Title: EPO Evening Primrose Oil, 500 mg

Allergy Research Group® / NutriCology®

Document No.: 71260CA

Revision: 09

Certificate of Analysis

Part Number	71260				
Lot Number	200662	200662			
Product Count & Form	120 Softgels	120 Softgels			
Packaging, Closure	175 cc bottle, desi	175 cc bottle, desiccant			
Storage Condition	Room Temperatu	re			
Shelf Life	36 months	36 months Expiration Date 08/2023			
Label Number, Size	71260L, F				
Retain Sample	2 bottles total				

ANALYSIS

Physical

Attribute	Method / SOP	Specification	Test Result
Product Color, Size	QC011	Yellow, Clear, Oval 10, S2	Complies
Product Weight Variation	QC012	Meets USP <2091> Expected 620-840 mg	Pass, 784 mg
Disintegration time	QC018	Meets USP <2040>	Pass, 3 min*

*tested by CMO

Purity/Strength per Softgel

Active Ingredients	Label Claim	Release Limits	Test Result	
Evening Primrose Oil	500 mg	100 - 150% of LC by input	100 - 150% of LC by input*	
Gamma-Linolenic Acid	45 mg	100 - 150% of LC by GC	48 mg/softgel	
Linoleic Acid	365 mg	100 - 150% of LC by GC	375 mg/softgel	
Oleic Acid	35 mg	100 - 150% of LC by GC	100 - 150% of LC by GC*	
Other Ingredients: Gelatin, Glycerin, Purified water				

*Production and Process Control

Identification \ Chemical \ Heavy Metals

Identification (Chemical) Heavy Metals					
Attribute	Method / SOP	Specification	Test Result		
Identity	Bulk Certificate of Analysis Review	Approved	Approved		
Arsenic (inorganic)	ICP-MS per USP <2232>	\leq 15 µg/day Ψ	0.014 µg/day		
Cadmium	ICP-MS per USP <2232>	\leq 5 µg/day Ψ	0.005 µg/day		
Mercury (total)	ICP-MS per USP <2232>	\leq 15 µg/day Ψ	0.030 µg/day		
Lead	ICP-MS per USP <2232>	$\leq 10 \ \mu g/day \ \Psi$	0.089 µg/day		

 Ψ Permitted Daily Exposure, BQL- Below Quantitative Limit

Microbiology

Attribute	Method / SOP	Specification	Test Result
Total Aerobic Count	USP/ AOAC/ FDA BAM	$\leq 1 \ge 10^4 \text{CFU/g}$	<10 CFU/g
Total Yeast & Mold	USP/ AOAC/ FDA BAM	$\leq 1 \text{ x } 10^3 \text{ CFU/g}$	<10 CFU/g
Coliform	USP/ AOAC/ FDA BAM	$\leq 1 \ge 10^2 \text{ CFU/g}$	<10 CFU/g
Escherichi coli	USP/ AOAC/ FDA BAM	None Detected	None Detected
Staphylococcus aureus	USP/ AOAC/ FDA BAM	None Detected	None Detected
Salmonella	USP/ AOAC/ FDA BAM	None Detected	None Detected

Approval:	Cy dag	Casey Van Wagoner 2020.12.16 10:56:09 -07'00'	Date: 16DEC2020
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40 West Louise Ave., Salt Lake City, UT 84115 Phone: (801) 485-1800 Fax: (801) 484-9211 Email: utlab@advancedlabsinc.com FDA Registration #14353128308

Test Certificate

Description:	Evening Primrose Oil
Sample ID:	41260
Lot No:	VS200662
Location: Received: Completed:	10/13/2020 10/20/2020

Client: Allergy Research Group 2300 S Main St Salt Lake City, UT 84115

Lab No: 204744-03

Analysis	Result	Per Unit	Specifications	Method
Mercury	0.019	µg/g	< 0.5 µg/g	ICP-MS USP <730>
Lead	0.057	μg/g	$< 0.5 \ \mu g/g$	ICP-MS USP <730>
Arsenic	0.009	μg/g	$< 0.5 \ \mu g/g$	ICP-MS USP <730>
Cadmium	0.003	μg/g	$< 0.5 \ \mu g/g$	ICP-MS USP <730>
Total Aerobic Microbial Count	<10	CFU/g	<= 10,000 CFU/g	USP <2021>
Coliform	<10	CFU/g	<= 100 CFU/g	AOAC 991.14
E.Coli	Absent	per 10 grams	Absent	USP <2022>
Staphylococcus aureus	Absent	per 10 grams	Absent	USP <2022>
Salmonella	Absent	per 10 grams	Absent	USP <2022>
Total Yeast & Mold	<10	CFU/g	<= 1,000 CFU/g	USP <2021>
Yeast *	<10	CFU/g	<= 1,000 CFU/g	USP <2021>
Mold *	<10	CFU/g	<= 1,000 CFU/g	USP <2021>

* For informational purposes only.

THESE RESULTS APPLY ONLY TO THE SAMPLE SUBMITTED AND NOT TO THE PRODUCT FROM WHICH IT WAS TAKEN. THESE RESULTS ARE PROVIDED ONLY FOR THE BENEFIT OF CLIENT, WITHOUT REPRESENTATION OR WARRANTY OF ANY KIND, EXCEPT FOR THE EXPRESS LIMITED WARRANTY PROVIDED SOLELY TO CLIENT IN ADVANCED LABORATORIES' TERMS OF SERVICE.

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Dated: 10/20/2020

Jeni Miller-Administrative Assistant

Printed: 10/20/2020 4:52:46 PM

Results Approved By:



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Micro Quality Labs, Inc.

Specializing in Pharmaceutical, Dietary Supplements, Toys and Cosmetics Testing 3125 N. Damon Way • Burbank, California 91505 (818) 845-0070 • Fax: (818) 845-0030 E-Mail: <u>Karine@MicroQualityLabs.com</u>

Customer:Allergy Research Group, LLCAddress:2300 S. Main stSalt Lake City, UT 84115

ANALYTICAL/CHEMICAL CERTIFICATE OF ANALYSIS

Sample Name: EVENING PRIMROSE OIL Product Code: 41260 Batch/Lot: VS200662 MQL Accession: 201013-0172

PO#: 78223-1012 Sample Description: BULK Rush: N/A Received Date: 10/13/20

Test Requested:	Test Method:	Specification:	Results:
Linolenic Acid	MQLTM-0201	NLT 365 mg/softgel	375 mg/softgel
	By GC-FID		
Gamma Linoleic Acid	MQLTM-0201	NLT 45 mg/softgel	48 mg/softgel
(GLA)	By GC-FID		
Prepared(By: Diana Zavala/Document Control Specialist		OCT 1 9 2020	
Reviewed By:		OCT 1 9 2020	
Esme Arellano/QA Assistant		10/19/20	

The aforementioned results on this report are representative of the samples submitted and may not be indicative of the entire manufacture, batch, and/or lot. Applicable current GMP's shall always be used when sampling. GLP's shall always be practiced by Micro Quality Labs to ensure the most accurate results.

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