	USP <233>	LOD of As = 8 ppt
	Supplemental, and Nutraceutical	Mercury		LOD of Cd = 4 ppt LOD of Pb = 4 ppt
	Nutraccutical			LOD of $Hg = 4 \text{ ppt}$
		pH	USP <791>	LOQ = 0.01 %
		Caffeine	CTLA: M061	LOQ = 0.000 2907 %
		Cannabinoids: Total Cannabidiol (CBD) Total Tetrahydrocannabinol (THC) CBD CBDA Δ9-THC THCA Δ8-THC THCV CBDV	CTLA: M052	LOQ = 0.062 5 %
		CBDVA CBGA		
		CBG CBN		
		CBC		
		CBL		
		Mineral Analysis:	CTLA: M068	LOD for Mg, Fe, and
		Chromium (Cr) Iron (Fe)		Zn = 5.000 ppb LOD for $Cr = 0.500 \text{ ppb}$
		Magnesium (Mg)		LOD 101 C1 - 0.300 pp0
		Zinc (Zn)		

1. The presence of a superscript F means that the laboratory performs testing of the indicated parameter at its fixed location. Example: Outside Micrometer F would mean that the laboratory performs this testing at its fixed location.



CTLA ID	88169	Sample Name	30330 Omega	
Customer	Origin	Lot Number	2329301	
Date Received	11/3/2023	Date Complete	11/8/2023	
Customer Address:	151 E 3450 N, Spanish Fork, UT 84660			

ANALYSIS	SPECIFICATION	RESULT	METHOD	MDL	UNITS
Total Aerobic Microbial Count		<100	USP <2021>	100	cfu/g
Total Coliforms		<10	USP <2021>	10	cfu/g
E. coli		Absent	USP <2021>		
Salmonella		Absent	USP <2021>		
Staphylococcus aureus <2022>		Absent	USP <2021>		
Rapid Yeast Mold		<10	AOAC 2014.05	10	cfu/g
Arsenic		0.01	<usp 233=""></usp>	0.001	ppm
Cadmium		0.01		0.001	
Mercury		<0.001		0.001	
Lead		0.03		0.001	
Other Ingrdients: Cellulose		By Input			

Serving Size= 2

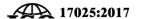
COA Note: Capsules

Approved By:

Date: 11/8/2023



Specifications provided by the Customer. Results with an asterisk (*) denote Specification should be reviewed by the Customer. This Certificate of Analysis represents the sample submitted and does not constitute a guarantee of quality for the entire product from which it was taken. The results are provided for the benefit of the Customer. Results using the "by input" method are calculated using information provided by the Customer. MDL = Method Detection Limit





