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- **MANUFACTURER ACCREDITATION(S):**

- **BLEND:** REVIVE
- **LOT:** 15515
- **PURPOSE:** Certificate of Analysis (COA)
- **MANUFACTURED:** USA

cGMP Certified Good Manufacturing Practices

FDA REGISTERED

NSF

ISO 9001 2015

usp

Pharma Natural® Natural Health Solutions

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

Product Name:	REVIVE	CoA Number:	MEU003-2023
Presentation:	30 Capsules	Bulk Number:	BK3484
Serving Size:	1 Capsule	Manufacture Date:	11/2023
Servings Per Unit:	60	Expiration Date:	11/2026
Lot Number:	15515	Testing Completed	12/08/2023
P.O:	MEU-BIO-002		

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 (*Pleurotus 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	0.009	ICP-MS USP<2232>, SOP L052
Arsenic (As)	≤ 1.5	0.064	ICP-MS USP<2232>, SOP L052
Cadmium (Cd)	≤ 0.5	0.013	ICP-MS USP<2232>, SOP L052
Mercury (Hg)	≤ 1.5	0.077	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 Wilmer Torres
QUALITY SPECIALIST
12/11/2023
DATE

Approved by Rosa Lydia Solis
QUALITY ASSURANCE MANAGER
12/11/2023
DATE

FORM 25.1

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- **INGREDIENT ACCREDITATION(S):**

- **BLEND:** REVIVE
- **INGREDIENT:** Almond (*Agaricus blazei*) Mushroom
- **LOT:** L23021501A
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

USDA ORGANIC

KOSHER

GF GLUTEN FREE

CERTIFIED VEGAN VEGAN.ORG

BRCGS Food Safety CERTIFICATED

NON GMO Project VERIFIED nongmoproject.org

CERTIFICATE OF ANALYSIS

Product Name: Agaricus blazei
Mushroom mycelial biomass powder cultured on organic oats (*Avena sativa*)
Claim: Certified 100% Organic (USDA-NOP Standards) Product of USA (California)
Item #: 50010
Lot #: L23021501A
Manufacturing Date: 04/23/2023 (milling date)
Best Used By Date: 04/2026

Laboratory Analysis

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
QA Specialist

05-02-23
Date

12h 1816
Rq 15073

Agaricus L23021501A

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- **INGREDIENT ACCREDITATION(S):**

- **BLEND:** REVIVE
- **INGREDIENT:** King Trumpet (*Pleurotus eryngii*) Mushroom
- **LOT:** L23071508A
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

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NON GMO Project VERIFIED nongmoproject.org

CERTIFICATE OF ANALYSIS

Product Name: King Trumpet (*Pleurotus eryngii*) Mushroom mycelial biomass and fruit body powder cultured on organic oats (*Avena sativa*)

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

Item #: 50080

Lot #: L23071508A

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

Best Used By Date: 08/2026

Laboratory Analysis

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 (LOQ)

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*

12/1817
By 15072

King Trumpet L23071508A

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- **INGREDIENT ACCREDITATION(S):**

- **BLEND:** REVIVE
- **INGREDIENT:** Scarlet Caterpillarclub (*Cordyceps militaris*) Mushroom
- **LOT:** L23081504P
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

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NON GMO Project VERIFIED nongmoproject.org

CERTIFICATE OF ANALYSIS

Product Name: *Cordyceps militaris* Mushroom mycelial biomass, stroma and fruit body powder cultured on organic oats (*Avena sativa*)

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

Item #: 50040

Lot #: L23081504P

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

Best Used By Date: 10/2026

Laboratory Analysis

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 (LOQ)

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
Date

Rg 1812
Rg 15670

C. Militaris L23081504P

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- **INGREDIENT ACCREDITATION(S):**

- **BLEND:** REVIVE
- **INGREDIENT:** Ashwagandha (*Withania somnifera*) Whole Plant Extract
- **LOT:** C231027
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

CERTIFICATE OF ANALYSIS

Product Name	ASHWAGANDHA EXTRACT
Product Code	0130
Batch No.	C231027
TR No.	KL23F0416
Date of Manufacture	June 2023
Date of Expiry	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	90147-43-6
Plant part	Whole plant
Preparation type	Extraction
Solvent used for extraction	Methanol
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	Result	Limit	Reference
PHYSICAL			
Description	Complies	Brown to dark brown powder with characteristic odour, hygroscopic*	Visual and Organoleptic
Identification	Complies	To comply by TLC for a) Alkaloids and b) Withanolides	SLL/STP-A-012
Solubility			SLL/STP-S-028
-Water solubles (1% w/v solution in water)	85.31 % w/w	Not less than 70.0% w/w	
-Alcohol solubles (1% w/v solution in 50% v/v alcohol)	80.49 % w/w	Not less than 60.0% w/w	
Loss on drying	2.58 % w/w	Not more than 5.0% w/w (dried at 105°C)	USP <731> RM 0915
Ash Content	4.10 % w/w	Not more than 15.0% w/w	USP <561> RM 15014

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Fax: 801-465-8600

- **INGREDIENT ACCREDITATION(S):**

- **BLEND:** REVIVE
- **INGREDIENT:** Ashwagandha (*Withania somnifera*) Whole Plant Extract
- **LOT:** C231027
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

CERTIFICATE OF ANALYSIS

Product Name	ASHWAGANDHA EXTRACT
Product Code	0130
Batch No.	C231027
TR No.	KL23F0416
Date of Manufacture	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-A by HPLC	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	Complies	Not more than 0.2ppm	SLL/STP-P-122
MICROBIAL			

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- **INGREDIENT:** Ashwagandha (*Withania somnifera*) Whole Plant Extract
- **LOT:** C231027
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

CERTIFICATE OF ANALYSIS

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 aeruginosa	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 INFORMATION

Sanitising treatment	Non-irradiated and not treated with ETO
Certification status (Kosher/Halal)	Kosher and Halal certified
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 temperature
Manufactured By	Sami-Sabinsa Group Limited - Peenya 19/1, 19/2, I Main II Phase, Peenya Industrial Area, Bangalore. 560 058, Karnataka, India
Manufactured At	Sami-Sabinsa Group Limited - Kunigal Plot No. 40-41 & 30-35, KIADB Industrial Area,Tumkur District, Kunigal - 572130
Remarks	*Since it is a herbal product, there is likely to be minor color variation from batch to batch of the product ecause of the geographical and seasonal variations of the raw material *Natural products can be hygroscopic and tend to agglomerate sometimes. It is suggested to sift the productbefore use.

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- **LOT:** C231027
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

- **INGREDIENT ACCREDITATION(S):**

CERTIFICATE OF ANALYSIS

Product Name	ASHWAGANDHA EXTRACT
Product Code	0130
Batch No.	C231027
TR No.	KL23F0416
Date of Manufacture	June 2023
Date of Expiry	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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- **INGREDIENT ACCREDITATION(S):**
  - None.

- **BLEND:** ELEVATE
- **INGREDIENT:** Silica (*Silicon dioxide*)
- **LOT:** 18997
- **PURPOSE:** Ingredient COA
- **ORIGIN:** USA

KJC NUTRA HEALTHY PRODUCTS QUALITY

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
B	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_2SO4), %	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, Fl 33178
kcharfan@kjcnutra.com / cs@kjcnutra.com 786.462.2549 / 786.444.5905

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PM 0481
Ey 14989