CERTIFICATE OF ANALYSIS

Product Name:	**CoQ10 Daily with Resveratrol Capsules**
Product #:	**PY034** Manufactured for: Purity Products
Lot#:	211938
Date Manufactured:	10/2021
Product Appearance:	#0 green/green chlorophyll vegetarian capsule filled with a reddish orange paste. Result: Passed
Weight Variation:	Theoretical Weight: 590.00 mg Specification: 531-649 mg Results: 595.00 mg Method: Current USP
Disintegration Time:	Specification: NMT 30 mins. Result: 11 mins. Method: Current USP
Reference:	B1418p147, B1425p55, B1381p164, B1429p59,60

DIETARY INGREDIENTS

Ingredient Name	LC/2caps	Result	% of LC	Spec	Method
Vitamin A (as natural beta carotene)	600.00 mcg (2000.00 IU)	600.00 mcg	100.00	100-150%	**
Vitamin D (as Cholecalciferol)	25.00 mcg (1000.00 IU)	25.00 mcg	100.00	100-150%	**
Vitamin E (as d-alpha tocopheryl acetate)	40.20 mg (60.00 IU)	48.61 mg	120.92	100-150%	HPLC
Vitamin B-12 (as methylcobalamin)	300.00 mcg	Positive	100.00	Positive **	ID-HPLC
Coenzyme Q10 (as ubidicarenone)	100.00 mg	107.24 mg	107.24	100-150%	HPLC
Trans-Resveratrol (P cuspidatum, 99%)	30.00 mg	30.01 mg	100.03	100-150%	HPLC

OTHER INGREDIENTS

Rice bran oil,
Vegetable cellulose,
Vegetable wax,
Rosemary extract,
Sodium copper chlorophyllin

HEAVY METALS

Heavy Metal	Specification	Result/2 caps	Method
Lead:	2.75 µg/maximum daily dose	0.037 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	ND	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.004 mcg	ICP-MS

CERTIFICATE OF ANALYSIS

MICROBIOLOGY			
Micro Study#MB0021560	**Specification**	**Result**	**Method**
Total Plate Count:	<10,000 CFU/g	<10 cfu/g	Current USP
Yeast & Mold Count:	<1,000 CFU/g	5 cfu/g	Current USP
Escherichia Coli:	Negative	ND	Current USP
Salmonella:	Negative	ND	Current USP
Staphylococcus Aureus:	Negative	ND	Current USP
Prepared by: Rebecca Rivera			Date: 11/19/2021
Reviewed by: Rajvidya Thakker			Date: 11/19/2021
Approved by: Santosh Dawri			Date: 11/19/2021

**- 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). Also 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.