

# CERTIFICATE OF ANALYSIS

Product Name:	<b>AstaFX Plus</b>	
Product #:	<b>PY084</b>	Manufactured for: Purity Products
Lot#:	193309	
Date Manufactured:	8/2020	
Product Appearance:	#0 Green/Green opaque vege capsule filled with yellowish/red oil/paste.	
Weight Variation:	(Current USP) – Target Weight: 0.6783 g	Average Weight: 0.5974g
Disintegration Time:	(Current USP) – Specification: NMT 30 minutes	Result: 11 minutes
Reference:	B1334p105, B1349p99	

## DIETARY INGREDIENTS

Ingredient Name	LC/2caps	Result	% of LC	Spec	Method
Astaxanthin	5.00 mg	Positive	100.00	Positive	ID-HPLC**
Carotenoid Co-Factor Blend	195.00 mg	195.00 mg	100.00	NLT 100%	**
Carrot Powder	70.00 mg	70.00 mg	100.00	NLT 100%	**
Organic Spirulina	65.00 mg	65.00 mg	100.00	NLT 100%	**
Spinach powder	55.00 mg	55.00 mg	100.00	NLT 100%	**
Lycopene Concentrate	2.50 mg	Positive	100.00	Positive	ID-HPLC**
Lutein (from Marigold flower extract)	2.50 mg	Positive	100.00	Positive	ID-HPLC**
Bioperine® (Black pepper fruit extract)	5.00 mg	5.10 mg	101.92	100-150%	HPLC

## OTHER INGREDIENTS

Rice flour
Cellulose (vegetarian capsule)
Magnesium stearate
Silica
Titanium dioxide
Chlorophyll (color)

## HEAVY METALS

Heavy Metal	Specification	Result/2caps	Method
Lead:	≤ 2.75 µg/maximum daily dose	0.050 mcg	ICP-MS
Arsenic:	≤ 10 µg/maximum daily dose	0.013 mcg	ICP-MS
Cadmium:	≤ 4.1 µg/maximum daily dose	0.018 mcg	ICP-MS

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### MICROBIOLOGY

<b>Micro Study#MB0008365</b>	<b>Specification</b>	<b>Result</b>	<b>Method</b>
Total Plate Count:	< 10,000 CFU/g	250 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:			Date: 8/20/20
Reviewed By:			Date: 8/20/20
Approved By:			Date: 8/20/20

\*\* - 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.