



Micro Quality Labs, Inc.

Specializing in Pharmaceutical, Dietary Supplements, Toys and Cosmetics Testing
3125 N. Damon Way • Burbank, California 91505
(818) 845-0070 • Fax: (818) 845-0030
E-Mail: Karine@MicroQualityLabs.com



Customer: Blue Diamond Herbs
Address: 8029 Fairview Rd. STE E
Mint Hill, NC 28227

ANALYTICAL/CHEMICAL CERTIFICATE OF ANALYSIS

Sample Name: THE GODFATHER MAENG DA
Product Code: N/A
Batch/Lot: BDH107GF
MQL Accession: 210701-0218

PO#: N/A
Sample Description: RAW
Rush: N/A
Received Date: 07/01/21

| Test Requested: | Test Method: | Specification: | Results: |
|-----------------|-------------------------|----------------|-----------|
| Arsenic | MQLTM-0278 By ICP-MS | N/A | 0.055 ppm |
| Cadmium | MQLTM-0278 By ICP-MS | N/A | 0.023 ppm |
| Mercury | MQLTM-0278 By ICP-MS | N/A | 0.012 ppm |
| Lead | MQLTM-0278 By ICP-MS | N/A | 0.303 ppm |

Prepared By:


Stella Garibian/Document Control Specialist

JUL 09 2021

07/09/21

Reviewed By:


Krista Otanez/Quality Assurance Coordinator

JUL 09 2021

07/09/21

Micro Quality Laboratories, Inc. (MQL), is an A2LA ISO 17025 accredited testing laboratory (Certificate Number 3034.01). The requirements of ISO 17025 were followed for the test, results and preparation of this certificate of analysis. MQL's scope of accreditation may be found on A2LA or MQL websites.

The aforementioned results on this report are representative of the samples submitted and may not be indicative of the entire manufacture, batch, and/or lot. Applicable current GMP's shall always be used when sampling. GLP's shall always be practiced by Micro Quality Labs to ensure the most accurate results.

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Customer: Blue Diamond Herbs
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| | |
|----------------|---------------|
| Received From: | Mint Hill, NC |
| Received Date: | 06/30/21 |
| Release Date: | 07/13/21 |
| PO # | N/A |

MICROBIOLOGICAL CERTIFICATE OF ANALYSIS

Sample Name: THE GODFATHER MAENG DA 50G POWDER

Product Code: N/A

Batch/Lot #: BDH107GF

MQL Accession #: 1127068

Description: RAW

| Analyte: | Result: | Method: | Test Date: | Comment: |
|---------------------|----------------------------|----------------------------|------------|----------|
| TPC | 2.1x10 ⁴ cfu/gm | TM-01 (Modified USP61) | 06/30/21 | N/A |
| Yeast/Mold | 3.0x10 ³ cfu/gm | TM-01 (Modified USP61) | 06/30/21 | N/A |
| Coliforms | 10 ³ cfu/gm | TM-01A (Modified USP62) | 06/30/21 | N/A |
| E.coli | Absent | TM-01A (Modified USP62) | 06/30/21 | N/A |
| Pseudomonas spp. | Absent | TM-01A (Modified USP62) | 06/30/21 | N/A |
| S.aureus | Absent | TM-01A (Modified USP62) | 06/30/21 | N/A |
| Salmonella/Shigella | Absent | TM-01A (Modified USP62) | 06/30/21 | N/A |

All Products were tested in accordance with the USP Standard for Total Plate Count and Enrichment. Additional guidance was referenced by CTFA Microbiological Guidelines.

Prepared By: Erika Zayas/ Document Control Specialist

Reviewed By: Ani Zohrabyan/Microbiologist

JUL 13 2021

Date:

COMMENT REV.01: Mold comment removed upon client's request.

