CERTIFICATE OF ANALYSIS

Product Name: Pyci	nogenol + CoQ1	0					
Product #: PY0	89			Manufa	ctured for: Purity	/ Products	
Lot#: 2110)30						
Date Manufactured: 4/20	21						
Product Appearance: #0 c	lear/clear gelatin	capsules filled w	vith a light re	ddish-bro	wn powder. Res	ult: Passed	
Weight Variation: Theoretic	al Weight: 650 mg	Specification: 585-7	15 mg Resu	ılt: 672.95 ı	mg Method: Curr	rent USP	
Disintegration Time: Speci	fication: NMT 3	0 mins. Resu	It: 11 mins	. Meth	od: Current USP		
Reference: B13	87p164, B1394p	77, B1399p15, B	1388p114				
	D	IETARY INGR	EDIENTS				
Ingredient Name		LC/2caps	Result	% of LC	C Spec	Method	
Vitamin D (as Cholecalciferol)		400.00 IU	567.81 IU	141.95	100-150%	HPLC	
Pycnogenol (Pinus pinaster) (bark)		50.00 mg	60.50 mg	120.99	100-150%	UV	
Coenzyme Q10 (as Kaneka Q10 Ubiquinone)		100.00 mg	122.53 mg	122.53	100-150%	HPLC	
Purity's BioAbsorb Complex		13.00 mg	13.00 mg	100.00	NLT 100%	**	
(Bioperine; Lactospore)	9.50 mg	11.00 mg	115.79	100-150%	HPLC		
		DTHER INGRE					
Rice flour, gelatin (capsul	e), magnesium st						
		HEAVY ME					
Heavy Metal	Specification			lt/2caps	Method		
Lead:		ximum daily dos		0	ICP-MS		
Arsenic:	\leq 10 µg/maximum daily dose		0.201 mcg		ICP-MS		
Cadmium:	\leq 4.1 µg/maximum daily dose		0.022 mcg		ICP-MS		
		MICROBIOL	LOGY				
Micro Study#MB0016327	7 Specificati	on I	Result		Method		
Total Plate Count:	N/A	1	N/A		Current USP		
Yeast & Mold Count:	< 1,000 CFU/	g 1	10 cfu/g		Current USP		
Escherichia Coli:	Negative	<u> </u>	ND		Current USP		
Salmonella:	Negative	۱	ND		Current USP		
Staphylococcus Aureus:	Negative	1	ND		Current USP		
Prepared by: Repared	Rivera				Date: 5/18/2021		
Reviewed by: Rujvidger Thaker				Date: 5/18/2021			
Approved by: Jantosh Jaw	, ,				Date: 5/18/2021		

**-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.