CERTIFICATE OF ANALYSIS

Product Name:	**MyBiotin Proclinical Vege Capsules**
Product #:	**PY111** Manufactured for: Purity Products
Lot#:	220362
Date Manufactured:	3/2022
Expiration Date:	3/2024
Product Appearance:	#0 clear/clear vege capsules filled with a light pink to light reddish powder. Result: Passed
Weight Variation:	Theoretical Weight: 540 mg Specification: 486-594 mg Result: 535.50 mg Method: Current USP
Disintegration Time:	Specification: NMT 30 mins. Result: 20 mins. Method: Current USP
Reference:	ATDS# 0133-00/22, 0133-01/22

DIETARY INGREDIENTS

Ingredient Name	LC/1cap	Result	% of LC	Spec	Method
Biotin (from Magnesium biotinate)	10,000.00 mcg	12442.53 mcg	124.43	100-150%	HPLC
Lustriva™ (Inositol-stabilized Arginine Silicate, Magnesium biotinate)	160.00 mg	160.00 mg	100.00	NLT 100%	**
Astaxanthin (as AstaREAL™)	4.00 mg	4.27 mg	106.75	100-150%	UV

OTHER INGREDIENTS

Vegetable cellulose (capsule), rice flour and magnesium stearate

HEAVY METALS

Heavy Metal	Specification	Result/1cap	Method
Lead:	2.75 µg/maximum daily dose	0.059 mcg	ICP-MS
Arsenic:	10 µg/maximum daily dose	0.031 mcg	ICP-MS
Cadmium:	4.1 µg/maximum daily dose	0.063 mcg	ICP-MS

GLUTEN

Allergen	Method	Specification	Result
Gluten	ELISA	< 20 ppm	< 20 ppm

MICROBIOLOGY

Micro Study# MB0025355	Specification	Result	Method
Total Plate Count:	< 10,000 CFU/g	20 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: 4/21/2022
Reviewed By: Rajvidya Thakur			Date: 4/21/2022
Approved By: Santosh Dawri			Date: 4/21/2022

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.