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United States Pharmacopoeia (USP) Limit for Nutritional Supplements:

- Arsenic 3.000 ppm / 3000.000 ppb
- Cadmium 3.000 ppm / 3000.000 ppb
- Lead 10.000 ppm / 10000.000 ppb
- Mercury 3.000 ppm / 3000.000 ppb

California Proposition 65 Daily Limits for Heavy Metals:

- Arsenic 10 ppm / 10,000.000 ppb
- Cadmium 4.1 ppm / 4,100.000 ppb
- Lead 0.5 ppm / 500.000 ppb**
- Methyl Mercury 0.3 ppm / 300.000 ppb

For labels with a single daily dose of 500mg, the PPM in products cannot be higher than:

- Arsenic = 20.000 ppm / 20,000.000 ppb
- Cadmium = 8.000 ppm / 8,000.000 ppb
- Lead = 0.900 ppm / 900.000 ppb
- Mercury = 0.600 ppm / 600.000 ppb

FDA Tolerable Daily Diet Lead intake:

- Children <6 years old = 6.000 ppm / 6000.000 ppb
 - Pregnant women = 25.000 ppm / 25000.000 ppb
 - Adults = 75.000 ppm / 75000.000 ppb
-



Recommended Microbial Limits for Botanical Ingredients (in colony-forming units (cfu)/g)
© AHPA 2014

[Current as of July 2012]

| Organization | AHPA | NSF/ANSI | USP | WHO | EHIA | EP | AHPA | USP |
|-------------------------------------------------------------|------------------------------------------------------------------------|-----------------------------------|------------------------------|-----------------------------------------------------------------------|------|----|-----------------------------------------------|-----------------------------|
| Plant material | Dried, unprocessed herbs for use as ingredients in dietary supplements | Botanical ingredient, non-extract | Dried or powdered botanicals | Untreated crude intended for further processing | NA | NA | Powdered botanical extracts and soft extracts | Powdered botanical extracts |
| Total aerobic microbial count | 10^7 | 10^7 | 10^6 | NA or 10^6 - 10^7 as per specific monographs | NA | NA | 10^4 | 10^4 |
| Total combined yeast & mold count | 10^6 | 10^6 | 10^3 | 10^5 (mold propagules); Occasionally 10^4 for specific monographs | NA | NA | 10^3 | 10^3 |
| Enterobacteria count (bile-tolerant Gram-negative bacteria) | 10^4 (Total coliforms) | 10^4 | 10^3 | 10^3 | NA | NA | 10^2 (Total coliforms) | NA |
| <i>Escherichia coli</i> | Not detected in 10 g* | 10^{2**} | Absence in 10 g | 10^4 | NA | NA | Not detected in 10 g* | Absence in 10 g |
| <i>Salmonella</i> spp. | Not detected in 25 g* | Not detected in 10 g | Absence in 10 g | NA or absent | NA | NA | Not detected in 25 g* | Absence in 10 g |
| <i>Staphylococcus aureus</i> | NA | Not detected in 10 g | NA | NA or absent | NA | NA | NA | NA |

AHPA – American Herbal Products Association, Guidance, 8630 Fenton St. #918, Silver Spring, MD 20910; 301-588-1171.

EHIA – European Herbal Infusions Association

EP – European Pharmacopoeia

NSF/ANSI – NSF International Standard/American National Standard for Dietary Supplements 173 – 2006

USP – United States Pharmacopeial Convention, USP-NF 35-30, 2012

WHO – World Health Organization, *Quality control methods for medicinal plant materials*, Geneva, 1998

NA – Not Assigned

*Sample size may vary depending on the method used.

**If the presence of *Escherichia coli* is confirmed, then testing shall be performed based on the USFDA *Bacteriological Analytical Manual* in Chapter 4A to determine whether the colonies are pathogenic enterovirulent *Escherichia coli* (EEC), not limited to O157:H7. There is a zero tolerance for the presence of EEC.

(a) (i) for dried, unprocessed herbs for use as ingredients in dietary supplements, and (ii) for herbal supplements in solid form consisting of dried, unprocessed herbs:

- Total aerobic plate count: 10^7 colony forming units/gram
- Total yeasts and molds: 10^5 colony forming units/gram
- Total coliforms: 10^4 colony forming units/gram
- Salmonella spp.: not detected in 25 grams
- Escherichia coli: not detected in 10 grams
- Total aflatoxins (B1 + B2 + G1 + G2): 20 $\mu\text{g/kg}$ (ppb)
- Aflatoxin B1: 5 $\mu\text{g/kg}$ (ppb)