EDITION No: 2



Version 4

456 Silver Creek Industrial Drive, TECUMSEH, ONTARIO N8N 4Y3 Tel:(519) 727-4618; Fax:(519)727-4619

CERTIFICATE OF ANALYSIS EMPTY HARD CAPSULES OF VEGETABLE ORIGIN

LOT No.: K2208002208		PRODUCT CODE	KC0-1000	SIZE:	0	
CAPSULE COLOR:	CAP -	NATURAL 1-0K	/ BODY -	NATURAL	1-0K	
PRINT: N/A	TEXT: N/A		INK COL	OR: N/A		
CapsCanada® capsules are preservative free and manufactured under strict cGMP conditions. All empty capsules are manufactured from pharmaceutical cellulose ethers, which are polymers derived from vegetable sources. Cellulose used to manufacture empty hard capsules of vegetable origin by CapsCanada® meet the current USP/NF and EP requeriments. CapsCanada® may blend pharmaceutical cellulose.						
(% Ingredients to % Cellulose)						
Сар		%	Body	%		
- ·						
the consistency of the finished produ	ct.	rovements, the color formula	ation reprasents target values only. The	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
Date of Manufacture: 2022-				Expiration Da		
CRITERIA	METH	OD / REFERENCE	SPECIFICATIONS		RESULTS	
ORGANOLEPTIC			Class amphy capaula shalls mast	ing the		
Physical Form	tr	itemal / Visual	Clean empty capsule shells meeti specified color and size		Passes	
Color /			Meets the specified requireme	nts	Passes	
Odor	Inter	nal / Organoleptic	Typical - Cellulose		Passes	
PHYSICAL			O Company of A	 - 	Passes	
Visual Defects	lr.	ternal / Visual	Conforms to the established AC	ils	107.5	
Average Capsule Weight		Internal	101,0 - 113,0 mg		4.3	
Loss on drying		USP	4.0% - 8.0%		Passes	
Disintegration		USP	N.M.T. 15 min		L93069	
ANALYTICAL			Meets USP Requirements		Passes	
Identification of HPMC		USP	N.M.T. 1.5% Transparent caps	ulae	1 23303	
Residue on Ignition *		USP	N.M.T. 6.0% colored capsule		Passes	
Arsenic *		SP (External)	N.M.T. 0.8 ppm		Passes	
Cadmium *		SP (External)	N.M.T. 0.5 ppm		Passes	
Lead *		SP (External)	N.M.T. 0.5 ppm		Passes	
Mercury *		SP (External)	N.M.T. 0.1 ppm		Passes	
Cobalt *		ISP (External)	N,M.T. 5.0 ppm		Passes Passes	
Vanadium *		ISP (External)	N.M.T. 10.0 ppm		Passes	
Nickel *		ISP (External)	N.M.T. 20.0 ppm		<10	
Total Aerobic Microbial Count		USP	N.M.T. 1000 cfu/g N.M.T. 100 cfu/g		<10	
Total Yeasts and Molds count		USP	Absence in 10 g		Absence	
Salmonella		USP	Absence in 1g		Absence	
Escherichia Coli		USP	Absence in 1g		Absence	
Staphylococcus aureus		USP	Absence in 1g		Absence	
Pseudomonas aeruginosa Capsules are certified as Kosher and i	Holal	1	· ·			
Storage Conditions: Temperature: 15°		midity: 35% - 70% RH				
*Reduced Frequency Testing.		⇒ No More Than				
reduced Frequency results	1 301310 1 0					
Auxin Epha						
Quality Control		advantus Cur-		Date:	2022-10-11	



Sample Information

CTLA ID: 83864

Date Received: 9/14/2023

Sample Name: 10449 Capsule, HPMC, 0, Clear, K-Cap

Lot Number: 47001

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	97.6	%
Total Aerobic Microbial Count	USP <2021>	100 Report	100	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Rapid Yeast and Mold	AOAC 2014.05	10 Report	<10	cfu/g
FTIR Spectra	FTIR	Report		

9/18/2023

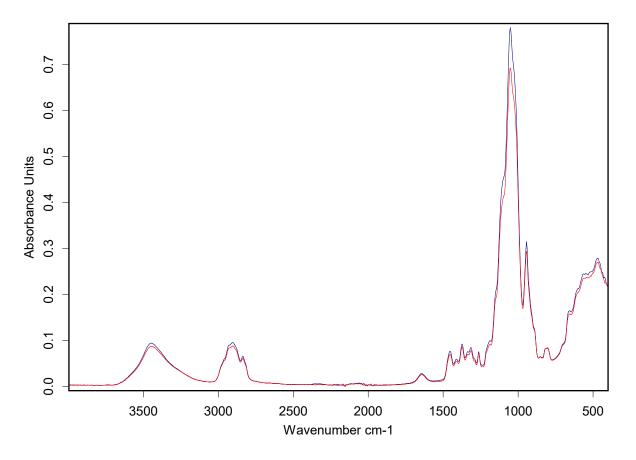
Specifications provided by the Customer. Results with an asterisk (*) Tenote Specifications should be reviewed by the Customer.

This Certificate of Analysis represents data for the sample submitted and does not constitute a guarantee of guality for the entire pro

This Certificate of Analysis represents data for the sample submitted and does not constitute a guarantee of quality for the entire product from which it was taken. These results are provided for the benefit of the Customer. MDL = Method Detection Limit. This document is not to be altered or reproduced except by the original authorizing body (CTLA)

151 E 3450 N, Ste 202, Spanish Fork, UT 84660

Search Library 9/15/2023 4:23:29 PM



Product Number	10449 CPS O K-Cap HPMC 12092 Standard 3
Entry No.	1401
Library name	ORIGIN STANDARDS LIBRARY.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Color	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
	976	10449 CPS O K-Cap HPMC 12092 Standard 3			

Cold	or	File	Path	Spectrum Type
		83864-ON 10449 Capsule, HPMC, 0, Clear, K-Cap (47001).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum



NANJING NUTRIHERB BIOTECH CO., LTD

Tel: (86)25 58862502 Http://www.naturemfg.com Mob: (86)17368474896 E-mail:sales@naturemfg.com

Certificate of Analysis

Product Name Gotu Kola Extract

Manufacturing Date 2023-06-18

Testing Date 2023-06-18

Botanical Source Centella Asiatica

Expiration Date

2025-06-17

Batch Number

20230618

Batch Quantity

25kgs

Analysis Item	Specification	Analysis Result	Method
Appearance	Light Yellow Powder	Complied	Visual
Odour	Characteristic	Characteristic	Organoleptic
Taste	Characteristic	Characteristic	Organoleptic
Loss on drying (%)	≤5.0	2.17	USP
Particle Size	100%Through 80 mesh	100%	USP < 786 >
Part Used	Whole	Complied	N/A
Method of Extraction	Soak and Extraction	Complied	N/A
Extraction Solvents	Food Grade Water&Alcohol	Complied	N/A
Identification	Complies by TLC	Complied	TLC
·			
Assay Total triterpenes (%)	≥ 80%	81.52%	HPLC
Asiaticoside(%)	≥15%	17.33%	HPLC
Madecassoside (%)	≥30%	35.81%	HPLC
Asiaticoside B (%)	≥20%	28.38%	HPLC
Heavy Metal Analysis			
Residual solvent ethanol	< 5000ppm	< 10ppm	GC
Lead (Pb)	≤2.0ppm	Complied	ICP-OES(CQ-MO-247)
Arsenic (As)	≤2.0ppm	Complied	ICP-OES(CQ-MO-247)
Mercury (Hg)	≤0.1ppm	Complied	ICP-OES(CQ-MO-247)
Cadmium (Cd)	≤1.0ppm	Complied	ICP-OES(CQ-MO-247)
Microbial Analysis			
Total Plate Count (CFU/g)	≤1,000	Complied	FDA-BAM
Yeast and Moulds (CFU/g)	≤100	Complied	FDA-BAM
Coliforms (MPN/g)	Not detected	Not detected	FDA-BAM
E. Coli (/g)	Not detected	Not detected	FDA-BAM
Salmonella (/25g)	Not detected	Not detected	FDA-BAM
Staphylococcus(/25g)	Not detected	Not detected	FDA-BAM

The raw plant material complies to EP regarding Heavy Metals ,mycotoxins and Pesticides Residues,according to EC396/2005,EC178/2006,and all related regulations regarding maximum residue levels (MRL) of pesticides in or on food products.

This product is not irradiated and is non-GMO according to EC 1829/2003 and EC 1830/2003 .

This products contains no ingredients which might pose a BSE/TSE risk.

Store in sealed containers at cool & dry place.

2 years if sealed and store away from direct sun light.

Tested by:

Safety Information

Storage

Shelf Life







Sample Information

CTLA ID: 82324

Date Received: 8/25/2023

Sample Name: 11040 DRM Gotu Kola 80% Triterpenes

Lot Number: 49969

Customer: Origin Nutraceutical

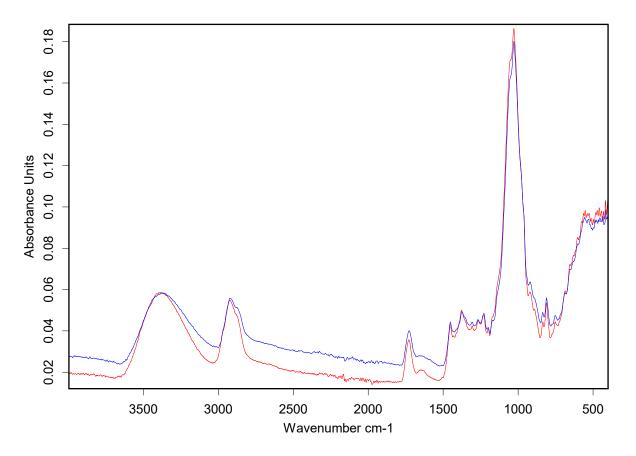
Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	94.8	%
Total Aerobic Microbial Count	USP <2021>	100 Report	100	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Rapid Yeast and Mold	AOAC 2014.05	10 Report	<10	cfu/g

8/29/2023

Specifications provided by the Customer. Results with an asterisk (A) The Customer Specifications should be reviewed by the Customer.

