amare®

CERTIFIED

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Due to a confidentiality agreement signed between Amare Global and the companies that manufacture our products, the names of those companies have been withheld.

amare

UL Verification Services — GMP Inspection Certificate
(UL — Certificate No: i10-34169)

Sunset product is manufactured issued by UL (Underwriters Laboratories), this inspection certificate confirms full compliance with 21 CFR Part 111 — the most current U.S. federal cGMP regulation for dietary supplements. The scope covers manufacturing, packaging, and warehousing of liquid, powder, tablet, capsule, and softgel dosage forms.

UL Solutions

UL VERIFICATION SERVICES INC. ISSUES THIS

CERTIFICATE OF INSPECTION

[redacted]

FOLLOWING INSPECTION OF ITS GOOD MANUFACTURING PRACTICE & QUALITY SYSTEM AND FINDING IT IN CONFORMANCE WITH:

Retail Certification Program Requirements and

ISO 22716:2007 and 2013 FDA Cosmetic Guidelines & ANSI 455-3 GMP for Cosmetics: 2021

COSMETIC GUIDELINES ON GOOD MANUFACTURING PRACTICES FOR THE FOLLOWING SCOPE OF INSPECTION:

The Manufacturing, Packaging and Warehousing of Cosmetic Softgel Capsules

Certificate Number: i10-34169-1
Issue Number: 8
Certificate Issue Date: 8/7/2024
Expiration Date: 3/30/2028
Audit Due Date: Decemer 27

Authorized by:

Karen Sak, Business Manager

UL Verification Services Inc.
7036 Snowdrift Road, Suite 200
Allentown, PA 18106
United States of America
800-903-5660

The UL Solutions Logo, Enhanced Certification Mark, and Accreditation Marks indicate satisfactory assessment against the above noted standard / requirements in accordance with the GMP Procedure for Certification, the UL Verification Services LLC Agreement for Use of Certification Symbols, and the scope of assessment. This certificate remains the property of UL Solutions, to whom it must be returned upon request. Revision 4/1/2024

UL Solutions

UL VERIFICATION SERVICES INC. ISSUES THIS

CERTIFICATE OF CONFORMANCE

[redacted]

FOLLOWING ASSESSMENT OF ITS GOOD MANUFACTURING PRACTICE & QUALITY SYSTEM AND FINDING IT IN CONFORMANCE WITH:

Retail Certification Program Requirements and

ISO 22716:2007 and 2013 FDA Cosmetic Guidelines & ANSI 455-3 GMP for Cosmetics: 2021

COSMETIC GUIDELINES ON GOOD MANUFACTURING PRACTICES FOR THE FOLLOWING SCOPE OF CERTIFICATION:

The Manufacturing, Packaging and Warehousing of Cosmetic Softgel Capsules

Certificate Number: a10-34169-1
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amare

California Organic Processed Product Registration
(California Department of Public Health — No: 20895)

This is from the facility where our **Sunset** product is manufactured registered under the California Organic Food and Farming Act as organically processed products. This registration confirms that those products are legally authorized to carry an organic label under California state law.

STATE OF CALIFORNIA

DEPARTMENT OF PUBLIC HEALTH
FOOD AND DRUG BRANCH

ORGANIC PROCESSED PRODUCT REGISTRATION

[redacted]

REGISTRATION NUMBER: 20895
EXPIRATION DATE: 7/8/2026

THE BUSINESS NAMED HEREIN IS REGISTERED IN THE STATE OF CALIFORNIA TO PROCESS AND/OR HANDLE PROCESSED PRODUCTS "SOLD AS ORGANIC". THIS REGISTRATION IS ISSUED IN ACCORDANCE WITH THE CALIFORNIA HEALTH AND SAFETY CODE AND IS NOT TRANSFERABLE TO ANY OTHER PERSON OR PLACE. THE REGISTRANT IS REQUIRED BY LAW TO IMMEDIATELY NOTIFY THE CALIFORNIA DEPARTMENT OF PUBLIC HEALTH OF ANY CHANGE IN THE INFORMATION REPORTED IN THE APPLICATION.

Food and Drug Branch, 1500 Capitol Avenue, MS 7602, PO Box 997435, Sacramento, CA 95899-7435 (916) 650-6500

Printed: 7/11/2025

OSP 07 102308

FDA Food Facility Registration (Dietary Supplement Manufacturer)
(FDA Registration No: 13618051326)

The facility where our **Sunset** product is manufactured, the facility producing our dietary supplement products is registered with the FDA under the Food Safety Modernization Act (FSMA). This registration confirms that the facility's food safety systems meet federal oversight standards.

