CERTIFICATE OF ANALYSIS

Product Name:	**My™ Metabolism™ Pro Clinical Vege Capsules**
Product #:	**PY117** Manufactured for: Purity Products
Lot#:	221544
Date Manufactured:	9/2022
Expiration Date:	9/2024
Product Appearance:	#00 clear/clear oblong vege capsules filled with a beige to brown powder. Result: Passed
Weight Variation:	Theoretical Weight: 920 mg Specification: 828-1012 mg Result: 924.56 mg Method: Current USP
Disintegration Time:	Specification: NMT 30 mins. Result: 7 mins. Method: Current USP
Reference:	ATDS#1638-00/22, 1638-03/22

DIETARY INGREDIENTS

Ingredient Name	LC/1cap	Result	% of LC	Spec	Method
Vitamin D (as cholecalciferol, D3)	2000.00 IU (50 mcg)	2452.27 IU	122.61	100-150%	HPLC
Chromium (from Chromax® chromium picolinate)	200.00 mcg	245.00 mcg	122.50	100-150%	ICP
Sinetrol® -XPurC Citrus grandis Osbeck,	630.00 mg	630.00 mg	100.00	NLT 100%	**
Grapefruit extract (Citrus paradisi Macfad), Citrus sinensis, Guarana seed (Paullinia cupana) and Orange juice concentrate (Citrus sinensis)					
[std. to 42% total flavanones	264.60 mg	264.60 mg	100.00	100-150%	**
and 3.5% Caffeine	22.05 mg	22.79 mg	103.35	100-150%	HPLC

OTHER INGREDIENTS

Vegetable cellulose, rice flour, magnesium stearate and silicon dioxide.

HEAVY METALS

Heavy Metal	Specification	Result/1cap	Method
Lead:	2.75 µg/maximum daily dose	0.470 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	0.298 mcg	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.012 mcg	ICP-MS

MICROBIOLOGY

Micro Study#MB0028908	Specification	Result	Method
Total Plate Count:	< 10,000 CFU/g	125 cfu/g	Current USP
Yeast & Mold Count:	< 1,000 CFU/g	15 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: 10/4/2022
Reviewed By: Rajvidya Thakker			Date: 10/5/2022
Approved By: Santosh Lawni			Date: 10/5/2022

**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.