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Search Library 9/14/2023 9:47:36 AM



Product Number	ON 11040 DRM Gotu Kola 80% Triterpenes (3
Entry No.	2261
Library name	ORIGIN STANDARDS LIBRARY.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Co	olor	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
		948	ON 11040 DRM Gotu Kola 80% Triterpenes (30968) Standard 3			

Colo	r	File	Path	Spectrum Type
		82324-ON 11040 (DRM Gotu Kola 80% Triterpenes (49969) C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\Data\MEAS Q		Query Spectrum



Sample Information

CTLA ID: 83861

Date Received: 9/14/2023

Sample Name: 11040 Gotu Kola 80% Triterpenes

Lot Number: 49969

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
Triterpene	HPLC	0.002 80	81.1	%
Gotu Kola Extract	HPTLC	Report	Characteristic	

HPTLC: Fingerprint of test solution is consistent with reference standard Centella Asiatica (gotu kola extract).

10/5/2023

Specifications provided by the Customer. Results with an asterisk (A) Tenote Specifications should be reviewed by the Customer. This Certificate of Analysis represents data for the sample submitted and does not constitute a guarantee of quality for the entire product from which it was taken. These results are provided for the benefit of the Customer. MDL = Method Detection Limit. This document is not to be altered or reproduced except by the original authorizing body (CTLA)



Les Dérivés Résiniques et Terpéniques

30 rue Gambetta - BP 90206 - 40105 DAX Cedex - FRANCE Tél: +33 5 58 56 62 00 - eMail: drtsales@drt.fr - web: www.drt.fr SAS au capital de 19 961 200 € - Siret 985 520 154 00016 R.C.S. DAX - APE 2014Z - n° TVA: FR 86985520154

Certificate of Analysis

Date: 2023/08/24

Material Reference: 8200.831 OLIGOPIN

Customer Material Reference: 11041 OLIGOPIN PUREXPERT Lot Number: 1000211631 Quantity: 2.000 KG

Manufacturing Date: 2023/03/13 Best Before Date: 2028/03/11 Customer Number: 1000008205 Order Reference: 62478

Customer Order Reference: 8210 Delivery Note Number: 80142092

Registration: EL168CQ

Customer: ORIGIN NUTRACEUTICAL, INC.

151 E 3450 N

84660 SPANISH FORK USSFU UT US USA

Email: purchasing@originnutra.com

Extract type: French Maritime Pine bark ext.

. (Pinus pinaster)

Plant Extract Ratio: 1000:1

Extraction type: Please refer to flowchart Additive in the extract: No additives

Restriction on use : No Pesticides : EU Pines

. . Pine bark extract not listed within Regulation (EC) 396/2005. By default, Ph. Eur is the reference

Contaminants : EU

.. Dioxins, PAH, Mycotoxins, Heavy Metals: Complies with Regulation (EC) No 1881/2006 and its successive modifications

Shelf life: 5 years
Country of origin: France
Professional use: Yes

... Product intended to be used as food supplement ingredient

Characteristic	Unit	Value	Type Value	Lower Limit	Upper Limit	Method of Analysis
Appearance	- (CONFORM				Visual
Color: brownish red	- (CONFORM				DRT 6903
Odour: Woody caract.	- (CONFORM				DRT 6917
pH 20°C (4% deionised w.)	-	2.9		2,5	4,5	DT_3301
Insoluble in water (20°C)	%	1.6			5,0	DRT 3882
Insoluble in THF	%	0.4			1,0	DRT 3884
Sulphuric ashes	%	0.01			0,50	PH.EUR 2.4.14
Absorbance UV at 281 nm	-	150		145	175	PH.EUR.2.2.25
Procyanidin - Porter	%	72	65			DT_2152

^{*} Annual monitoring

DRT methods : available upon request

This document has been produced electronically and is effective without a signature

This product was produced in accordance with the guidelines and monitored in every manufacturing stage



Les Dérivés Résiniques et Terpéniques

30 rue Gambetta - BP 90206 - 40105 DAX Cedex - FRANCE Tél: +33 5 58 56 62 00 - eMail: drtsales@drt.fr - web: www.drt.fr SAS au capital de 19 961 200 € - Siret 985 520 154 00016 R.C.S. DAX - APE 2014Z - n° TVA: FR 86985520154

Certificate of Analysis

Date: 2023/08/24

Material Reference: 8200.831 OLIGOPIN

Customer Material Reference: 11041 OLIGOPIN PUREXPERT Lot Number: 1000211631 Quantity: 2.000 KG

Manufacturing Date: 2023/03/13 Best Before Date: 2028/03/11 Customer Number: 1000008205

Order Reference: 62478 Customer Order Reference: 8210 Delivery Note Number: 80142092

Registration: EL168CQ

Customer: ORIGIN NUTRACEUTICAL, INC.

