

# MANUFACTURER ACCREDITATION(S):











### Auria Health, LLC



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BLEND: REVIVE

LOT: 15817

PURPOSE: Certificate of Analysis (COA)

MANUFACTURED: USA



#### Pharma Natural, Inc.

14500 NW 60th Ave, Building 7F, FI 33014 USA Ph.:305 231 8877 Fax::866 526 4796 www.pharmanatural.com

#### CERTIFICATE OF ANALYSIS

REVIVE Product Name: Presentation: 30 Capsules Serving Size: 1 Capsule Servings Per Unit: 30 Lot Number: 15817

MEU-BIO-002

MEU005-2023 CoA Number: Bulk Number: BK3484 Manufacture Date: 11/2023 **Expiration Date:** 11/2026 **Testing Completed** 12/08/2023

Physical Analysis			
Test	Specification	Result	Method
Appearance and Color	Veggie Capsule "00" Clear filled with light brown powder	Pass	Visual, SOP 024
Weight (mg)	760 mg ± 10 %	771 mg	USP<2091>, SOP L047
Disintegration	< 30 min	12 min	USP<2040>, SOP L048

Chemical Analysis				
Active Ingredients per Capsule	Composition	Specification	Assay Result	Method
Almond Mushroom (Agaricus blazei) (fruiting body)	250 mg	r > 0.90 Correlation to Standard	Present by input	Input by batch record review
King trumpet Mushroom ( <i>Pleurotus</i> eryngii) (fruiting body)	160 mg	r > 0.90 Correlation to Standard	Present by input	Input by batch record review
Scarlet Caterpillarclub Mushroom (Cordyceps militaris) (fruiting body)	100 mg	r > 0.90 Correlation to Standard	Present by input	Input by batch record review
Ashwagandha (Withania somnifera) Extract (whole plant)	120 mg	r ≥ 0.90 Correlation to Standard	Present by input	Input by batch record review

		Heavy Metals	
Test	Specification (ppm) (mcg/g)	Results (ppm) (mcg/g)	Method
Lead (Pb)	≤ 0.5	0.008	ICP-MS USP<2232>, SOP L052
Arsenic (As)	≤ 1.5	0.066	ICP-MS USP<2232>, SOP L052
Cadmium (Cd)	≤ 0.5	0.007	ICP-MS USP<2232>, SOP L052
Mercury (Hg)	≤ 1.5	0.100	ICP-MS USP<2232>, SOP L052

Microbiological Analysis			
Test	Specification	Result	Method
Rapid Aerobic Count Plate	≤10,000 cfu/g	<10,000 cfu/g	USP-NF<2021>, SOP L056
Rapid Yeast & Mold Count Plate	≤1,000 cfu/g	<1,000 cfu/g	USP-NF<2021>, SOP L056
Escherichia coli	Absence / 10g	Negative	USP-NF<2022>, SOP L056
Salmonella spp.	Absence / 10g	Negative	USP-NF<2022>, SOP L056

Prepared by

QUALITY SPECIALIST 12/11/2023 DATE

Approved by

Rosa Lydia Solis QUALITY ASSURANCE MANAGER 12/11/2023 DATE FORM 25.1















### Auria Health, LLC



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BLEND: REVIVE

• INGREDIENT: Almond (Agaricus blazei) Mushroom

**LOT:** L23021501A

• PURPOSE: Ingredient COA

**ORIGIN:** USA

### CERTIFICATE OF ANALYSIS

Product Name: Agaricus blazei

Mushroom mycelial biomass powder cultured on organic oats (Avena sativa)

Certified 100% Organic (USDA-NOP Standards) Product of USA (California) Claim:

50010 Item #:

Lot#: L23021501A

04/23/2023 (milling date) Manufacturing Date:

