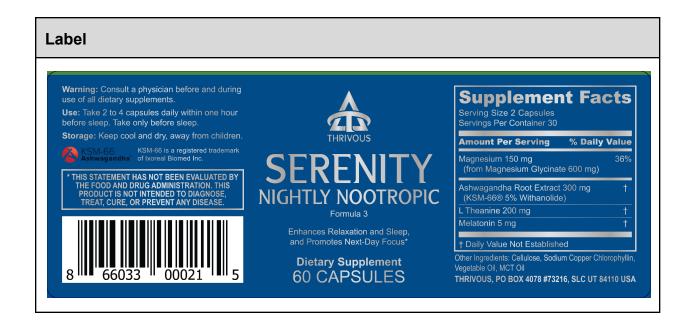


CERTIFICATE OF ANALYSIS AND QUALITY

Product	Serenity Nightly Nootropic
sku	SERENITY
Barcode	866033000215
Formula	3
Date	10 August 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
Excipient MCT Oil Certificate of Analysis from Supplier
Excipient MCT Oil Certificate of Analysis from Third-Party Testing
Excipient Vegetable Oil Certificate of Analysis from Supplier
Excipient Vegetable Oil Certificate of Analysis from Third-Party Testing
Ashwagandha Certificate of Analysis from Supplier
Ashwagandha Certificate of Analysis from Third-Party Testing
L Theanine Certificate of Analysis from Supplier
L Theanine Certificate of Analysis from Third-Party Testing
Magnesium Glycinate Certificate of Analysis from Supplier
Magnesium Glycinate Certificate of Analysis from Third-Party Testing
Melatonin Certificate of Analysis from Supplier
Melatonin Certificate of Analysis from Third-Party Testing



10 August 2023

RE: Letter of Guarantee for Thrivous Serenity Nightly Nootropic

To whom it may concern,

The undersigned, Lincoln Cannon LLC DBA Thrivous ("Thrivous"), hereby guarantees as follows regarding Serenity Nightly Nootropic ("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.
- 3. 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.

LITAH DEPARTMENT O	F 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:

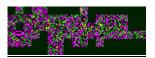
ISO/IEC 17025:2017

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:



Tracy Szerszer President

Perry Johnson Laboratory Accreditation, Inc. (PJLA) 755 W. Big Beaver, Suite 1325 Troy, Michigan 48084 Initial Accreditation Date: Issue Date: Expiration Date:

March 31, 2021 March 24, 2023 June 30, 2025

Accreditation No.: Certificate No.: 102267 L23-261

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	Mercury		LOD of $Cd = 4 ppt$
	Nutraceutical			LOD of Pb = 4 ppt
		***	LICE 701	LOD of Hg = 4 ppt
		рН	USP <791>	LOQ = 0.01 %
		Caffeine	CTLA: M061	LOQ = 0.000 2907 %
		Cannabinoids:	CTLA: M052	LOQ = 0.062 5 %
		Total Cannabidiol (CBD)		
		Total		
		Tetrahydrocannabinol		
		(THC)		
		CBD		
		CBDA		
		Δ9-THC		
		THCA		
		Δ8-THC THCV		
		CBDV		
		CBDVA	7	
		CBGA		
		CBG		
		CBN		
		CBC		
		CBL		
		Mineral Analysis:	CTLA: M068	LOD for Mg, Fe, and
		Chromium (Cr)		Zn = 5.000 ppb
		Iron (Fe)		LOD for $Cr = 0.500 \text{ ppb}$
		Magnesium (Mg)		
		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	80270	Sample Name	Serenity
Customer	Thrivous	Lot Number	2320001
Date Received	7/31/2023	Date Complete	8/3/2023
Customer Address:	PO Box 4078 #73216, SLC Utah 84110		

