CERTIFICATE OF ANALYSIS

Product Name: Green T	ea CR Phytosome	<u>)</u>					
Product #: PY100	ŭ ŭ						
Lot#: 212825							
Date Manufactured: 9/2021							
Product Appearance: #0 green/green Chlorophyll capsules filled with a brownish orange to yellowish orange powder. Result: Passed							
Weight Variation: Theoretical V	Veight: 590 mg Spe	ecification: 531-0	649 mg Result: 5	81.41 mg Me	thod: Current USP		
Disintegration Time: Specification: NMT 30 mins. Result: 15 mins. Method: Current USP							
Reference: B1424p17,18, B1381p149, B1420p78, B1408p123							
DIETARY INGREDIENTS							
Ingredient Name	LC/2caps	Result	% of LC	Spec	Method		
Vitamin C (Ascorbic acid)	60.00 mg	61.02 mg	101.07	100-150%	HPLC		
Green Tea GreenSelect®320.00 mg320.00 mg100.00NLT 100%**Phytosome® (Camellia sinensis leaves/sunflower lecithin)100.00100.00100.00100.00							
Providing min. 41 mg EGCG	41.00 mg	49.21 mg	120.02	100-150%	HPLC		
Meriva® Curcumin Phytosome® Complex (Curcuma longa rhizome	250.00 mg /sunflower lecithin)	250.00 mg	100.00	NLT 100%	**		
Providing 45 mg Curcuminoids	45.00 mg	45.17 mg	100.38	100-150%	HPLC		
Grapeseed extract (Vitis vinifera fi	ruit) 10.00 mg	10.00 mg	100.00	NLT 100%	**		
Citrus Bioflavonoids	10.00 mg	10.00 mg	100.00	100-150%	**		
Trans-Resveratrol (from Polygonum cuspidatum root	40.00 mg	40.74 mg	101.85	100-150%	HPLC		
OTHER INGREDIENTS							
Hypromellose							
Rice flour							
Silicon dioxide							
Vegetable stearate							
Chlorophyllin							
HEAVY METALS							
Heavy Metal S	Specification		Result/2caps	Meth	nod		
	2.75 μg/maximum daily dose		0.070 mcg ICP-MS				
Arsenic: 10	10 µg/maximum daily dose		0.044 mcg ICP-MS				
Cadmium: 4	4.1 µg/maximum daily dose		0.073 mcg ICP-MS				

CERTIFICATE OF ANALYSIS

MICROBIOLOGY					
Micro Study#MB0020700	Specification	Result	Method		
Total Plate Count:	< 10,000 CFU/g	15 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:	RUNDA		Date: 10/27/2021		
Reviewed By: Rujvidyer Th	Date: 10/27/2021				
Approved By: Jantosh Jaumi			Date: 10/27/2021		

**-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.