

CERTIFICATE OF ANALYSIS

Product Name:	CoQ10 Daily with Resveratrol Capsules		
Product #:	PY034	Manufactured for: Purity Products	
Lot#:	193305		
Date Manufactured:	8/2020		
Product Appearance:	#0 green/green chlorophyll vegetarian capsule filled with a reddish orange paste. Result: Passed		
Weight Variation:	Theoretical Weight: 590.00 mg Results: 602.30 mg	Specification: 531-649 mg Method: Current USP	
Disintegration Time:	Specification: NMT 30 minutes	Result: 11 mins.	Method: Current USP
Reference:	B1354p51,52, B1349p100		

DIETARY INGREDIENTS

Ingredient Name	LC/2caps	Result	% of LC	Spec	Method
Vitamin A (as natural beta carotene)	600.00 mcg (2000.00 IU)	834.10 mcg	139.07	100-150%	HPLC
Vitamin D (as Cholecalciferol)	25.00 mcg (1000.00 IU)	29.55 mcg	118.20	100-150%	HPLC
Vitamin E (as d-alpha tocopheryl acetate)	40.20 mg (60.00 IU)	44.28 mg	110.15	100-150%	HPLC
Vitamin B-12 (as methylcobalamin)	300.00 mcg	300.00 mcg	100.00	100-150%	**
Coenzyme Q10 (as ubiquinone)	100.00 mg	100.00 mg	100.00	100-150%	**
Trans-Resveratrol (P cuspidatum, 99%)	30.00 mg	34.88 mg	116.25	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	0.038 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	0.005 mcg	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.004 mcg	ICP-MS

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MICROBIOLOGY

Micro Study#MB0008366	Specification	Result	Method
Total Plate Count:	<10,000 CFU/g	<10 cfu/g	Current USP
Yeast & Mold Count:	<1,000 CFU/g	<10 cfu/g	Current USP
Escherichia Coli:	Negative	ND	Current USP
Salmonella:	Negative	ND	Current USP
Staphylococcus Aureus:	Negative	ND	Current USP
Prepared by:			Date: 8/20/20
Reviewed by:			Date: 8/20/20
Approved by:			Date: 8/20/20

** - 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.