ANALYSIS	SPECIFICATION	RESULT	METHOD	MDL	UNITS
Rapid Complete Micro					
Total Plate Count	Report	900	USP<2021>	100	cfu/g
Total Coliforms	Report	<10	USP<2021>	10	cfu/g
E. coli	Report	Absent	USP<2022>		
Salmonella	Report	Absent	USP<2022>		
S. aureus	Report	Absent	USP<2022>		
Yeast & Mold	Report	<10	AOAC 2014.05	100	cfu/g
Heavy Metal					
Arsenic	Report	0.035	USP <2232>	0.001	ppm
Cadmium	Report	0.016	USP <2232>	0.001	ppm
Mercury	Report	<0.001	USP <2232>	0.001	ppm
Lead	Report	<0.001	USP <2232>	0.001	ppm
Magnesium (from Magnesium Glycinate 600mg)	Report	150	By Input		mg
Ashwagandha Root Extract (KSM-66® 5% Withanolide)	Report	300	By Input		mg
L-Theanine	Report	200	By Input		mg
Melatonin	Report	5			mg
Other Ingredients: Cellulose, Sodium Copper Chlorophylln, Vegetable Oil, MCT Oil					

COA Note:

Approved By:

Date: 8/3/2023



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

151 E 3450 N, Ste 202 Spanish Fork, UT 84660 (385) 477-4999



ISO 17025 Accreditation No: 102267

CERTIFICATE OF ANALYSIS

Issued date: Apr. 12, 2022

Product Description

Empty Hard Capsule From Hypromellose (HPMC)

Customer:

SUHEUNG-AMERICA

Manufacturing Date:

Jul. 23, 2021

Capsule Type:

EMBO CAPS® VG-PRO "Kosher and Halal Certified" Expiration Date :

Jul. 22, 2026

Capsule Item No.: VP0L17L17

Quantity:

Lot Number:

VP0L17L17 - 13908

7.020.000 PCS (108 Cartons)

Product Size :

Color

Carton No.: 1 - 108

Product Code:

CAP

BODY

L17 (DK. GREEN TR.)

L17 (DK. GREEN TR.)

Composition

Cap Sodium Copper Chlorophyllin 0.4000% Body Sodium Copper Chlorophyllin 0,4000% Hypromellose qsp 100 Hypromellose qsp 100

Anglytical Results

Test Items		<u>Unit</u>	<u>Standard</u>	Results	Test Method
Length	Cap	mm	10.3 ~ 11.1	10.7	In-house Spec.
	Body	mm	18.0 ~ 18.8	18.4	In-house Spec.
Weight		mg	84.6 ~ 105.4	97.2	In-house Spec.
ldentification	of Hypromellose*		Positive	Positive	USP/EP
	of Colorant*		Positive	Positive	USP/EP
Disintegration		min/sec	NMT 15	13'50"	USP/EP
Loss on Drying	3	%	3.0 ~ 7.0	4.9	USP/EP
Residue on 1gn	ition (Ash)*	%	NMT 3.0	1.2	USP/EP
TAMC		CFU/g	NMT 500	<10	USP/EP
TYMC		CFU/g	NMT 100	<10	USP/EP
E.Coli			Negative/10g	Negative	USP/EP
Salmonella			Negative/10g	Negative	USP/EP
Staphylococcus	s aureus		Negative/10g	Negative	USP/EP
Pseudomonas a	eruginosa		Negative/10g	Negative	USP/EP

^{*} Reduced frequency testing

Elemental Impurities

Suheung Empty Hard Capsules comply with ICH Q3D for elemental impurities and meeting below acceptance level.

Elemental impurities test results based on continuous monitoring program.

<u>Elemente</u>	Unit	Acceptance level
Arsenic	ppm	NMT 1.0
Lead	ppm	NMT 1.0
Cadmium	ppm	NMT 0.5
Mercury	ppm	NMT 0.1
Cobalt	ppm	NMT 5
Vanadium	ppm	NMT 10
Nickel	ppm	NMT 20

Ingredients List

Ingredients Name	E Nr	Cl Nr	Function	Regulatory Reference
Sodium Copper Chlorophyllin	E141	75815	Colorant	(EU) 231/2012, 21 CFR
Hypromellose			Structure	EP, JP, USP/NF

⁻ Storage Conditions: recommended 15-25°C and 35-65%RH

* ORDER NO. HC6701

Quality Assurance Manager

⁻ We, Suheung Co., Ltd., hereby certify that the Empty Hard Capsules manufactured by us are free from Preservatives and no Ethylene Oxide or/and Irradiation Treatment.