2026

CERTIFICATE OF REGISTRATION

This certifies that:

[redacted]

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.:	**13618051326**
U.S. FDA UFI (DUNS) No.:	**831560578**
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, 2026, 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 Lemarz
Executive Director
Registrar Corp
Dated: August 28, 2025

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NSF/GRMA Certificate of Conformance — Dietary Supplement GMP (NSF International — Certificate No: C0040870)

Issued by NSF International, one of the world's most trusted independent certification bodies, this certificate confirms that our **Sunset** manufacturing facility complies with NSF/ANSI 455-2 — the Good Manufacturing Practices standard for Dietary Supplements.

GRMA AuditSphere 455-2 DS

CERTIFICATE	CERT-000366
Audit Number	AO-000663
Issue Date:	03-25-2024
Expiration:	08-06-2026
First Certification:	03-25-2024
Annual Surveillance Date	01-29-2025

CERTIFICATION	
Certification Date:	03-25-2024
Expiration:	08-06-2026

CERTIFICATE OF CONFORMANCE

[redacted]

Facility Type & Description: Campus/Multi-facility, Distributor Onsite, Manufacturing, Primary Facility

Following The Audit Of The Organization's Quality System, Good Manufacturing Practices, And Finding Conformance With:
Global Retailer & Manufacturer Alliance, Inc. Certification Program v1.2
Standard: NSF/ANSI 455-2 - 2021 for Dietary Supplements
Product Types and Manufacturing Stages listed on Page 2
Audit Identification: Onsite

Authorised by David Trosin, Managing Director, Health Sciences Certification

The GRMA Logo, GRMAuditSphere Logo, NSF International's Logo, etc. indicate satisfactory assessment against the Global Retailer & Manufacturer Alliance, Inc.'s Certification Program & NSF/ANSI 455-2 - 2021 for Dietary Supplements. The certification remains the property of NSF International, to whom it must be returned upon request.
Verify Current Certification Status Through Retailer Member Directory

Certificate Issuer
NSF International
789 N. Dixboro Road
Ann Arbor, Michigan, 48105, UNITED STATES
Accreditation # 0216

ANAB ANSI National Accreditation Board ACCREDITED ISO/IEC 17065 PRODUCT CERTIFICATION BODY

Powered by Intact Platform

Page 1|3
Valid certificate consists of 3 pages.
Audit Number | AO-000663

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NSF Certificate of Conformity — Dietary Supplement GMP (NSF/ANSI 455-2:2021)

This is NSF's complementary Conformity Certificate covering an extensive range of product technologies: liquid formula, encapsulation, packaging, labeling, warehousing, mixing, dry formula, coating, and more — all audited and confirmed compliant.

NSF Certificate of Conformity

Print Date
March 02, 2026

Certification Number
C0040870-HSCDS-13

Initial Certification
March 19, 2021

Expiration Date
March 25, 2027

Signed on behalf of NSF International

David Trosin
Senior Director, Nutrition and Wellness

NSF International has assessed and confirmed compliance of

[redacted]

Scope: NSF/ANSI 455-2 - 2024
which includes 21CFR Part 111, 21 CFR Part 117, 21 CFR Part 11, 21 CFR Part 1.5 Subpart L & 21 CFR Part 1.9 Subpart O

Product Technologies:
Coating, Dry Formulation, Encapsulation, Liquid Formulation, Mixing, Packaging/Labeling Operation, Packaging/Labeling Operation - Bulk Packaging, Packaging/Labeling Operation - Dispensing, Packaging/Labeling Operation - Primary Packaging, Packaging/Labeling Operation - Secondary Packaging, Quality Unit Operations, Warehousing

Product Categories:
Ingestible Liquid, Ingestible Oil, Soft Gel

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PROCESS CERTIFICATION BODY

NSF International
789 N. Dixboro Road, Ann Arbor, MI 48105 USA

This certificate is the property of NSF International and must be returned upon request.