Best Used By Date: 04/2026

	Laboratory A	nalysis	
Test	Method	Specification	Actual Reported Value
Identification	DNA sequencing of master tissue culture, taxonomic and visual monitoring of morphology and growth metrics during growth cycle. Annual HPTLC testing of finished product.	Complies to species positive ID specifications	Complies to species positive ID specifications
% Moisture	Constant Weight Moisture Meter	< 6% moisture	3.7 % moisture
Particle Size	Screen / Sieve	≥ 95% through 60 Mesh	Complies
Gluten	Agrastrip Gluten G12 Test Kit	< 20 ppm	Complies
TPC	FDA BAM Chapter 3 or equivalent	≤ 10,000 cfu/g	< 10 cfu/g
Yeast & Mold	FDA BAM Chapter 18 mod. or equivalent	≤ 1,000 cfu/g	< 10 cfu/g
Coliforms	CMMEF Chapter 9.933 or equivalent	≤ 100 cfu/g	< 10 cfu/g
Salmonella	AOAC-RI 121501 or equivalent	Not detected / 25 g	Not detected / 25 g
Staphylococcus aureus	PF-AOAC 2003.07, 2003.08, 2003.11 or equivalent	< 10 cfu/g (Not Detected)	< 10 cfu/g (Not Detected)
E. coli	CMMEF Chapter 9.933 or equivalent	< 10 cfu/g (Not Detected)	< 10 cfu/g (Not Detected)
Listeria spp.	AOAC-RI 061702 or equivalent	Not detected / 25 g	Not detected / 25 g
Arsenic (As)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤ 0.25 ppm	0.0104 ppm
Cadmium (Cd)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤ 0.1 ppm	0.0112 ppm
Lead (Pb)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤ 0.1 ppm	< 0.00500 ppm
Mercury (Hg)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤ 0.1 ppm	< 0.00500 ppm

	Sensory Analysis	
Test	Specification	Results
Appearance	Powder	Complies
Color	Brown	Complies
Aroma	Mild / Earthy	Complies
Flavor	Slightly Sweet / Nutty / Earthy	Complies
Texture	Powdery	Complies

Rachel Warner **OA Specialist** 

05-02-23

Agaricus L23021501A















### Auria Health, LLC



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BLEND: REVIVE

• INGREDIENT: King Trumpet (Pleurotus eryngii) Mushroom

**LOT:** L23071508A

• PURPOSE: Ingredient COA

ORIGIN: USA

### CERTIFICATE OF ANALYSIS

King Trumpet (Pleurotus eryngii) **Product Name:** 

Mushroom mycelial biomass and fruit body powder cultured on organic oats (Avena

Certified 100% Organic (USDA-NOP Standards) Product of USA (California) Claim:

Item #:

Lot#: L23071508A

Manufacturing Date: 08/24/2023 (milling date)

Best Used By Date: 08/2026

	Laboratory A	nalysis	
Test	Method	Specification	Actual Reported Value
Identification	DNA sequencing of master tissue culture, taxonomic and visual monitoring of morphology and growth metrics during growth cycle. Annual HPTLC testing on milled powder.	Complies to species positive ID specifications	Complies to species positive ID specifications
% Moisture	Constant Weight Moisture Meter	< 6% moisture	2.9 % moisture
Particle Size	Screen / Sieve	≥95% through 60 Mesh	Complies
Gluten	ELISA with RIDASCREEN® Total Gluten - R7041	<20 ppm	Complies
TPC	FDA BAM Chapter 3 or equivalent	≤10,000 cfu/g	20 cfu/g
Yeast & Mold	FDA BAM Chapter 18 mod. or equivalent	≤1,000 cfu/g	10 cfu/g
Coliforms	CMMEF Chapter 9.933 or equivalent	≤100 cfu/g	< 10 cfu/g
E. coli	CMMEF Chapter 9.933 or equivalent	< 10 cfu/g (Not Detected)	< 10 cfu/g (Not Detected
Salmonella	AOAC-RI 121501 or equivalent	Not detected / 25 g	Not detected / 25 g
Listeria spp.	AOAC-RI 061702 or equivalent	Not detected / 25 g	Not detected / 25 g
Arsenic (As)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤0.25 ppm	0.0103 ppm
Cadmium (Cd)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤ 0.1 ppm	0.00620 ppm
Lead (Pb)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	≤0.1 ppm	< 0.00500 ppm
Mercury (Hg)	AOAC 2011.19 and 993.14 (modified) ICP-MS Instrumentation	Not Detected at LOQ	< 0.00500 ppm (LOQ)

