## **CERTIFICATE OF ANALYSIS**

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Product Name:	Asta-FX Liquid Ve	ege Capsules					
	roduct Code: PY041			Manufactured for: Purity Products			
Lot#:	213270						
Date of Manufacture	: 1/2022						
Product Appearance	/Color: #0 Green/Green	n Opaque veg	ge-capsule filled	d with yellowis	h/red oil-paste	•	
e	Theoretical Weight: 590 Method: Current USP	) mg Spec	eification: 531-6	649 mg Rest	ult: 594.35 mg		
Disintegration Time:	: Specification: NMT 3	0 mins. Re	esult: 6 mins.	Method Curren	t USP		
Notebook Reference	: B1442p7,8, B1441p	31, B1432p11	2				
DIETARY INGREDIENTS							
Ingredient Name		LC/2caps	Result	% of LC	Spec	Method	
Astaxanthin (AstaRI	EAL® Astaxanthin)	4.00 mg	5.86 mg	146.50	100-150%	UV	
EVNol <sup>TM</sup> Tocotriend	x 54.04 mg	54.04 mg	100.00	NLT 100%	**		
Mixed Tocotrienols (from EVNOL <sup>TM</sup> )		20.00 mg	27.35 mg	136.75	100-150%	HPLC	
d-alpha tocopherol (from EVNOL <sup>TM</sup> )		5.12 mg	7.28 mg	142.19	100-150%	HPLC	
Black Pepper Extrac	5.00 mg	5.00 mg	100.00	NLT 100%	**		
(Bioperine)		4.75 mg	5.40 mg	11368	100-150%	HPLC	
		OTHER ING	REDIENTS				
Organic extra virgin Oil, and Chlorophyll	olive oil, Vegetable Colin.	ellulose, Vege	table Wax, d-al	pha Tocophery	l Acetate, and	Rosemary	
HEAVY METALS							
Heavy Metal	Specification		Result /2cap	sult /2caps Method			
Lead:	2.75 µg/maximum d	aily dose	0.339 mcg	ICP-	ICP-MS		
Arsenic:	10 μg/maximum dai	ly dose	0.155 mcg	ICP-	ICP-MS		
Cadmium:			0.028 mcg	ICP-	ICP-MS		
MICROBIOLOGY							
Micro Study#MB00	023487 Specificati	on	Result	Meth	hod		
Total Plate Count:	otal Plate Count: <10,000 CFU/g		155 cfu/g	Curre	Current USP		
Yeast & Mold Count: <1,000 CFU/g		/g	<10 cfu/g	Curre	Current USP		
Escherichia Coli: Negative		-	ND	Curre	Current USP		
Salmonella: Negative			ND	Curre	Current USP		
Staphylococcus Aureus: Negative			ND	Curre	Current USP		
Prepared by:		Date: 2/7/2022					
Reviewed by: Rajvidyer Theker					Date: 2/7/2022		
Approved by: Jantoshdauni					: 2/7/2022		
	21 CFR 111.75(d) (1) the	6 - 11	ad diatawa in anad				

\*\*- 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.