CERTIFICATE OF ANALYSIS

	PY029					
	1102)		1	Manufactured	l for: Purity Pre	oducts
Lot#:	213762					
Date Manufactured:	2/2022					
Product Appearance:	#0 Green vegetarian	capsule filled with	th reddish-oran	ge paste. Re	sult: Passed	
Weight Variation: Theoret	tical Weight: 590 mg S	pecification: 531-649	mg Result: 54	43.76 mg Met	hod: Current USP)
Disintegration Time: S	Specification: NMT 3	30 mins. Result:	12 mins. N	lethod: curre	nt USP	
Reference:	B1443p157,158, B1	444p43,44, B143	5p145, B1445p	21-33, B143	2p46	
DIETARY INGREDIENTS						
Ingredient Name		LC/2caps	Result	% of LC	Spec	Method
Vitamin A (as beta car	rotene) 600.00 r	ncg (2000.00 IU)	779.04 mcg	129.84	100-150%	HPLC
Vitamin D (as choleca	lciferol) 10.00 m	cg (400.00 IU)	14.64 mcg	146.40	100-150%	HPLC
Vitamin E (as d-alpha t	.00 mg (60.00 IU)	45.65 mg	114.13	100-150%	HPLC	
Vitamin B-12 (as Methylcobalamin)300.00 mcg			Positive	100.00	Positive **I	D-HPLC
Coenzyme Q10 (as ubidicarenone) 100.00 mg			100.45 mg	100.45	100-150%	HPLC
Virtiva® 60.00 mg			60.00 mg	100.00	NLT 100%	**
[Minimum 12% phosp		soybean phosphol	lipids), 5% ginl	kgo flavonglu	ucosides and 0.	5%
ginkgo terpenes (leaf)]		2 00	2 00	100 ((100 1500/	
5% ginkgo flavongluce	osides	-	3.98 mg	132.66	100-150%	HPLC
Trans-Resveratrol	950/ antro at (ma at		10.86 mg	108.60	100-150%	HPLC
(Polygonum cuspidatu) OTHER INGRE	DIENTC			
Rice bran oil, vegetable				nner chloronh	vllin	
Rice of all on, vegetable	centulose, vegetable w	HEAVY ME		pper emotoph	ly IIII	
Heavy Metal	leavy Metal Specification		Result/2caps	Method		
Lead	$\leq 2.75 \mu g/maximu$	um daily dose	0.405 mcg	ICP-MS		
Arsenic	$\leq 10 \mu g/maximum$ daily dose		0.091 mcg	ICP-MS		
Cadmium	\leq 4.1 µg/maximu	im daily dose	0.036 mcg	ICP-I	MS	
		MICROBIOI	LOGY			
Micro Study#MB002	4073 Specificati	ion	Result	Meth	od	
Total Plate Count:	<10,000 CFU	U/g 1	0 cfu/g	Curre	ent USP	
Yeast & Mold Count:	: <1,000 CFU/g		l0 cfu/g	Current USP		
Escherichia Coli:	Negative		ND	Current USP		
Salmonella:	Negative		ND	Current USP		
Staphylococcus Aureus: Negative			ND	Current USP		
Prepared by:	Date: 3/1/2022					
	Date: 3/1/2022					
Reviewed by: Kajui						

CERTIFICATE OF ANALYSIS

**- 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.