Limit of quantification (LOO)

	Sensory Analysis	
Test	Specification	Results
Appearance / Color	Light Brown Powder	Complies
Aroma	Mild / Earthy	Complies
Flavor	Nutty / Earthy	Complies
Texture	Powdery	Complies

Hector Ramirez Quality Responsible

09-06-23

Date

King Trumpet L23071508A















Last Updated · 04/12/2025

### Auria Health, LLC



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BLEND: REVIVE

INGREDIENT: Scarlet Caterpillarclub (Cordyceps militaris) Mushroom

**LOT**: L23081504P

• PURPOSE: Ingredient COA

ORIGIN: USA

#### CERTIFICATE OF ANALYSIS

Product Name: Cordyceps militaris

Mushroom mycelial biomass, stroma and fruit body powder cultured on organic oats

(Avena sativa)

Certified 100% Organic (USDA-NOP Standards) Product of USA (California) Claim:

50040 Item #: Lot#: L23081504P

Manufacturing Date: 10/08/2023 (milling date)

10/2026 Best Used By Date:

	Laboratory A	nalysis	
Test	Method	Specification	Actual Reported Value
Identification	DNA sequencing of master tissue culture, taxonomic and visual monitoring of morphology and growth metrics during growth cycle. Annual HPTLC testing on milled powder.	Complies to species positive ID specifications	Complies to species positive ID specifications
% Moisture	Constant Weight Moisture Meter	< 6% moisture	3.8 % moisture
Particle Size	Screen / Sieve	≥ 95% through 60 Mesh	Complies
Gluten	ELISA with RIDASCREEN® Total Gluten - R7041	<20 ppm	Complies
TPC	FDA BAM Chapter 3 or equivalent	≤10,000 cfu/g	10 cfu/g
Yeast & Mold	FDA BAM Chapter 18 mod. or equivalent	≤1,000 cfu/g	10 cfu/g
Coliforms	CMMEF Chapter 9.933 or equivalent	≤ 100 cfu/g	<10 cfu/g
E. coli	CMMEF Chapter 9.933 or equivalent	< 10 cfu/g (Not Detected)	< 10 cfu/g (Not Detected)
Salmonella	AOAC-RI 121501 or equivalent	Not detected / 25 g	Not detected / 25 g
Listeria spp.	AOAC-RI 061702 or equivalent	Not detected / 25 g	Not detected / 25 g
Arsenic (As)	ICP-MS	≤0.25 ppm	0.0124 ppm
Cadmium (Cd)	ICP-MS	≤0.1 ppm	0.00681 ppm
Lead (Pb)	ICP-MS	≤0.1 ppm	0.0164 ppm
Mercury (Hg)	ICP-MS	Not Detected at LOQ	< 0.00500 ppm (LOQ)

Limit of quantification (LOO)

	Sensory Analysis	
Test	Specification	Results
Appearance / Color	Light Brown Powder	Complies
Aroma	Mild / Earthy	Complies
Flavor	Nutty / Earthy	Complies
Texture	Powdery	Complies

Hector Ramirez

Quality Responsible

10-18-23

C. Militaris L23081504P







#### Auria Health, LLC

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**BLEND:** REVIVE

**INGREDIENT:** Ashwagandha (Withania somnifera) Whole Plant Extract

**LOT**: C231027

**PURPOSE:** Ingredient COA

ORIGIN: USA

CERTI	FICATE	OF ANA	LYSIS
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**Product Name Product Code** 

ASHWAGANDHA EXTRACT

Batch No.