Sample Information

CTLA ID: 70053

Date Received: 3/15/2023

Sample Name: 10448 CPS Capsule, HPMC, 0, Chlorophyll

Lot Number: 47445

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID	FTIR	Report	98.1	%
FTIR Spectra	FTIR	Report		
Rapid Complete Micro				
Total Plate 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

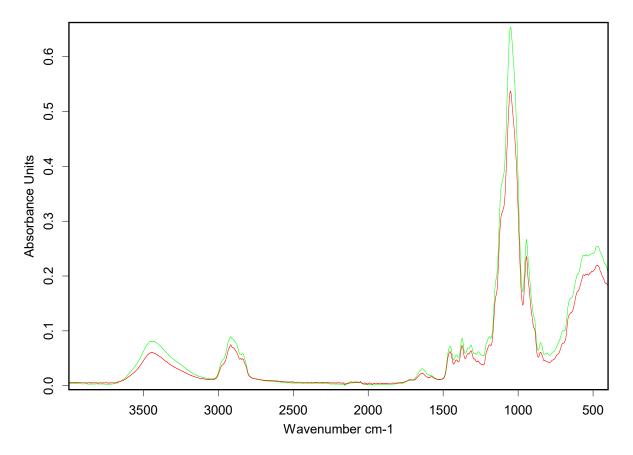
Amended report: Test added

3/23/2023

Specifications provided by the Customer. Results with an asterisk of denote Specifications should be reviewed by the Customer.

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Search Library 3/17/2023 3:18:34 PM



Product Number	ON 10448 CPS Capsule HPMC 0 Chlorophyll 3
Entry No.	2448
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
	981	ON 10448 CPS Capsule HPMC 0 Chlorophyll 35035 Standard 3			

Color	File	Path	Spectrum Type
	70053-ON 10448 CPS Capsule, HPMC, 0, Chlorophyll (4744	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\Data\MEAS	Query Spectrum



Material#: 26154

Material Name: MCT Oil

Allergens: None

Batch: F060822-26154

Mfg. Date: 06/08/2022

Retest Date: 06/08/2025

Characteristic	Units	Value	Specifications	Test Method		
Acid Value	mg KOH/g	0.02	Max 0.1	AOCS Te 2a-64		
Saponification Value	mg KOH/g	334	Min. 325 Max. 345	AOCS Tl 1a-64		
Iodine Value	$gI_2/100g$	0.03	Max. 0.5	AOCS Tg 1a-64		
Hydroxyl Value	mg KOH/g	3	Max. 5	AOCS Cd 13-60		
Peroxide Value	meq/kg	NIL	Max. 1	AOCS Cd 8-53		
C8 Caprylic Acid	%	59.1	Min. 55 Max. 65	ISO 12966-2		
C10 Capric Acid	%	40.7	Min. 35 Max. 45	ISO 12966-2		
C12 Lauric Acid	%	0.1	Max. 1	ISO 12966-2		
Chromium	ppm	< 0.05	Max. 0.05	ICPMS		
Copper	ppm	< 0.1	Max. 0.1	ICPMS		
Lead	ppm	< 0.1	Max. 0.1	ICPMS		
Nickel	ppm	< 0.1	Max. 0.1	ICPMS		
Tin	ppm	< 0.1	Max. 0.1	ICPMS		
*Typical Microbial D	ata:					
Aerobic Plate Count	cfu/g	<10	Max. 5000	FDA-BAM		
E. Coli	cfu/g	<10	Max. 10	FDA-BAM		
Yeast & Mold	cfu/g	<10	Max. 100	FDA-BAM		
Salmonella cfu/g Negative Negative FDA-BAM * testing is done on an annual basis						

Results Reviewed by	Results Verified by
Signature: Lexi Zaremba	Signature Outh Kingsburg
Print Name: Lexi Zaremba	Print Name: Arielle Klingenberg
Date: 06/08/2022	Date: 6/29/22



Sample Information

CTLA ID: 69969

Date Received: 3/14/2023

Sample Name: 10842 MCT Oil 100%

Lot Number: 44891

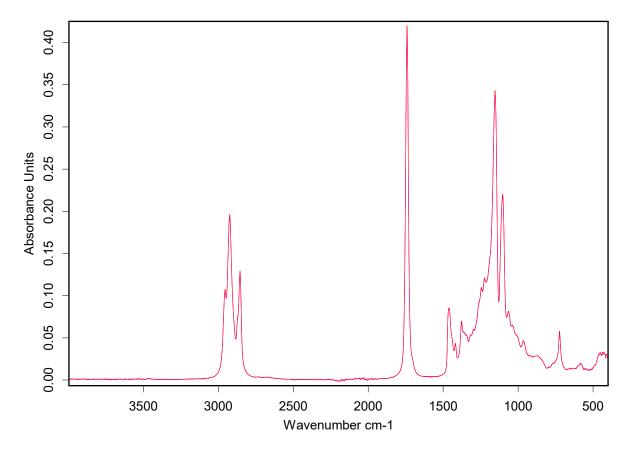
Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID, Rapid Complete Micro Combo				
ID	FTIR	Report	99.6	%
Total Plate 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		

