CERTIFICATE OF ANALYSIS

Product Name: Advanced IP-6 Ve	ege Capsules						
Product # PY025	Manufactured for: Purity Products						
Lot#: 214356							
Date Manufactured: 3/2022							
Product Appearance: #00 Clear/Clear Ve	ege capsule filled	with light yellow	v powder. R	esult: Passed			
Weight Variation:Theoretical Weight: 840 mg Result: 842.66 mgSpecification: 756-924 mg Method: Current USP							
Disintegration Time: Specification: NMT	30 mins. Res	sult: 10 mins. M	lethod: Curre	ent USP			
Notebook Reference: B1441p130, B1435p186, B1444p94							
DIETARY INGREDIENTS							
Ingredient Name	LC/4caps	Result	% of LC	Spec	Method		
Vitamin D (as cholecalciferol)	1000.00 IU	1038.94 IU	103.89	100-150%	HPLC		
Calcium	350.00 mg	354.00 mg	101.14	100-150%	ICP		
(as calcium magnesium phytate, calcium carbonate)							
Magnesium	160.00 mg	237.76 mg	148.60	100-150%	ICP		
(as magnesium oxide, magnesium citrate, calcium magnesium phytate)							
Coenzyme Q10 (as ubidicarenone)	20.00 mg	20.51 mg	102.54	100-150%	HPLC		
Green Tea Extract	48.00 mg	48.00 mg	100.00	NLT 100%	**		
(Camellia sinensis) (dried leaves) (decaffeinated) [Standardized for 60% Polyphenols (28.8 mg]							
Inositol Hexaphosphate	1600.00 mg	1600.00 mg	100.00	NLT 100%	**		
(as calcium magnesium phytate)							
Milk Thistle Extract (Silybum marianus) (seeds) [Standardized for 80% silymarin (100.00 mg (80 mg)]	100.00 mg	100.00	100-150%	**		
and 30% silymarin by HPLC	30.00 mg	30.00 mg	100.00	100-150%	**		
Quercetin (as dehydrate)	100.00 mg	100.00 mg	100.00	100-150%	**		
(Dimorphandra gardeniaria Fam. Leguminosae) (seeds)							
OTHER INGREDIENTS							
Vegetable cellulose							
Magnesium stearate							
Rice flour							
HEAVY METALS							
Heavy Metal Specification	Specification		Method				
Lead: $\leq 2.75 \ \mu g/maximum$	\leq 2.75 µg/maximum daily dose		ICP-MS				
	$\leq 10 \ \mu g/maximum$ daily dose		ICP-MS				
Cadmium: $\leq 4.1 \ \mu g/maximum$	a daily dose	0.060 mcg	ICP-N	МS			

CERTIFICATE OF ANALYSIS

MICROBIOLOGY					
Micro Study#MB0024553	Specification	Result	Method		
Total Plate Count:	<10,000 CFU/g	135 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: Rebecca Rivera			Date: 3/21/2022		
Reviewed by: Rujvidger Thaker			Date: 3/21/2022		
Approved by: Santost Janu		Date: 3/21/2022			

**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.