0130 C231027

TR No. Date of Manufacture Date of Expiry

KL23F0416 June 2023 May 2028

Sold To: PHARMA NATURAL, INC

PO No.: PN-67461

Quantity: 10.000 KG

Category

Intended for Nutraceutical application

Botanical/Scientific name

Withania somnifera

CAS No Plant part 90147-43-6 Whole plant

Preparation type Solvent used for extraction Methanol

Extraction

Solvent used in manufacture

None

Final extract ratio

5:1 to 8:1

Standardization

Alkaloids, Withanolides, Withaferin-A

**Excipient Used** 

Maltodextrin

Excipient name

%Used

CAS No.

Maltodextrin

25% to 50%

9050-36-6

**Parameters** 

Identification

Result

Limit

Reference

PHYSICAL

Description

Complies

Complies

Brown to dark brown powder with characteristic Visual and Organoleptic

To comply by TLC for a)

odour, hygroscopic\*

SLL/STP-A-012

Alkaloids and b) Withanolides

SLL/STP-S-028

Solubility

-Water solubles (1% w/v

85.31 % w/w

Not less than 70.0% w/w

solution in water) -Alcohol solubles (1% w/v

80.49 % w/w

Not less than 60.0% w/w

solution in 50% v/v

alcohol)

Loss on drying

2.58 % w/w

Not more than 5.0% w/w (dried at 105°C)

Ash Content

4.10 % w/w

Not more than 15.0% w/w USP <561>

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East Windsor

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## Auria Health, LLC





• **BLEND**: REVIVE

• INGREDIENT: Ashwagandha (Withania somnifera) Whole Plant Extract

• LOT: C231027

• PURPOSE: Ingredient COA

• ORIGIN: USA

	CERTIFICAT	E OF ANALYSIS	Page 2 of 4
Product Code	ASHWAGANDHA EXTRACT 0130		5
	C231027		
	KL23F0416		
	June 2023		
Date of Expiry	May 2028		
Tapped bulk density	0.84 g/ml	Between 0.60g/ml and 0.85g/ml	USP <616>
Loose bulk density	0.56 g/ml	Record	USP <616>
Sieve Test (Passes Through)			USP <786>
- 20 Mesh	100.00 % w/w	Not less than 100% w/w	
- 40 Mesh	100.00 % w/w	Not less than 85% w/w	
- 80 Mesh	92.21 % w/w	Not less than 70% w/w	
CHEMICAL			
Assay			
-Content of Alkaloids by Gravimetry	1.18 % w/w	Not less than 1.0% w/w	SLL/STP-A-051
-Content of total Withanolides (free Withanolides and Glycowithanolides) by Gravimetry	7.18 % w/w	Not less than 7.0% w/w and not more than 10.0% w/w	SLL/STP-W-005
-Content of Withaferine- by HPLC	A 0.44 % w/w	Not less than 0.25% w/w	SLL/STP-W-004
Elemental Contaminants	(Heavy Metals)		
Lead	<0.2 ppm (µg/g)	Not more than 3ppm (µg/g)	USP <2232>
Arsenic	<0.2 ppm (µg/g)	Not more than 1ppm (µg/g)	USP <2232>
Cadmium	<0.2 ppm (µg/g)	Not more than 1ppm (µg/g)	USP <2232>
Mercury	<0.02 ppm (µg/g)	Not more than 0.1ppm (µg/g)	USP <2232>
OTHERS			
Residual solvents	Complies	To comply as per USP	USP <467>
Residual pesticides	Complies	To comply as per USP	USP <561>
Glyphosate MICROBIAL	Complies	Not more than 0.2ppm	SLL/STP-P-122
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East Windsor			750 Innovation Circle
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### Auria Health, LLC