3/16/2023

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)

Search Library 3/15/2023 10:50:35 AM



Product Number	ON 10842 LIQ MCT Oil 100% 18053 Standard
Entry No.	1721
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
	996	ON 10842 LIQ MCT Oil 100% 18053 Standard 1			

Color	File	Path	Spectrum Type
	69969-ON 10842 MCT Oil 100% (44891).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum



SM Ingredients, Inc. 27132 B Paseo Espada Suite 1221 San Juan Capistrano, CA 92675

Phone: 714-602-7006 Fax: 714-685-7723

CERTIFICATE OF ANALYSIS

VEGATEX

Product Name:

Vegatex™

Lot Number:

22-7703

Date of Manufacture:

April 28, 2022

Date of Retest/Expiration:

April 27, 2025

Manufacturer:

LipidChem SDN BHD

Country of Origin:

Malaysia

TEST	TEST SPECIFICATION	TEST RESULT
Appearance	White to Off White Powder	Complied
Free Fatty Acid (%)	≤ 0.25	0.06
lodine Value (gl2/100g)	≤ 15.0	13.98
Slip Melting Point (°C)	57 min	58.2
Moisture Content (%)	≤ 0.25	0.04

Note: This analysis is only fit for this product lot.

Authorized Signature,

Stephen Marantette

Rev.000

Date Issued: July 22, 2022

Vegatex[™] COA Lot # 22-7703 Page 1 of 1



Sample Information

CTLA ID: 69970

Date Received: 3/14/2023

Sample Name: 10867 EXC Vegatex

Lot Number: 46244

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID	FTIR	Report	99.3	%
FTIR Spectra	FTIR	Report		

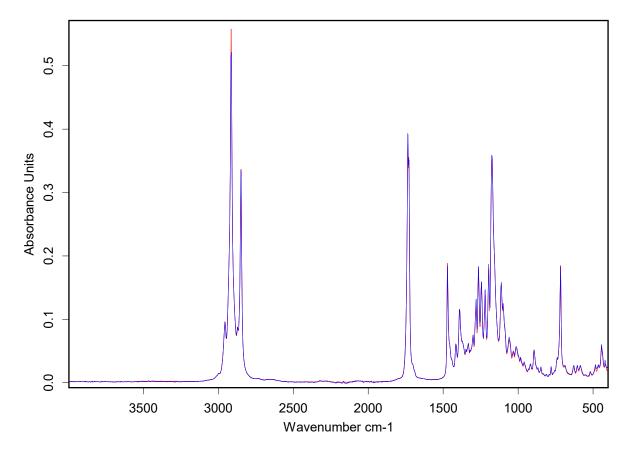
3/16/2023

Specifications provided by the Customer. Results with an asterisk (*) denote 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 3/15/2023 11:00:52 AM



Product Number	ON 10867 DRM EXC Vegatex Vegetable Oil Po
Entry No.	2841
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
	993	ON 10867 DRM EXC Vegatex Vegetable Oil Powder (43744) Standard 2			

Color	File	Path	Spectrum Type
	69970-ON 10867 EXC Vegatex (46244).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum



SHRI KARTIKEYA PHARMA



DEPARTMENT OF QUALITY CONTROL

Certificate Of Analysis Product Name: KSM-66 Ashwagandha Root Extract-Vegan (Withania Somnifera) Issued To: Origin Nutraceutical, Inc. Batch No: KSM/VG/23/S037 Mg. Date: Jan 2023 Exp. Date: Dec 2025 No. Tests Certificate Of Analysis Stability: 3 Years from the date of manufacture. Country of Origin: India Report No: SKP/S037 Lab Ref No: QC/037 Result Limits (specification)