151 E 3450 N

84660 SPANISH FORK USSFU UT US USA

Email: purchasing@originnutra.com

Characteristic	Unit	Value	Type Value	Lower Limit	Upper Limit	Method of Analysis
Procyanidin content (GPC)	%	71.7		67,0		DRT 1610
Monomeres	%	28.3	28,0			DRT 1610
Dimeres	%	17.4	18,0	15,0		DRT 1610
Trimeres + Tetrameres	%	54.3	50,0			DRT 1610
Water (extract. solvent)	%	3.2			5,0	PH.EUR 2.2.32
Ethyl acetate	mg/kg	< 10			10	DT_1251
Co-solvent: alpha pinene	mg/kg	< 5			5	DRT 1253
Lead content	mg/kg	< 0.13			3,00	EXTERNE
Arsenic content	mg/kg	< 0.13			0,50	EXTERNE
Mercury content	mg/kg	< 0.01			0,10	EXTERNE
Cadmium content (mg/kg)	mg/kg	0.03			1,00	EXTERNE
TAMC (<1000 CFU/g)	-	CONFORM				PH.EUR.2.6.12
TYMC (<100 CFU/g)	-	CONFORM				PH.EUR.2.6.12
Salmonella (abs CFU/25g)	-	CONFORM				PH.EUR.2.6.31
Escherichia C. (abs CFU/g)	-	CONFORM				PH.EUR.2.6.13
Gram- bact (<100 CFU/g)	-	CONFORM				PH.EUR.2.6.31
Staph.Aureus (abs CFU/g)	-	CONFORM				PH.EUR.2.6.13

^{*} Annual monitoring

DRT methods : available upon request

I his document has been produced electronically, and is effective without a signature. This product was produced in accordance with the guidelines and monitored in every manufacturing stage.

Page 2 of 3



Les Dérivés Résiniques et Terpéniques

30 rue Gambetta - BP 90206 - 40105 DAX Cedex - FRANCE Tél: +33 5 58 56 62 00 - eMail: drtsales@drt.fr - web: www.drt.fr SAS au capital de 19 961 200 € - Siret 985 520 154 00016 R.C.S. DAX - APE 2014Z - n° TVA: FR 86985520154

Certificate of Analysis

Date: 2023/08/24

Material Reference: 8200.831 OLIGOPIN

Customer Material Reference: 11041 OLIGOPIN PUREXPERT Lot Number: 1000211631 Quantity: 2.000 KG

Manufacturing Date: 2023/03/13 Best Before Date: 2028/03/11 Customer Number: 1000008205

Order Reference: 62478
Customer Order Reference: 8210
Delivery Note Number: 80142092

Registration: EL168CQ

Customer: ORIGIN NUTRACEUTICAL, INC.

151 E 3450 N

84660 SPANISH FORK USSFU UT US USA

Email: purchasing@originnutra.com

Characteristic	Unit	Value	Type Value	Lower Limit	Upper Limit	Method of Analysis
Liste. Mono. (<100 CFU/g)	-	CONFORM				EXTERNE
Pseudomonas.Aerug (abs/g)	-	CONFORM				PH.EUR.2.6.13
Clostridium (abs CFU/g)	-	CONFORM				PH.EUR.2.6.13
Candida Albi. (abs CFU/g)	-	CONFORM				PH.EUR.2.6.13
Aflatoxin B1 (<2 μg/kg)*	-	CONFORM				EXTERNE
Aflat.B1+B2+G1+G2 (<4 μg/kg)*		CONFORM				EXTERNE
Ochratoxine A (<2 µg/kg)*	(-)	CONFORM				EXTERNE
Pesticide content*	-	CONFORM				PH.EUR.2.8.13
Benzo(a) pyren (<10 μg/kg)*	-	CONFORM				EXTERNE
Sum of PAH (<50 µg/kg)*	-	CONFORM				EXTERNE
Dioxins (<0,75 pg/g)*	-	CONFORM				EXTERNE
Sum of PCBs (<40 ng/g)*	-	CONFORM				EXTERNE
Dioxins + PCB (<1,25 pg/g)*	-	CONFORM				EXTERNE
Product File Validation	-	CONFORM				PROD

