CERTIFICATE OF ANALYSIS

Product Name:	**Vital Brilliance Formula Vege Capsules (Version 2)**	
Product #:	**PY109**	Manufactured for: Purity Products
Lot#:	214354	
Date Manufactured:	03/2022	
Product Appearance:	#0 clear/clear capsules filled with a brown powder. Result: Passed	
Weight Variation:	Theoretical Weight: 600 mg Result: 649.37 mg	Specification: 540-660 mg Method: Current USP
Disintegration Time:	Specification: NMT 30 mins.	Result: 11 mins. Method: Current USP
Reference:	B1441p88, 92	

DIETARY INGREDIENTS

Ingredient Name	LC/2caps	Result	% of LC	Spec	Method
Rhodiolife® Rhodiola (Rhodiola rosea) extract (root);	300.00 mg	300.00 mg	100.00	NLT 100%	**
9 mg Rosavins	9.00 mg	9.80 mg	108.88	100-150%	HPLC
Organic Schizandra (Schisandra chinensis) powder (whole berry)	150.00 mg	150.00 mg	100.00	NLT 100%	**
Eleuthero (Eleutherococcus senticosus) extract (root) [Standardized to min 0.8% eleutherosides]	150.00 mg	150.00 mg	100.00	NLT 100%	**
Providing 1.2mg eleutherosides	1.2 mg	Positive	100.00	Positive	**ID-HPLC
Vitality Booster Herbal Blend	200.00 mg	200.00 mg	100.00	NLT 100%	**
Asian ginseng (Panax ginseng extract (root) [Standardized to min 2% Ginsenosides (1.333mg)]					
Ashwagandha extract (Withania somnifera) (root) [Standardized to min 4% Withanolides (2.667mg)					
Goji berry (Lycium Barbarum) extract (ripe fruit) [Standardized to min 40% Polysaccharides (26.667 mg)]					

OTHER INGREDIENTS

Vegetable cellulose (capsule), rice flour and vegetable stearate.

HEAVY METALS

Heavy Metal	Specification	Result/2caps	Method
Lead:	2.75 µg/maximum daily dose	0.169 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	0.095 mcg	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.065 mcg	ICP-MS

MICROBIOLOGY

Micro Study# MB0024944	Specification	Result	Method
Total Plate Count:	< 10,000 CFU/g	155 CFU/g	Current USP
Yeast & Mold Count:	< 1,000 CFU/g	<10 CFU/g	Current USP
E. Coli:	Negative	ND	Current USP
Salmonella:	Negative	ND	Current USP
Staph Aureus:	Negative	ND	Current USP

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Prepared By: Salma Jahan	Date: 4/5/2022
Reviewed By: Rajvidya Thakker	Date: 4/5/2022
Approved By: Santosh Kanwar	Date: 4/5/2022

**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.