No.	Tests	Result	Limits (specification)
1	Description	Light Yellowish Brown Powder	
2	Plant Part Used	Roots	
3	Identification	Six spots at Rf of 0 61, 0.53, 0.46, 0.35, 0.28, 0.23	To Pass The Test (TLC)
4	Physico-chemical analysis		
	Moisture Content (%)	3.26	<5.0
	Ash Content	5.32	<15.0
	Acid insoluble ash (%w/w)	2 07	<5.0
	pH of 5% w/v solution	4 61	4.0-6.0
5	Partical Size		Name of the second
	Passing through 40 mesh sieve	Pass	>95%(w/w)
	Passing through 80 mesh sieve	Pass	>85%(w/w)
6	Density		
	Bulk density (g/cc)	0.57	0.2-0.6
	Tapped bulk density (g/cc)	0.76	0.4-0.8
7	Heavy Metals		
	Lead	BDL	<1ppm
	Cadmium	BDL	<1ppm
	Arsenic	BDL	<1ppm
	Mercury	BDL	<0.1ppm
8	Microbiological analysis as per USP		
	Total viable aerobic count	<10 ⁴ cfu/g	<10 ⁴ cfu/g
	Total enterobacteriaceae	<10 ² cfu/g	<10 ² cfu/g
	Total fungal count	<10 ³ cfu/g	<10 ³ cfu/g
9	Test for specific pathogens		
	E.Coli	Absent	Absent
	Salmonella sp. (cfu/10g)	Absent	Absent
	S.aureus (1g)	Absent	Absent
10	Aflatoxins (B1,B2,G1,G2)	BDL	<20ppb
11	Phyto-Chemical Analysis		1
	Total Withanolides (%w/w) by HPLC	5.19	≥5%
	(withaferin A) by HPLC	BDL	(<0.1%)

BDL-Below detection levels

Sample complies with the above said specifications.

Storage: Store in original, sealed containers at room temperature. Avoid light and keep away from excessive humidity.



THE WAR PHY WOUND

QC Manager

QC Analyst

Regd. Off.: 5-9-225, 3rd Floor, Sanali Estate, C A Lane, Abids, Hyderabad-500001, Telangana State, India. Corporate Off.: H.No 1057F2, 3rd Floor, PBN Centre, Jubilee Hills, Road No:45, Hyderabad - 500033, Telangana, India.

Ph: +91 40 23554386



Sample Information

CTLA ID: 73042

Date Received: 4/28/2023

Sample Name: 11087 KSM-66 Ashwagandha

Lot Number: KSM/VG/23/5037

Customer: Origin Nutraceutical

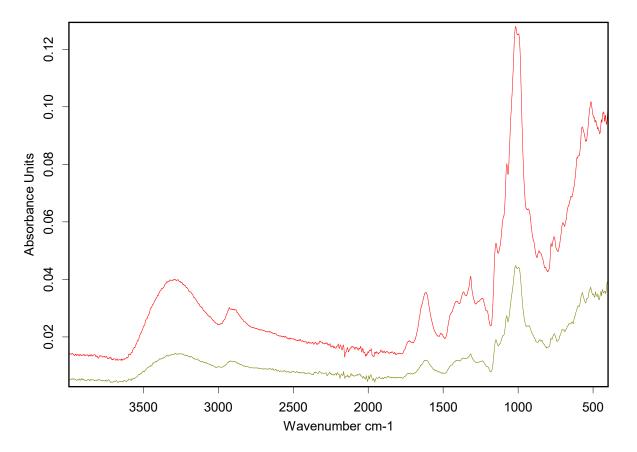
Analysis	Method	MDL Specification	Result	Units
ID, Complete Micro Combo				
ID	FTIR	Report	94.3	%
Total Aerobic Microbial Count	USP <2021>	100 Report	900	cfu/g
Total Coliforms	USP <2021>	10 Report	<10	cfu/g
E. coli	USP <2022>	Report	Absent	
Salmonella	USP <2022>	Report	Absent	
Staphylococcus aureus <2022>	USP <2022>	Report	Absent	
Yeast & Mold	USP <2021>	100 Report	<100	cfu/g
FTIR Spectra	FTIR	Report	Attached	
Ashwagandha	HPTLC	Report	Characteristic	

HPTLC: Fingerprint of test solution is consistent with Withania Somnifera (Ashwaganda root).