For the most current and complete information, please access NSF's website (nsf.org)

NSF
GMP CERTIFIED
NSF/ANSI 455-2
Dietary Supplements

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SQF Food Safety Code — Food Manufacturing Edition 9
(Eurofins Food Assurance — Certificate No: 52985)

This is the facility where our **Sunset** product is manufactured,the SQF (Safe Quality Food) certification is one of the most widely recognized food safety standards in global retail. Product scope: Dietary Supplements (FSC 31) — specifically liquid and powder collagen. Achieved through Unannounced Recertification, meaning production processes are maintained at audit-ready levels at all times.

eurofins | Food Assurance

Eurofins Food Assurance
2120 Rittenhouse Street, Suite A
Des Moines, IA 50321, USA
Ph: (515) 299-6979
www.eurofinsus.com/food-safety

DATES OF AUDIT:
06/13/2022 - 06/15/2022

NEXT RE-CERTIFICATION DATE:
07/10/2023

DATE OF DECISION:
07/11/2022

EXPIRATION DATE:
09/23/2023

CERTIFICATE NUMBER:
52985

CERTIFICATION TYPE:
Unannounced Recertification

Certificate of Registration

This acknowledges that

[redacted]

is registered as meeting the requirements for the

SQF Food Safety Code for Food Manufacturing, Edition 9

Registration schedule

Scope of registration [food sector categories and products]:

Food sector category: FSC 31: Dietary Supplements

Products: Liquid collagen & Powder collagen

Signature of issuing officer
Brian Neal
Technical Manager

Signature of authorized officer
Chuck Russo
Vice President

SQF INSTITUTE — One world. One standard.
JAS-ANZ

amare

Production/quality management certifications of the facility where our Nitro product is manufactured.

SQF Level 3 Certification

This certification represents the highest distinction attainable from SQF. The SQF Food Safety and Quality Program, formerly known as SQF Level 3 certification, includes all the criteria for levels 1 and 2 SQF certification, while adding defined parameters for achieving consistent quality food grade products.

Current Good Manufacturing Practice Regulations

CGMP refers to Current Good Manufacturing Practice regulations enforced by the FDA. Our CGMP designation assures you of our proper design, monitoring, and control of manufacturing processes and facilities.

NSF

GMP Regulatory Compliance – Good Manufacturing Practices (GMPs) and NSF/ANSI 455 provide a system of processes, procedures and documentation for the dietary supplement, personal care and OTC industries.

amare

Production/quality management certifications of the facility where our Nitro product is manufactured.

USDA Organic Certification

Our recognized organic certification is your verifiable assurance that we've used only produce grown on soil with no prohibited substances applied for three years before harvesting, as required by USDA organic regulations.

Non GMO

An important distinction and recognized industry standard, non GMO refers to a product produced without genetic engineering or ingredients derived from genetically modified organisms.

Non-GMO Project

Non-GMO Project Verified means that a product is compliant with the Non-GMO Project Standard, which includes stringent provisions for testing, traceability, and segregation. Non-GMO Project Standard is a consensus-based set of criteria crafted with the insight from dozens of industry experts, reflecting a dynamic range of perspectives.

amare

Production/quality management certifications of the facility where our Nitro product is manufactured.

Gluten Free

Gluten free has become a popular distinction for discerning consumers hoping to achieve improved health, weight loss, and increased energy. Our gluten free distinction lets the consumer know that your products are capable of meeting these strict demands.

Marine Stewardship Council

MSC meets international best practice guidelines and industry standards to ensure that you can trust seafood with the blue MSC label.

Kosher Certification

Kosher certification indicates that a product has been verified by a rabbinic agency to ensure all ingredients, derivatives, tools, and machinery have no trace of non-kosher substances.

amare

Production/quality management certifications of the facility where our Nitro product is manufactured.

OU Kosher

The OU (Orthodox Union) Kosher is the world's largest and most widely recognized international kosher certification agency. They certify over 1,000,000 products and hold the highest standard of kosher certification. The OU symbol is one of the world's best-known trademarks and offers high quality food standard assurance to customers.

Halal

The word, "halal" means "permissible" in Arabic. This is not the same as kosher. A halal certificate, issued by IFANCA (Islamic Food and Nutrition Council of America), guarantees that a food grade product meets the requirements of Islamic law and has been certified suitable for consumption. Unlike halal, kosher certified foods conform only to Jewish law, and not Islamic.

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FDA Drug Establishment Registration / FEI (iLABS Korea Co., Ltd.)
(FEI: 3020318393)

The facility where our **Sunrise, Energy+, Edge+, HL5, On Shots, and MentaBiotics** products are manufactured is also registered with the U.S. FDA as a drug manufacturing establishment. This indicates that the quality systems applied at the facility for supplement and cosmetic production are aligned with international pharmaceutical standards.

2024

CERTIFICATE OF COSMETIC FACILITY REGISTRATION

This certifies that:

iLABS KOREA Co., Ltd.
61 Hambangmoe-ro 401beon-gil, Namdong-gu
Incheon, Incheon 21638
South Korea

has submitted its facility registration statement for the cosmetic product facility identified below with the U.S. Food and Drug Administration pursuant to Chapter VI of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 361 et seq.) and such registration has been verified as currently effective on the date hereof by Registrar Corp.