^{*} Annual monitoring

DRT methods : available upon request



Sample Information

CTLA ID: 83603

Date Received: 9/11/2023

11041 DRM Oligopin PUR'expert Sample Name:

50293 Lot Number:

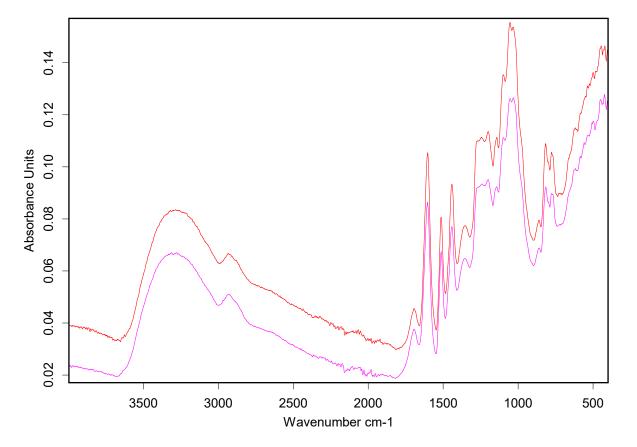
Origin Nutraceutical Customer:

Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	94.5	%
Total Aerobic Microbial Count	USP <2021>	100 Report	100	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Ranid Yeast and Mold	AOAC 2014 05	10 Report	<10	cfu/a

9/13/2023

Specifications provided by the Customer. Results with an asterisk of the Specifications should be reviewed by the Customer. This Certificate of Analysis represents data for the sample submitted and does not constitute a guarantee of quality for the entire product from which it was taken. These results are provided for the benefit of the Customer. MDL = Method Detection Limit. This document is not to be altered or reproduced except by the original authorizing body (CTLA)

Search Library 9/14/2023 9:43:22 AM



Product Number	ON 11041 DRM Oligopin PUR Expert (31406)
Entry No.	2319
Library name	ORIGIN STANDARDS LIBRARY.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Color	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
	945	ON 11041 DRM Oligopin PUR Expert (31406) Standard 1			

Color	File	Path	Spectrum Type
	83603-ON 11041 DRM Oligopin PURexpert (50293).2	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\Data\MEAS	Query Spectrum



Sample Information

CTLA ID: 83862

Date Received: 9/14/2023

Sample Name: 11041 Oligopin PUR'expert

Lot Number: 50293

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
Procyanidin	HPLC	0.002 67	69.3	%
French Maritime Pine Bark Extract	HPTLC	Report	Characteristic	

HPTLC: Fingerprint of test solution is consistent with reference standard Pinus Maritima (pine bark extract).

10/5/2023

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5625 Daniels Street, Chino, CA 91710

Tel: 909-628-2600 Fax: 909-628-8110 www.AuNutra.com

Certificate of Analysis

Product Name	Deodorized Garlic P.E.	Botanical Source	Allium Sativum L.
Part Used	Garlic Bulb	Country of Origin	China
Solvent Used	Water	Carrier Used	None
Batch Number	GA230212	Manufacture Date	02.12.2023

ANALYSIS	SPECIFICATION	RESULTS	TEST METHODS
Assay	NLT1.00% Allicin	1.20%	HPLC
	NLT2.00% Alliin	2.52%	HPLC
Chemical Physical Control			
Characters/Appearance	Fine Powder	Conforms	Visual
Color	White to Pale Yellow	Conforms	Visual
Odor	Deodorized	Conforms	Organoleptic
Taste	Characteristic	Conforms	Organoleptic
Mesh Size/Sieve Analysis	NLT90% through 80 mesh	Conforms	80 Mesh Sieves
Loss on drying	NMT10%	4.80%	USP<731>
Heavy Metals	NMT10ppm	Conforms	ICP-MS
Arsenic (As)	NMT2ppm	0.310ppm	ICP-MS
Lead (Pb)	NMT2ppm	0.900ppm	ICP-MS
Mercury (Hg)	NMT2ppm	0.060ppm	ICP-MS
Cadmium (Cd)	NMT2ppm	0.420ppm	ICP-MS
Pesticide Residue	Meets the requirement	Conforms	USP<561>
Solvents Residue	NMT5,000ppm	Conforms	USP<467>
Sterilization Method	High Temperature Pressure	Conforms	PSL
Microbiology Control			
Total Plate Count	NMT10,000cfu/g	650cfu/g	AOAC
Total Yeast & Mold	NMT1,000cfu/g	40cfu/g	AOAC
E.Coli	Negative	Conforms	AOAC
Salmonella	Negative	Conforms	AOAC
Staphylococcus	Negative	Conforms	AOAC

