CERTIFICATE OF ANALYSIS

Product Name:	**Vitamin C&D Super Immune**	
Product #:	**PY010**	Manufactured for: Purity Products
Lot#:	201156	
Date Manufactured:	4/2020	
Product Appearance:	# #00 clear/clear gelatin capsules filled with a light brown powder. Result:	
Weight Variation:	(Current USP) – Target Weight: 934.20-1,141.80 mg	Average Weight: 1034.43mg
Disintegration Time:	(Current USP) – Specification: NMT 30 minutes	Result: 10 minutes
Reference:	B1335p144, 145, B1318p192	

DIETARY INGREDIENTS

Ingredient Name	LC/1cap	Result	% of LC	Spec	Method
Vitamin C (from Pureway-C®)	500.00 mg	537.81 mg	107.56	100-150%	Titration
Proprietary formulation of ascorbic acid USP combined with lipid metabolites (fatty acids from vegetable waxes)					
Vitamin D (Cholecalciferol)	25.00 mcg (1000 IU)	25.00 mcg (1000 IU)	100.00	100-150%	**
Acerola extract 4:1 (Malpighia punicifolia L.) (Fruit) (Equivalent to 10mg Acerola powder)	2.50 mg	2.50 mg	100.00	NLT 100%	**
Citrus Bioflavonoid Complex	170.00 mg	170.00 mg	100.00	NLT100%	**
(85 mg total bioflavonoids, including Hesperidin)	85.00 mg	113.42 mg	133.43	100-150%	HPLC
Rose Hips extract 4:1 (Rosa canina) (fruit) (equivalent to 10mg rose hips powder)	2.50 mg	2.50 mg	100.00	NLT 100%	**
Rutin	10.00 mg	11.04 mg	110.40	100-150%	HPLC

OTHER INGREDIENTS

Gelatin (capsule), Rice flour, Vegetable stearate

HEAVY METALS

Heavy Metal	Specification	Result	Method
Lead:	≤ 2.75 µg/maximum daily dose	0.932 mcg	ICP-MS
Arsenic:	≤ 10 µg/maximum daily dose	0.065 mcg	ICP-MS
Cadmium:	≤ 4.1 µg/maximum daily dose	0.006 mcg	ICP-MS

MICROBIOLOGY

Micro Study#MB0004652	Specification	Result	Method
Total Plate Count:	< 10,000 CFU/g	<10 cfu/g	Current USP
Yeast & Mold Count:	< 1,000 CFU/g	5 cfu/g	Current USP
Escherichia Coli:	Negative	ND	Current USP
Salmonella:	Negative	ND	Current USP
Staphylococcus Aureus:	Negative	ND	Current USP
Prepared by:			Date: 4/20/20
Reviewed by:			Date: 4/20/20
Approved by:			Date: 4/20/20

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