NAME OF FACILITY: iLABS KOREA Co., Ltd.

FACILITY REGISTRATION NUMBER: 3029936534

This certificate affirms that the above company has filed its Facility Registration Statement for the referenced cosmetic product facility with the U.S. Food and Drug Administration pursuant to the Federal Food, Drug, and Cosmetic Act, as amended by the Modernization of Cosmetics Regulation Act of 2022, (21 U.S.C. 361 et seq.), such registration having been verified as effective by Registrar Corp as of the date hereof. Registrar Corp will confirm that such filing remains effective upon request and presentation of this certificate until December 31, 2024, unless such filing is 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. Registration of a cosmetic product facility or assignment of a registration number does not in any way denote approval by the Food and Drug Administration of the company or the facility. Any representation in labeling or advertising that creates an impression of official approval because of such filing, or such number could be considered misleading and misbranding.

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 a private registration agent not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
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info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: February 16, 2024

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FDA Cosmetic Facility Registration (iLABS Korea Co., Ltd.) (Registrar Corp)

The facility where our **Sunrise, Energy+, Edge+, HL5, On Shots, and MentaBiotics** products are manufactured, the cosmetic components of our products is registered with the U.S. Food and Drug Administration (FDA). This registration confirms that our manufacturing site operates to auditable standards under U.S. federal law.

FY2024

CERTIFICATE OF REGISTRATION

This certifies that:

iLABS KOREA Co., Ltd.
61 Hambangmoe-ro 401beon-gil, Namdong-gu
Incheon, Incheon 21638
South Korea

is registered with the U.S. Food and Drug Administration for the statutory filing period applicable to U.S. FY 2024 pursuant to part 207 of Title 21, U.S. Code of Federal Regulations.

DUNS Number:	**69-418-5577**
FEI:	**3020318393**
U.S. Agent/Registrant Contact:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

Filing was performed during the October 1 - December 31, 2023 statutory period, and renewal is not required until the next statutory period of October 1 - December 31, 2024. Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate, until the end of the year stated above, unless 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. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of registration or possession of registration number or NDC number is misleading and constitutes misbranding. 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.

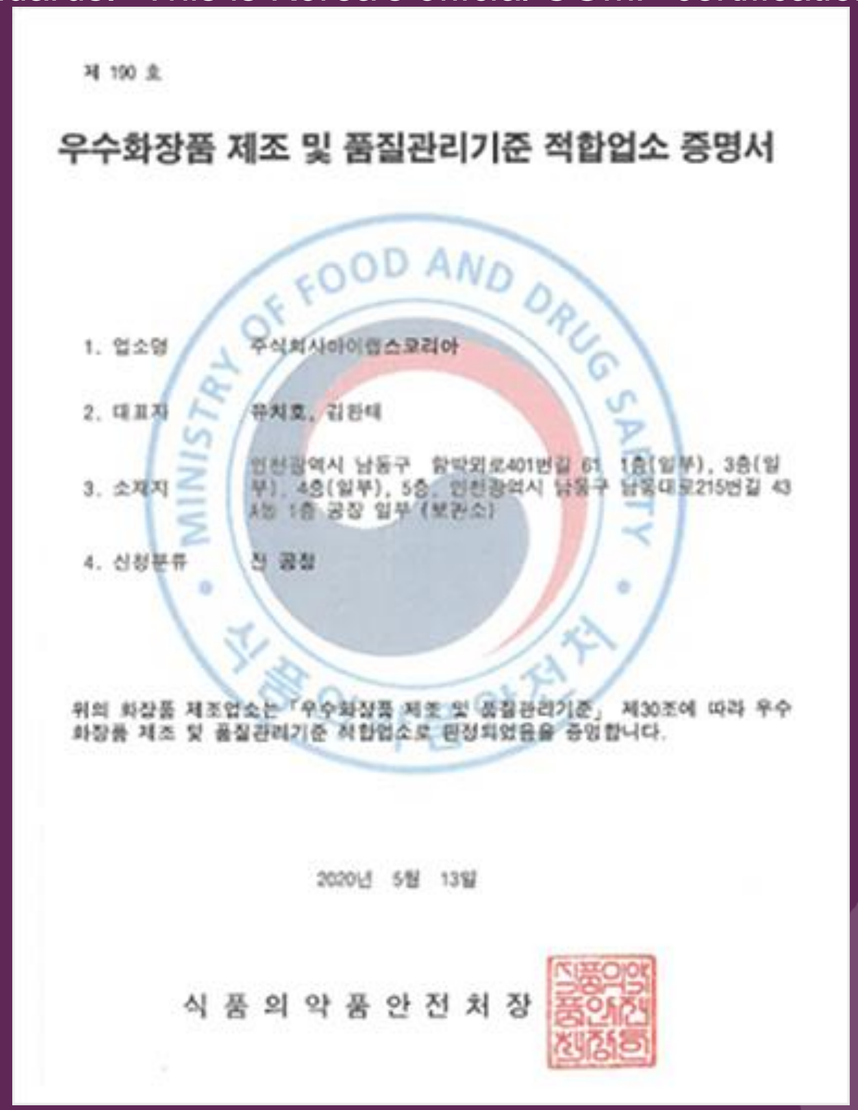
Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: August 5, 2024

