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CERTIFIED

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Amare Global ile ürünlerimizi üreten şirketler arasında imzalanan gizlilik anlaşması nedeniyle, bu şirketlerin isimleri açıklanmamıştır.

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UL Verification Services — GMP Denetim Sertifikası
(UL, Sertifika No: i10-34169)

Sunset ürünümüzün üretildiği tesise ait, UL (Underwriters Laboratories) tarafından düzenlenen bu denetim sertifikası, 21 CFR Part 111 kapsamında — ABD Takviye Edici Gıda cGMP yönetmeliğinin en güncel versiyonuna — tam uyum sağlandığını belgeler. Sıvı, toz, tablet, kapsül ve softgel formlarındaki ürünlerin üretimi, paketlenmesi ve depolanması denetim kapsamındadır.

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
Issue Number: 8
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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

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California Organic Processed Product Registration
(California Sağlık Bakanlığı, No: 20895)

Sunset ürünümüzün üretildiği tesise ait olan, California Organik Gıda ve Tarım Yasası kapsamında "organik" olarak işlenmiş ürün kaydına tabidir.
Bu kayıt, söz konusu ürünlerin organik etiketini yasal güvenceyle taşıyabildiğini belgeler.

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 (Takviye Gıda Üreticisi)
(FDA Kayıt No: 13618051326)

Sunset ürünümüzün üretildiği tesis, ABD Gıda Güvenliği Modernizasyon Yasası (FSMA) kapsamında FDA'ya kayıtlıdır.
Bu kayıt, gıda güvenliği sistemlerinin federal denetim standartlarını karşıladığını gösterir.

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

© GOES 961 LITHO IN U.S.A.

NSF/GRMA GMP Uygunluk Belgesi
(NSF International, Sertifika No: C0040870)

Bağımsız denetim kuruluşu NSF International tarafından düzenlenen bu belge,
Sunset ürünümüzün üretildiği tesisimizin NSF/ANSI 455-2 standardına —
yani Takviye Edici Gıdalar için İyi Üretim Uygulamaları'na — uygun olduğunu onaylar.
NSF, dünyada en güvenilir üçüncü taraf denetim kuruluşlarından biridir.

NSF

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 — Takviye Gıda GMP (NSF/ANSI 455-2:2021)

NSF'nin bir önceki belgeyi tamamlayan ikinci Uygunluk Sertifikası'dır. Ürün teknolojileri kapsamı oldukça geniştir: sıvı formül, kapsülleme, ambalajlama, depolama, kaplama ve kuru formül dahil tüm üretim aşamaları denetlenmiştir.

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 — Gıda Üretimi Ed. 9
(Eurofins Food Assurance, Sertifika No: 52985)

Sunset ürünümüzün üretildiği tesise ait SQF (Safe Quality Food) sertifikası, gıda güvenliği alanında küresel perakende sektörünün en yaygın kabul gören belgelerinden biridir. Ürün kategorisi: Takviye Edici Gıdalar (FSC 31) — özellikle sıvı ve toz kolajen. Denetimsiz yeniden sertifikalandırma (Unannounced Recertification) ile elde edilmiştir; bu da üretim süreçlerinin her an denetlenebilir düzeyde olduğu anlamına gelir.

eurofins | Food Assurance

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

DATES OF AUDIT:
07/17/2025 – 07/18/2025

NEXT RE-CERTIFICATION DATE:
07/10/2026

DATE OF DECISION:
07/29/2025

EXPIRATION DATE:
09/23/2026

CERTIFICATE NUMBER:
FSS202005_11

CERTIFICATION TYPE:
Announced Recertification

Certificate of Registration

This acknowledges that

[redacted]

is registered as meeting the requirements for the
SQF Food Safety Code for Dietary Supplements Manufacturing, Edition 9

Registration schedule

Scope of registration [food sector categories and products]:

Food sector category: FSC 31: Dietary Supplements Manufacturing

Products: Tablets, Capsules, Softgels, Liquids, and Powders

Signature of issuing officer
Brian Neal

SQF INSTITUTE
One world. One standard.
SQF Institute is a division of FMI.

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PRODUCT CERTIFICATION BODY
9166

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Nitro Ürünümüzün üretildiği tesise ait
üretim/kalite yönetimi sertifikaları

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.

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Nitro Ürünümüzün üretildiği tesise ait
üretim/kalite yönetimi sertifikaları

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.

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Nitro Ürünümüzün üretildiği tesise ait
üretim/kalite yönetimi sertifikaları

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.

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Nitro Ürünümüzün üretildiği tesise ait
üretim/kalite yönetimi sertifikaları

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.

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

Sunrise, Energy+, Edge+, HL5, On Shots, MentaBiotics ürünlerimizin kozmetik bileşenlerini üreten tesisimiz,
ABD Gıda ve İlaç İdaresi'ne (FDA) kayıtlıdır.
Bu kayıt, üretim tesisimizin ABD federal mevzuatı kapsamında denetlenebilir standartlarda faaliyet gösterdiğini belgeler.

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
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: February 16, 2024

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FDA Drug Establishment Registration / FEI (iLABS Korea Co., Ltd.)
(Registrar Corp)

Sunrise, Energy+, Edge+, HL5, On Shots, MentaBiotics ürünlerimizin üretildiği tesis, ABD FDA'ya ilaç üretim tesisi olarak da kayıtlıdır. (FEI: 3020318393).Bu, tesisin takviye edici gıda ve kozmetik üretiminde uyguladığı kalite sistemlerini uluslararası ilaç standartlarıyla uyumlu olduğuna işaret eder.