5/10/2023

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)

Search Library 5/3/2023 9:51:42 AM



Product Number	39834-ON 11087 DRM KSM-66 Ashwagandha
Entry No.	368
Library name	COMPARISON TESTS.S01
Library description	Origin Nutraceutical
Copyright	Origin Nutraceutical Materials Only

Color	Hit Quality	Compound name	CAS Number	Molecular formula	Molecular weight
	943	39834-ON 11087 DRM KSM-66 Ashwagandha (38810)			

Color	File	Path	Spectrum Type
	73042-ON 11087 KSM-66 Ashwagandha (KSM-VG-23-503	7 C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum



Product: KSM-66 Ashwagandha Extract Certificate No.: P7142166 Version 1

Lot No: KSM/VG/23/S037 Issued for: Origin Nutraceutical

Test / Date / Method	Specification	Results
Total Withanolides 2023-04-10 HPLC	≥ 5%	5.6%

Issuance Date: 2023-04-24

Approved By: Mo Lababidi Senior Analyst

M

Requisition: 1165R14588 Received On 2023-04-03 Code No: N/A For Origin Nutraceutical, 151 E 3450 N Ste 201, Spanish Fork, 84660, Tel:. Fax:..



SICHUAN TONGSHENG AMINO ACID CO., LTD

Manufacture address:No.23 of Yongdinghe West Road, Tianyuan development zone, Deyang, Sichuan, China

Certification of Analysis

Product Name: L-Theanine Molecular Formula: C₇H₁₄N₂O₃ Manufacture time: 2022-06-06 Testing time: 2022-06-14

Bulk Density: 0.32g/mL

Specification:(Enterprise Standard)

CAS#: 3081-61-6

Molecular Weight:174.2 Batch No.: 010095-H2022061401

Expiry time: 2025-06-05 Tapped Density: 0.49g/mL

Titem	Standard Analysis data		Test Method
Appearance	White crystalline powder	Conforms	Visual
Specific rotation(a)D20 (C=2, H2O)	+7.5° to +8.5°	+7.91°	AJ197
Solubility	Freely soluble in water, insoluble in EtOH	Conforms	Visual
Assay	98.0-102.0%	99.41%	AJI97
Melting Point (℃)	200.0℃ to 210 .0℃	201.4-201.7℃	AJI97
Chloride(C1)	Not more than 0.02%	<0.02%	AJI97
Iron(Fe)	Not more than 10ppm	<10ppm	AJI97
Heavy metals(Pb)	Not more than 10ppm	<10ppm	Colorimetry
Arsenic(As2O3)	Not more than 4ppm	<4ppm	AFS
Loss on drying	Not more than 1.0%	0.16%	AJI97
Residue on ignition	Not more than 0.2%	0.07%	AJI97
Total Plate Count	Not more than 1000cfu/g	Conforms	AOAC
Yeat& mould	Not more than 100cfu/g	Conforms	AOAC
Ethyl alcohol	Not more than 3000ppm	Conforms	AOAC
E.Coli	Negative	Negative	AOAC
Salmonella	Negative	Negative	AOAC
PH	5.0-6.0	5.77	AJI97
Mesh Size	90% through 60mesh	90% through 60mesh	Mesh screen
Lead	Not more than 1ppm	Conforms	AAS
Cadmium	Not more than 0.3ppm	Conforms	AAS
Mercury	Not more than 0.1 ppm	Conforms	AAS
Conclusion	The product conforms to t	the Standard	基根本

Inspector:王明芳

Manager:肖宏林

Sheet Made on June 21, 2022





Sample Information

CTLA ID: 69966

Date Received: 3/14/2023

Sample Name: 10310 L-Theanine

Lot Number: 46997

Customer: Origin Nutraceutical

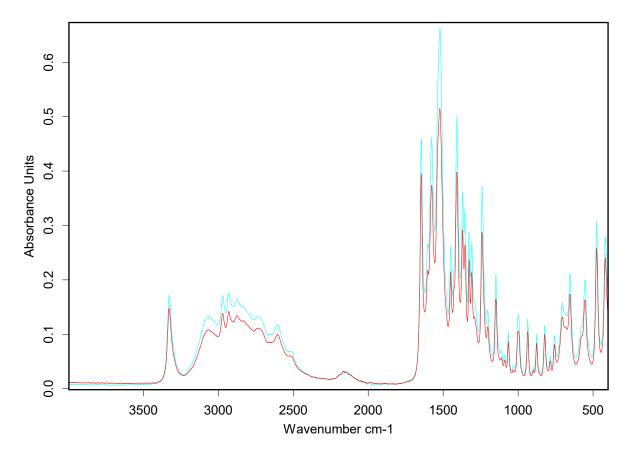
Analysis	Method	MDL Specification	Result	Units
ID	FTIR	Report	97.5	%
FTIR Spectra	FTIR	Report		
L-Theanine	HPLC	0.0750 Report	99.313	%