Package & Storage	Packed in paper-drums and two plastic bags inside. Store in a well-closed
	container away from moisture
Shelf Life	Three years if sealed and store away from direct sun light.
	SC: TH-GE220215

Approved by QA/QC PW Date: 03.15.2023



Sample Information

CTLA ID: 73994

Date Received: 5/11/2023

Sample Name: 10620 DRM Garlic 1% Allicin/2% Allin

Lot Number: 48251

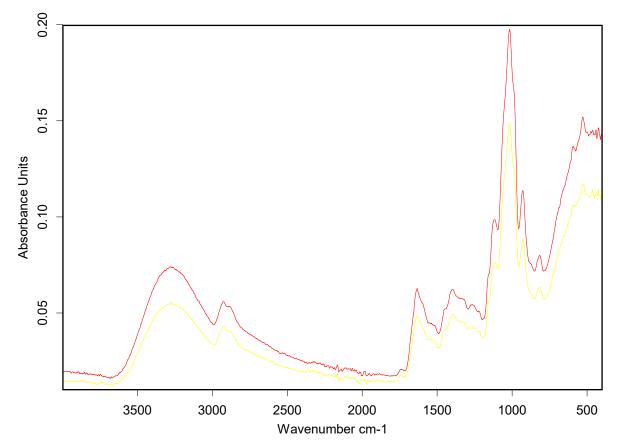
Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	95.5	%
Total Aerobic Microbial Count	USP <2021>	100 Report	1,200	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Rapid Yeast and Mold	AOAC 2014 05	10 Report	<10	cfu/a

5/15/2023

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Search Library 9/15/2023 4:13:37 PM



Product Number	ON 10620 DRM Garlic 1% Allicin 2% Allin (482
Entry No.	3176
Library name	ORIGIN STANDARDS LIBRARY.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Color	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
	970	ON 10620 DRM Garlic 1% Allicin 2% Allin (48251) Standard 2			

Color	File	Path	Spectrum Type
	83860-ON 10620 Garlic 1% Allicin 2% Allin (48251).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum



Sample Information

CTLA ID: 83860

Date Received: 9/14/2023

Sample Name: 10620 Garlic 1% Allicin/2% Allin

Lot Number: 48251

Customer: Origin Nutraceutical

Analysis	Method	MDL	Specification	Result	Units
ID	FTIR		Report	97.0	%
FTIR Spectra	FTIR		Report	Attached	
Garlic Extract	HPTLC		Report	Characteristic	mg/g
Allicin	HPLC	0.002	1	1.06	%

HPTLC: Fingerprint of test solution is consistent with reference standard Allium Sativum (garlic bulb extract).

10/5/2023

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Page 1 of 2

Evonik Rexim S.A.S., 80400 Ham, France

Certific	ate of Analysis
Date	Sep 2, 2021

Product

AvailOm® 50 High EPA

CAS nº127964-03-8

Manufactured according to HACCP 90258091A2

Batch

8095201003

Manufacturing date

Oct 01, 2020

Recommended date of

Oct 01, 2024

retesting

			•	Specification	
Property	Test method	Unit	Value	Min.	Max.
Appearance			Complies	Yellowis powder	sh to brownish
Identification			Complies	Retention	on time EPA A
Sum EPA + DHA as FFA (% w/w)		%	53	46	57
EPA as FFA (% w/w)		%	36	31	37
DHA as FFA (% w/w)		%	16	15	20
L-Lysine (% w/w)		%	32	30	34
Other fatty acids (*)			Approx. 9 - 24 (%W/W)	Approx. W)	9 - 24 (%W/
TOTOX Value (AV + 2PV)			< 3		26
Water (% w/w)		%	0,2		1,0
Residue on ignition (% w/w)		%	0,1		0,5
Sum Dioxins&Furans (PCDDs and PCI	DFs)(*)		<=1.75		1,75
Dioxin-like PCBs (pg WHO-TEQ/g) (*)			<=3		3
Sum PCBs28,52,101,138,153,180(ng/s	g)(*)		<=90		90
Sum dioxins&Furans+Dioxin-like PCBs	; (*)		<=3		3