amare

Korean MFDS CGMP Certificate — 우수화장품 제조 및 품질관리기준 (Ministry of Food and Drug Safety, Korea)

The facility where our **Sunrise, Energy+, Edge+, HL5, On Shots, and MentaBiotics** products are manufactured has been certified by the Korean Ministry of Food and Drug Safety as compliant with the "Superior Cosmetic Manufacturing and Quality Management Standards." This is Korea's official CGMP certification.



제 190 호

우수화장품 제조 및 품질관리기준 적합업소 증명서

1. 업소명 주식회사아이엠스코리아

2. 대표자 유치호, 김원태

3. 소재지 인천광역시 남동구 함박외로401번길 61, 1층(일부), 3층(일부), 4층(일부), 5층, 인천광역시 남동구 남동대로215번길 43 A동 1층 공장 일부 (보관소)

4. 신청분류 전 공정

위의 화장품 제조업소는 「우수화장품 제조 및 품질관리기준」 제30조에 따라 우수화장품 제조 및 품질관리기준 적합업소로 판정되었음을 증명합니다.

2020년 5월 13일

식 품 의 약 품 안 전 처 장

MINISTRY OF FOOD AND DRUG SAFETY

amare

ISO 14001:2015 — Environmental Management System (Kiwa)

The facility where our **Sunrise, Energy+, Edge+, HL5, On Shots, and MentaBiotics** products are manufactured, holds ISO 14001 certification, an internationally recognized standard for environmental management systems. This demonstrates a systematic approach to waste management, energy consumption, and minimizing environmental impact across production operations.

CERTIFICATE

kiwa

Registration no. KE 221809 IAF Code 12
First issue date 2017-05-07 Issue date 2023-06-08
Expiry date 2026-06-07 Release date 2023-06-23

Environmental Management System Certificate

KS I ISO 14001:2015/ISO 14001:2015

We certify that the Environmental Management System of the Organization:

(주)아이앱스 코리아

Is in compliance with the standard KS I ISO 14001:2015/ISO 14001:2015
for the following products/services:

화장품(스킨케어, 메이크업, 향수, 바디클렌징, 헤어제품)의 개발 및 제조

General Manager
Seung-Chan Lee

인증 사업장

IAF KAB

KAB-QC-21

amare

ISO 9001:2015 — Quality Management System (Kiwa)

The same manufacturer holds ISO 9001 quality management certification. This confirms that all processes — from product development through to manufacturing — are managed in accordance with internationally recognized quality standards.

CERTIFICATE

kiwa

Quality Management System Certificate

KS Q ISO 9001:2015/ISO 9001:2015

We certify that the Quality Management System of the Organization:

(주)아이랩스 코리아

is in compliance with the standard KS Q ISO 9001:2015/ISO 9001:2015
for the following products/services:

화장품(스킨케어, 메이크업, 향수, 바디클렌징, 헤어제품)의 개발 및 제조

General Manager
Seung-Chan Lee

amare

ISO 22716:2007 — Cosmetic GMP (CGMP)
(Kiwa — Certificate KC 22107)

ISO 22716 is the international Good Manufacturing Practice standard for cosmetics. This certificate confirms that good manufacturing practices are fully implemented across skincare, makeup, and hair product production.
It is a core requirement for European and global cosmetic markets.

Certificate

Certificate
KC 22107

kiwa

First issue 2019-08-27 Replaces 2025-08-04
Valid from 2025-08-27 Valid until 2028-08-26

Cosmetic-Good Manufacturing Practices Certificate

ISO 22716:2007

Kiwa has determined that the Cosmetic-Good Manufacturing Practices of

㈜아이랩스 코리아

complies with the standard ISO 22716:2007 for the scope

스킨케어, 페이셜 워시, 메이크업, 파운, 바디 클렌징 및 헤어제품의 제조

인증 사업장

General Manager
Seung-Cheon Lee

KC 22107

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