

CERTIFICATE OF ANALYSIS

Product Name:	Vital Focus Capsules		
Product #:	PY067	Manufactured for: Purity Products	
Lot#:	211727		
Date Manufactured:	6/2021		
Product Appearance:	#1 Clear/Clear vegetarian capsule filled with tan to light brown powder. Result: Passed		
Weight Variation:	Theoretical Weight: 445 mg Specification: 400.50-489.50 mg Result: 449.20 mg Method: Current USP		
Disintegration Time:	Specification: NMT 30 minutes Result: 12 mins. Method: Current USP		
Reference:	B1397p98, B1408p1,2, B1399p118		

DIETARY INGREDIENTS

Ingredient Name	LC/2caps	Result	% of LC	Spec	Method
Synapsa™ Bacopa Monniera Extract (leaf)	150.00 mg	150.00 mg	100.00	NLT 100%	**
Standardized to 55% Bacosides (82.50 mg)	82.50 mg	82.50 mg	100.00	NLT 100%	**
8% Bacosides by HPLC	12.00 mg	17.62 mg	146.83	100-150%	HPLC
CereBoost American Ginseng (Panax quinquefolius) extract (root)	200.00 mg	200.00 mg	100.00	NLT 100%	**
10% Ginsenosides	20.00 mg	29.89 mg	149.45	100-150%	HPLC
Schisandra Extract (fruit and seeds)	50.00 mg	50.00 mg	100.00	NLT 100%	**
9% Schizandrins	4.50 mg	4.50 mg	100.00	100-150%	**
Vinpocetine	10.00 mg	10.13 mg	101.30	100-150%	HPLC

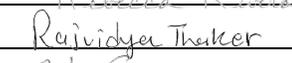
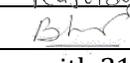
OTHER INGREDIENTS

Rice Flour, Vegetable Cellulose (capsule) and Vegetable Stearate
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HEAVY METALS

Heavy Metal	Specification	Result (2caps)	Method
Lead:	2.75 µg/maximum daily dose	0.045 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	0.221 mcg	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.011 mcg	ICP-MS

MICROBIOLOGY

Micro Study#	Specification	Result	Method
MB0017476			
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: 6/28/2021
Reviewed By: 			Date: 6/28/2021
Approved By: 			Date: 6/28/2021

** - 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.