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• **BLEND**: REVIVE

INGREDIENT: Ashwagandha (Withania somnifera) Whole Plant Extract

LOT: C231027

• PURPOSE: Ingredient COA

• ORIGIN: USA

		TE OF ANALYSIS	Page 3 of 4
Product Name	ASHWAGANDHA EXTRACT		
Product Code	0130		
Batch No.	C231027		
TR No.	KL23F0416		
Date of Manufacture	June 2023		
Date of Expiry	May 2028		
Total aerobic microbial count	800 cfu/g	Not more than 5000cfu/g	USP <2021>
Total yeasts and molds count	<10 cfu/g	Not more than 100cfu/g	USP <2021>
Escherichia coli	Complies	Negative/10g	USP <2022>
Salmonella	Complies	Negative/10g	USP <2022>
Staphylococcus aureus	Complies	Negative/10g	USP <2022>
Pseudomonas aeruginos	a Complies	Negative/10g	USP <62>
Bile tolerant gram negative bacteria	<10 cfu/g	Not more than 100cfu/g	USP <2021>
Coliforms	<10 cfu/g	Less than 10cfu/g	BAM Chapter 4
ADDITIONAL INFORMATI	ON		
Sanitising treatment	Non-irradiated and not t	reated with ETO	
Certification status (Kosher/Halal)	Kosher and Halal certifie	d	
BSE/TSE status	BSE/TSE free		
Genetic modification status	GMO free		
Country of origin	India		
Cultivated or wild crafted	Cultivated		
Storage condition	Store at room temperatu	ire	
Manufactured By	Sami-Sabinsa Group Limi 19/1, 19/2, I Main II Phas India	ted - Peenya se, Peenya Industrial Area, Bar	ngalore. 560 058, Karnataka,
Manufactured At	Sami-Sabinsa Group Limi Plot No. 40-41 & 30-35, I	ted - Kunigal KIADB Industrial Area,Tumkur	District, Kunigal - 572130
Remarks .	to batch of the product e raw material	uct, there is likely to be minor cause of the geographical and hygroscopic and tend to aggluctbefore use.	I seasonal variations of the

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### Auria Health, LLC





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BLEND: REVIVE

INGREDIENT: Ashwagandha (Withania somnifera) Whole Plant Extract

**LOT**: C231027

• PURPOSE: Ingredient COA

**ORIGIN: USA** 

## CERTIFICATE OF ANALYSIS

Page 4 of 4

Product Name Product Code

ASHWAGANDHA EXTRACT 0130

Batch No. TR No. Date of Manufacture

Date of Expiry

C231027 KL23F0416 June 2023

May 2028

Dr. Hari Ramachandran QA/ QC Manager

The above certificate of analysis is based on Specifications Issue No: 14 Dated: April 20, 2022

\*\*In-House developed and validated methods NAP:Not Applicable NAV:Not Available

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· None.

### Auria Health, LLC





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BLEND: ELEVATE

**INGREDIENT:** Silica (Silicon dioxide)

**LOT**: 18997

• PURPOSE: Ingredient COA

ORIGIN: USA



HEALTHY PRODUCTS



Product Name: Silicon Dioxide Powder (Pirosil PS-200)

Manufacture Date: May, 2023 Expiration Date: May, 2025

Batch No: 18997 Quantity: 13.44mt

Characteristics	Test Results	FCC Limits
A	PASSES TEST	PASSES TEST
В	PASSES TEST	PASSES TEST
ASSAY (as SiO 2), %	96	94 min
Lead, ppm	1.0	5 Max.
Loss on Drying (105° C 2 hours), %	5.7	7 Max.
Loss on Ignition (1000° C 1 hour), %	6.0	8.5 Max.
Soluble Ionizable Salts (as Na 2 SO4), %	1.5	5 Max.
Tapped density, g/l	158	160 Max.

This product meets FCC specifications.

Note: The above information is based on the certificate of analysis received from the manufacturer of this product. It is not intended to be a substitute for strict quality control analysis by the purchaser of this product.



Quality Control

CONTACT INFO: 12601 NW 115 th ave, Ste A-103 Medley, FI 33178

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