Tel.: +33 3 23 81 00 25 Fax: +33 3 23 81 11 12 www.evonik.com



Product

AvailOm® 50 High EPA CAS n°127964-03-8 Manufactured according to HACCP 90258091A2

Date

Page 2 of 2 Sep 2, 2021

Batch

8095201003

				Sp	ecification
Property	Test method	Unit	Value	Min.	Max.
Benzo(a)pyrene (µg/kg) (*)			<=2.0		2,0
Sum benzo(a)pyrene & 3 other PAH (*)			<=10		10
Arsenic (*)		ppm	<=0.1		0,1
Cadmium (*)		ppm	<=0.1		0,1
Mercury (*)		ppm	<=0.1		0,1
Lead (*)		ppm	<=0.1		0,1
Iron (*)		ppm	<=30		30
Total aerobic germ at 30°C (CFU) (*)		/g	<=1000		1000
Yeasts and moulds (CFU) (*)		/g	<=100		100
Escherichia coli (CFU/10g) (*)			0		0
Pseudomonas aeruginosa (CFU) (*)		/g	0		0
Salmonella Species (CFU/25g) (*)			0		0
Staphylococcus aureus (CFU) (*)		/g	0		0
Guaranteed by regular monitoring (*):			Conform		

UD date

Aug 31, 2021

UD name

A29368 Audrey Anastase

UD date = Release date

UD name = Authorized personnel of the quality unit

Evonik Rexim S.A.S.; 33, Rue de Verdun; 80400 Ham; France;

Tel. +33 3 23 81 00 25

Product complies to the specification.

This document has been produced electronically and is effective without a signature

*** End ***



Sample Information

CTLA ID: 83243

Date Received: 9/6/2023

Sample Name: 11042 DRM AvailOm High EPA (30% EPA, 15% DHA)

Lot Number: 50242

Customer: Origin Nutraceutical

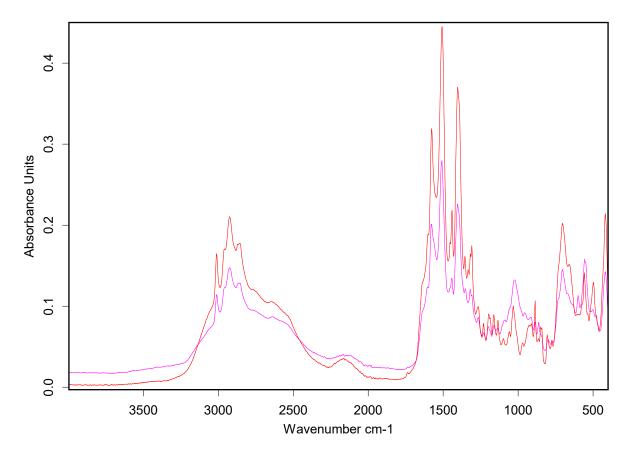
Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	86.5	%
Total Aerobic Microbial Count	USP <2021>	100 Report	<100	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Rapid Yeast and Mold	AOAC 2014.05	10 Report	<10	cfu/g

9/8/2023

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Search Library 9/14/2023 9:44:55 AM



Product Number	ON 11042 DRM AvailOm High EPA (30% EPA,
Entry No.	2246
Library name	ORIGIN STANDARDS LIBRARY.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Color	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
	865	ON 11042 DRM AvailOm High EPA (30% EPA, 15% DHA) (30626) Standard 1			

Color	File	Path	Spectrum Type
	83243-ON 11042 DRM AvailOm High EPA (30% EPA, 15% D C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\Data\MEAS		Query Spectrum



Sample Information

CTLA ID: 83863

Date Received: 9/14/2023

Sample Name: 11042 AvailOm High EPA (30% EPA, 15% DHA)

Lot Number: 50242

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
Total Omega 3	GC FID	0.10000 Report	70.643	%
DHA/EPA Combo				
EPA	GC-FID	0.10000 30	50.895 *	%
DHA	GC-FID	0.10000 15	19.590	%

9/19/2023

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