VITALABS®, INC. The Natural Choice

NSF GMP CERTIFIED NSF/ANSI 455-2 Dietary Supplements

SQF GFSI-Benchmarked Standard SQF Code Edition 9: FSC 31 Dietary Supplements Manufacturing

CERTIFICATE OF ANALYSIS

Manufactured For: Brand on Demand Inc

Label Name: UNLABELED

Customer Master Batch Lot Number: 35658

Vitalabs Lot Number: 232644

Product Name: Super Fat Burner Capsules

Product Code: VLI659

Date Manufactured: 12/2023

Expiration Date: 12/2025

Product Appearance: #00 green/green oblong bovine capsules filled with a reddish-orange to reddish-brown powder, which may be spotted ; Result: Passed

Actual Weight:	Target Weight:	Weight Range:	Method:
879.24 mg	870.00 mg	783.00 - 957.00 mg	Current USP
Disintegration Time:	**Specification:**	**Result:**	**Method:**
	NMT 30 Minutes	10 mins	Current USP

Serving Size: 4 capsules

Reference: ATDS# 9041-00/23, CKv4p72, 74, 9041-01/23

Vitalabs GMP Info: NSF/ANSI 455-2 Certification Number C019998-HSCDS-3

Vitalabs GFSI Info: SQF Certification Number 72660

COMMENTS

DIETARY INGREDIENTS

Active Ingredient Name	Label Claim (per 4 capsules)	Result	% of Label Claim	Spec	Test Method
Vitamin C (as ascorbic acid)	60.00 mg	60.00 mg	100.00	NLT 100%	†HPLC
Vitamin B-6 (as pyridoxine HCl)	25.00 mg	25.00 mg	100.00	NLT 100%	†HPLC
Choline (as choline bitartrate)	200.00 mg	200.00 mg	100.00	NLT 100%	**ID-HPLC
Chromium (as chromium polynicotinate)	200.00 mcg	200.00 mcg	100.00	NLT 100%	**ID-ICP
Medium Chain Triglycerides Oil	70.00 mg	70.00 mg	100.00	NLT 100%	**ID by GC
CLA (Conjugated Linoleic Acid)	100.00 mg	100.00 mg	100.00	NLT 100%	**ID by GC
GLA (Gamma-Linolenic Acid)	5.00 mg	5.00 mg	100.00	NLT 100%	**ID-GC
Bladderwrack Thallus Powder	50.00 mg	50.00 mg	100.00	NLT 100%	**ID by FTIR
Inositol	500.00 mg	500.00 mg	100.00	NLT 100%	†HPLC
Gymnema sylvestre Leaf (25% Extract)	25.00 mg	25.00 mg	100.00	NLT 100%	**ID-HPLC
Garcinia cambogia Fruit Extract (50% hydroxycitric acid)	200.00 mg	200.00 mg	100.00	NLT 100%	**ID-HPLC
L-Carnitine (as L-Carnitine Tartrate)	25.00 mg	33.68 mg	134.72	100-150%	HPLC
Turmeric Root (95% extract)	25.00 mg	32.60 mg	130.40	100-150%	HPLC
Coenzyme Q10	5.00 mg	5.66 mg	113.20	100-150%	HPLC
Proprietary Blend Consisting of: Spirulina Powder, L-Phenylalanine, L-Tyrosine, L-Methionine, Bromelain, Psyllium Husk Powder, Clove Bud Powder, Allspice, Kelp 10:1 Extract, Juniper Berry 4:1 Extract, Buchu Leaf 4:1 Extract, Uva Ursi Leaf (20% Extract), Cinnamon Bark 10:1 Extract (*Cinnamomum cassia*), Cranberry Fruit Concentrate, and Grapefruit Fruit 4:1 Extract	631.00 mg	641.05 mg	101.59	100-150%	**

OTHER INGREDIENTS

Gelatin (bovine)
Rice flour
Vegetable magnesium stearate
Silicon dioxide
Chlorophyll

REQUIRED ALLERGEN WARNINGS

Eggs: No	**Fish:** No	**Milk:** No	**Peanuts:** No
Sesame: No	**Shellfish:** No	**Soy:** No	**Tree Nuts:** No

Wheat: No

HEAVY METALS

Type of Heavy Metal	Limits	Results	Test Method
Lead	<= 2.75 mcg per serving	1.17 mcg / 4 Capsules	*(ICP-MS)
Arsenic	<= 10 mcg per serving	3.67 mcg / 4 Capsules	*(ICP-MS)
Cadmium	<= 4.1 mcg per serving	0.076 mcg / 4 Capsules	*(ICP-MS)

MICROBIOLOGY

Microbial Test	Limits	Results	Test Method
Standard Plate Count	< 10,000 cfu / g	720 cfu/g	USP<2021>
Yeast and Mold	< 1,000 cfu / g	10 cfu/g	USP<2021>
E. coli	Negative / 10g	Negative	USP<2022>
Salmonella	Negative / 10g	Negative	USP<2022>
Staphylococcus aureus	Negative / 10g	Negative	USP<2022>

SIGNATURE

APPROVED BY: Kirk A Brackett **DATE:** 02-05-24

*** - In Accordance with 21 CFR 111.75(d)(1) the following finished dietary ingredients product specifications are exempt from direct finished batch testing requirements as set forth in paragraph 21 CFR 111.75(c)(1). Possible exemption reasons include: instrumental quantification limits, unavailability of a scientifically valid test method, matrix interference, and the lack of biomarkers. These ingredients are confirmed by proper raw material identification, verified by production process controls, and QA batch record review and approval to ensure finished product meets all approved MMR in-process specifications.*

† - This Lot is a part of a rotational statistical testing program in accordance with 21CFR 111.75(c) for a subset of finished dietary supplement batches manufactured. Marked ingredients were not included in this rotation. These ingredients are confirmed by proper raw material identification, verified by production process controls and QA batch record review and approval to ensure finished product meets all approved MMR in-process specifications. Ref: SOP No. QC-3.

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