3/16/2023

Specifications provided by the Customer. Results with an asterisk (A) The Customer 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 3/15/2023 10:12:45 AM



Product Number	10310 L-Theanine 62382 Standard 3
Entry No.	292
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
	975	10310 L-Theanine 62382 Standard 3			

Color	File	Path	Spectrum Type
	69966-ON 10310 L-Theanine (46997).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS	Query Spectrum

6439 Alondra Blvd, Paramount, CA 90723, USA Tel (562) 633 8985, Fax (562) 633 8986

Website: nutrilandgroup.com

Email: nutrilandgroup@outlook.com



Certificate of Analysis

Product:

Magnesium Glycinate 25%

Batch:

12212809MGOG

Manufacturing Date:

December 2021

December 2024

Content

Specification

Results Conforms

Retesting Date:

Methods

Appearance: Particle Size:

Off-white powder 40-60 mesh

Conforms

Sieve

Magnesium (Mg):

≥25% <8% 26% 0.9% ICP 105°C

Loss on Drying: PH:

8 - 11

10.5

USP

Heavy Metals:

<10 ppm

Conforms

ICP-MS

Arsenic (As):

<1 ppm

Conforms

ICP-MS

Cadmium (Cd):

<1 ppm

Conforms

ICP-MS

Lead (Pb):

<1 ppm <1 ppm

Conforms Conforms ICP-MS

Mercury (Hg): Total Plate Count:

<1,000 cfu/g

Conforms

ICP-MS

Mold & Yeast:

<100 cfu/g

Conforms

USP USP

E. Coli:

Negative

Conforms

USP

Salmonella:

Negative Negative Conforms
Conforms

USP USP

GMO Statement:

Staphylococcus Aureus:

Non-GMO

Standard:

Food grade

Treatment:

None

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Digitally Signed By QC Director:

Micheal Ma



Sample Information

CTLA ID: 69968

Date Received: 3/14/2023

Sample Name: 10580 Magnesium Glycinate 25%

Lot Number: 43342/43419

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID	FTIR	Report	94.6	%
FTIR Spectra	FTIR	Report		

3/16/2023

from which it was taken. These results are provided for the benefit of the Customer. MDL = Method Detection Limit. This document is not

Specifications provided by the Customer. Results with an asterisk (4) The Customer 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

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Sample Information

CTLA ID: 55668

Date Received: 7/12/2022

Sample Name: 10580 DRM Magnesium Glycinate 25%

Lot Number: 43419

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
Mineral Analysis	ICP-MS	0.00000 25–29 5	Magnesium 27.9853	%

Customer note: retest

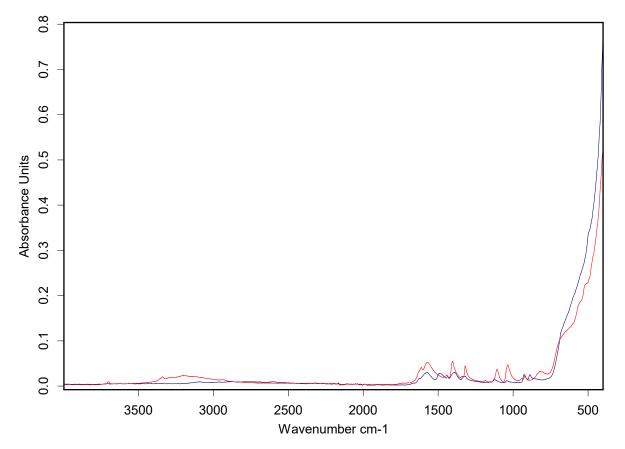
7/20/2022

DATE

Quality Manager

Specifications provided by the Customer. Results with an asterisk (*) denote 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.

