## **CERTIFICATE OF ANALYSIS**

Product Name:	Vital Super Reds Veg	e Capsules				
Product #:	PY096			Manufactured for: Purity Products		
Lot#:	183608					
Date Manufactured:	1/2019					
Product Appearance:	#00 clear/clear oblong	vege capsules fi	lled with a reddi	sh brown pow	der Result: Pass	sed
Weight Variation: (	Current USP) – Target V	Weight: 770.00 r	ng Average	Weight: 797.	00 mg	
Disintegration Time:	(Current USP) – Speci	fication: NMT 3	0 minutes R	esult: 15 min	nutes	
Reference:	B1280p23					
	]	DIETARY ING	REDIENTS			
Ingredient Name		LC/2caps	Result	% of LC	Spec	Method
Organic Cranberry (Organic PACran®) (Vaccinium macrocarpon) Fruit)		500.00 mg	500.00 mg	100.00	NLT 100%	**
Organic Schisandra Freeze Dried powder (Schisandar chinensis) (Whole berry)		400.00 mg	400.00 mg	100.00	NLT 100%	**
Wolfberry Goji Extract (fruit)	t (Lycium barbarum)	200.00 mg	200.00 mg	100.00	NLT 100%	**
Pomegranate P40p extr		40.00 mg	40.00 mg	100.00	NLT 100%	**
(fruit skin) 16mg/40%	punicosides	16.00 mg	17.06 mg	106.65	100-150%	HPLC
		OTHER INGR				
Hydroxypropyl methyle	cellulose, rice flour, mag	·		xtrin.		
		HEAVY M				
Heavy Metal	Specification		Result/2caps	Method		
Lead:	$\leq 2.75 \mu$ g/maximum daily dose		0.088 mcg	ICP-MS		
Arsenic:	$\leq$ 10 µg/maximum daily dose		0.099 mcg	ICP-MS		
Cadmium:	$\leq$ 4.1 µg/maximum dai		0.017 mcg	ICP	-MS	
		MICROBIC	DLOGY			
Micro Study#191766	Specification	Result	t	Method		
Total Plate Count:	NMT 10,000 CFU/g 180 CF		FU/g	Current USP Method		
Yeast & Mold Count:	NMT 1,000 CFU/g <10 CFU		U/g	Current USP Method		
Escherichia Coli:	Negative	Negative ND		Current USP Method		
Salmonella:	Negative	ND		Current USI	P Method	
Staphylococcus Aureus: Negative ND				Current USI	P Method	
Prepared By:		Date: 2/18	8/19			
Reviewed By:		Date: 2/18	8/19			
Approved By: & Ventates waren				Date: 2/18		
**- In Accordance with 2	21 CFR 111.75(d) (1) the	following finishe	d dietary ingredi	ents product sp	pecifications are	

\*\*- 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 and are therefore quantitatively verified to meet the dietary supplement specifications at the finished batch stage as per test method QP#1358.