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

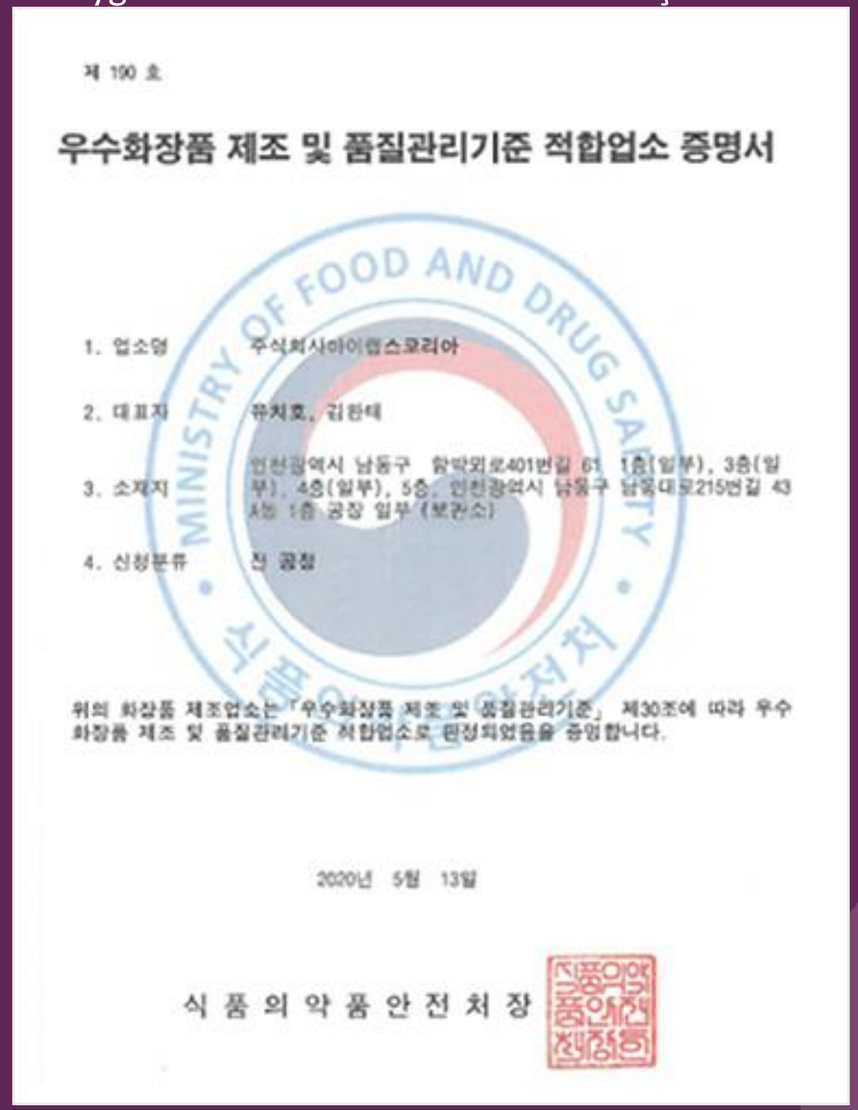
Filing was performed during the October 1 - December 31, 2023 statutory period, and renewal is not required until the next statutory period of October 1 - December 31, 2024. Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate, until the 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

Kore MFDS CGMP Sertifikası — 우수화장품 제조 및 품질관리기준
(Kore Gıda ve İlaç Güvenlik Bakanlığı)

Sunrise, Energy+, Edge+, HL5, On Shots, MentaBiotics ürünlerimizin üretildiği tesis,
Kore Gıda ve İlaç Güvenlik Bakanlığı tarafından "Üstün Kozmetik Üretim ve
Kalite Yönetimi Standartları"na uygun tesis olarak sertifikalandırılmıştır. Kore'nin resmi CGMP belgesidir.



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
식품의약품안전처

amare

ISO 14001:2015 — Çevre Yönetim Sistemi
(Kiwa)

Sunrise, Energy+, Edge+, HL5, On Shots, MentaBiotics ürünlerimizin üretildiği tesis, uluslararası geçerliliğe sahip ISO 14001 standardı kapsamında çevre yönetim sistemi sertifikasına sahiptir.
Bu belge; atık yönetimi, enerji tüketimi ve çevresel etki konularında sistematik bir yaklaşım uygulandığını gösterir.

CERTIFICATE

kiwa

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

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, 에이크업, 향수, 바디클렌징, 헤어제품)의 개발 및 제조

General Manager
Seung-Chan Lee

인증 사업장

IAF KAB

KAB-QC-21

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ISO 9001:2015 — Kalite Yönetim Sistemi
(Kiwa)

Aynı üretici, ISO 9001 kalite yönetim sistemi sertifikasına da sahiptir. Kozmetik ürünlerin geliştirilmesinden üretimine kadar tüm süreçlerin uluslararası kalite standartlarına göre yönetildiğini belgeler.


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

IAF

KAB

ISO 22716:2007 — Kozmetik GMP (CGMP)
(Kiwa)

Uluslararası kozmetik GMP standardı olan ISO 22716 sertifikası, cilt bakımı, makyaj ve saç ürünleri üretiminde iyi imalat uygulamalarının eksiksiz hayata geçirildiğini onaylar. Avrupa ve küresel kozmetik pazarlarında kabul gören temel bir belgedir.

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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