Search Library 3/15/2023 10:43:22 AM



Product Number	10580 Magnesium Glycinate 5309 Standard 1
Entry No.	932
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
	946	10580 Magnesium Glycinate 5309 Standard 1			

Color	r File	Path	Spectrum Type
	69968-ON 10580 Magnesium Glycinate 25% (43342-43419) C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\DATA\MEAS Q		Query Spectrum

6439 Alondra Blvd, Paramount, CA 90723, USA Tel (562) 633 8985, Fax (562) 633 8986

Website: nutrilandgroup.com

Email: nutrilandgroup@outlook.com



Certificate of Analysis

Product: Melatonin Chemical Name: N-Acetyl-5-Methoxytryptamine Batch: KH-MT300-220415-1 Manufacturing Date: April 2022 Retesting Date: April 2024 73-31-4 CAS: Nanjing Chrise Co, China Origin: Methods Results Content Specification Visual Off-white to white powder Conforms Appearance: Conforms 80 mesh screen 80 mesh Particle Size: Chemical Formula: $C_{13}H_{16}N_2O_2$ 232.28 Molecular Weight: Conforms IR Identification: **Positive** 99.5% USP40 ≥99% Assay: 0.14% USP40 <1% Loss on Drying: Conforms USP40 Sulphated Ash: <1% Solubility: Insoluble in water Conforms Conforms USP40 Acetic Acid: <0.1% Conforms USP40 Ethyl Acetate: < 0.1% USP40 Conforms Heavy Metals: <10 pm Conforms USP40 Arsenic: <1.0 ppm Conforms USP40 Cadmium: <0.3 ppm <0.5 ppm Conforms USP40 Lead: Conforms USP40 Mercury: <0.2 ppm Conforms USP40 Total Plate Count: <1,000 cfu/g Conforms USP40 Mold & Yeast: <50 cfu/g Conforms USP40 Coliforms: <10 cfu/g Conforms USP40 E. Coli: Negative Conforms USP40 Negative Salmonella: Conforms USP40 Staphylococcus Aureus: Negative Non-GMO **GMO Statement:** Standard: USP food grade Treatment: None None Peak E: Nutritional Information Vitamin A: 0 Calcium: 0 Calories: 0 Vitamin Bs: 0 0 Carbohydrate: 0 Potassium: 0 Sodium: 0 Vitamin C: 0 Fat: Other Vitamins: Fiber: 0 Other Minerals:

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Digitally Signed By QC Director:



Sample Information

CTLA ID: 72256

Date Received: 4/14/2023

Sample Name: 10395 Melatonin

Lot Number: 47444

Customer: Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
Rapid Complete Micro				
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
Melatonin	HPLC	0.075 Report	101.886	%

4/24/2023

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

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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)



Sample Information

CTLA ID:

70052

Date Received:

3/15/2023

Sample Name:

10395 DRM Melatonin

Lot Number:

47444

Customer:

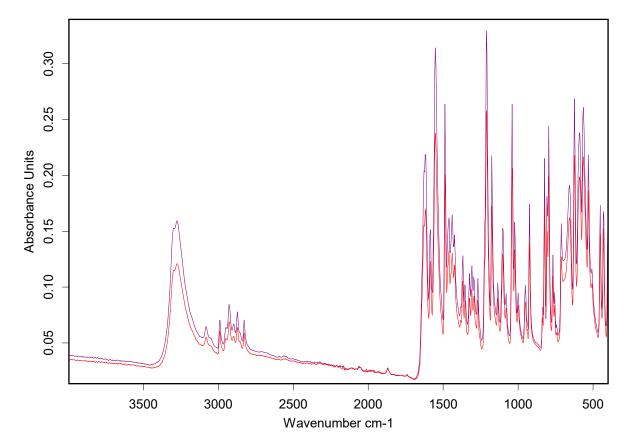
Origin Nutraceutical

Analysis	Method	MDL Specification	Result	Units
ID	FTIR	Report	95.5	%

3/20/2023

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)

Search Library 5/1/2023 11:58:23 AM



Product Number	ON 10395 Melatonin (40985) Standard 2
Entry No.	3113
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
	955	ON 10395 Melatonin (40985) Standard 2			

Color	File	Path	Spectrum Type
	70052-ON 10395 DRM Melatonin (47444).0	C:\Users\CTLA Admin\Documents\Bruker\OPUS_7.5.18\Data\MEAS	Query Spectrum