MiramarLab
CUSTOM MANUFACTURING & PRODUCT DEVELOPMENT

MANUFACTURE DECLARATION LETTER

Manufacture FDA Registration # 14574831328

Client: 1Q SOLUTIONS LLC

To Whom it May Concern:

The undersigned does hereby declare that the goods described herein are manufactured in and are products of the United States of America. The undersigned does also hereby declare that the goods described herein are freely sold in the United States of America in an FDA & GMP Inspected Facility #14574831328.

Miramar Lab is engaged in business with 1Q SOLUTIONS LLC in the development and manufacturing of 1Q SOLUTIONS LLC brand products. See below list of current products.

PRODUCT MANUFACTURE LIST:

- FOCUS (2oz, 60ml)
- IMMUNE (2oz, 60ml)
- RELAX (2oz, 60ml)

The undersigned grants distribution right and the full authority to process any necessary documentation on behalf of manufacture to be import the product to Venezuela. See below authorize imported.:

CASA DE REPRESENTACION COSMELAB VENEZUELA, C.A..

Address:

Urbanizacion Monte Rey Calle B,
Municipio Baruta.
Venezuela

For any question or concerns feel free to contact me at 305-455-5017 EXT 1001 or email vr@miramarlab.com.

Date: 7/3/25

Sincerely,
Victor M. Ramirez
CEO

2025

CERTIFICATE OF REGISTRATION

This certifies that:

Miramar Cosmetics Inc
2301 Nw 107th Ave Ste 101
Doral, FL 33172-2177
United States

is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as currently effective on the date hereof by Registrar Corp:

U.S. FDA Registration No.:	**14574831328**
U.S. FDA UFI (DUNS) No.:	**012873243**
U.S. Registration Agent:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

This certificate affirms that the above stated facility is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as effective by Registrar Corp as of the date hereof, and Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until December 31, 2025, unless such registration has been terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: November 14, 2024

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Audit recognition

This certificate has been awarded to

Miramar Cosmetics, Inc., dba Miramar Lab

after having conducted an audit at

2301 NW 107th Ave, Suite 101, Doral, FL 33172

Audit Date(s):
03/12/2025 – 03/13/2025

Certificate Number:
FSS201910-2964_3

Certificate Expiry Date:
03/13/2026

Audit Score:
84%

Eurofins GMP for Dietary Supplements

(Per requirements as listed in 21 CFR part 111)

Authorised by:

Brian Neal
Technical Manager

Eurofins Food Assurance
2120 Rittenhouse st., Suite A

Des Moines, IA 50321, USA

Ph:(515)299-6979

e-mail address

www.eurofins.com/foodassurance

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