Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

LICENSE TO OPERATE

as

Cosmetic |Wholesaler

is hereby granted to

FRONTROW ENTERPRISE PHILS. INC.

G/F Pacific Century Tower, 1472-1476 Quezon Avenue, Brgy. South Triangle, Quezon City, Metro Manila

Owner:	**FRONTROW ENTERPRISE PHILS. INC.**
License Number:	**LTO-3000002878127**
Application Type:	**Renewal**
Validity of License:	**09 March 2024**

This LTO shall be renewed within **90 days** before its expiration, upon compliance with relevant laws, rules and regulations and the payment of fees. This LTO may be suspended, cancelled or revoked by this Office for cause if found violating RA 9711 and related issuances.

Furthermore, with this LTO, the FDA allows the establishment to apply for a market authorization [i.e. registration (CPR) or notification] for health products prior to manufacture, importation, sale or offer for sale, distribution, transfer, advertisement and/or promotion as the case may be.

BY AUTHORITY OF THE DIRECTOR GENERAL

ENGR. ANA TRINIDAD F. RIVERA, MSc
Director IV
Center for Cosmetics and Household/Urban Hazardous Substances Regulation and Research

DISPLAY IN PUBLIC VIEW

Additional information required under FDA Circular 2016-006 are reflected at the second page of this LTO

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

LICENSE TO OPERATE

as

Drug Importer|Wholesaler

is hereby granted to

FRONTROW ENTERPRISE PHILS. INC.

Ground Floor, Pacific Century Tower, 1472-1476, Quezon Avenue, South Triangle 4, Quezon City, Metro Manila

Owner:	**FRONTROW ENTERPRISE PHILS. INC.**
License Number:	**LTO-3000006828520**
Application Type:	**Variation**
Validity of License:	**12 September 2022**

This LTO shall be renewed within **90 days** before its expiration, upon compliance with relevant laws, rules and regulations and the payment of fees. This LTO may be suspended, cancelled or revoked by this Office for cause if found violating RA 9711 and related issuances.

Furthermore, with this LTO, the FDA allows the establishment to apply for a market authorization [i.e. registration (CPR) or notification] for health products prior to manufacture, importation, sale or offer for sale, distribution, transfer, advertisement and/or promotion as the case may be.

BY AUTHORITY OF THE DIRECTOR GENERAL

JESUSA JOYCE N. CIRUNAY, RPh
Director IV
Center for Drug Regulation and Research

DISPLAY IN PUBLIC VIEW

Additional information required under FDA Circular 2016-006 are reflected at the second page of this LTO

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

LICENSE TO OPERATE

as

Food Importer|Exporter|Wholesaler

is hereby granted to

FRONTROW ENTERPRISE PHILS. INC.

Ground Floor Pacific Century Tower #1472-1476 Quezon Avenue, Brgy. South Triangle, Quezon City, Metro Manila

Owner:	**FRONTROW ENTERPRISE PHILS. INC.**
License Number:	**LTO-3000007306632**
Application Type:	**Renewal**
Validity of License:	**21 April 2026**

This LTO shall be renewed within **90 days** before its expiration, upon compliance with relevant laws, rules and regulations and the payment of fees. This LTO may be suspended, cancelled or revoked by this Office for cause if found violating RA 9711 and related issuances.

Furthermore, with this LTO, the FDA allows the establishment to apply for a market authorization [i.e. registration (CPR) or notification] for health products prior to manufacture, importation, sale or offer for sale, distribution, transfer, advertisement and/or promotion as the case may be.

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV
Center for Food Regulation and Research

DISPLAY IN PUBLIC VIEW

Additional information required under FDA Circular 2016-006 are reflected at the second page of this LTO

LUQMAN LUCMAN BIN USMAN IMAM, MA (Islamic Studies)
Shari'ah Counselor-at-law
President

Address: **31-B A. Lake St., Brgy. Balong-Bato, San Juan City, Metro Manila Philippines** Contact #: **09183367111 / 09172400478 / 02-5320126** Bank Account: **3883-0661-88** Bank of the Philippine Islands, N. Domingo St., San Juan City, Metro Manila
Facebook Page: **facebook.com/philcongressorginc**

منظمة مؤتمر العلماء بالفلبين
PHILIPPINE ULAMA CONGRESS ORGANIZATION, INC.
SEC. REG. NO. CN200520263
Excellence. Partnership. Service.

شهادة الحلال

HALAL CERTIFICATE

Control # 03/4FP- 05062022

TO WHOM THESE PRESENTS MAY COME, GREETINGS:

After the thorough inspection, study and deliberation of the products distributed by FRONTROW ENTERPRISE PHILIPPINES INC., with office address at Unit 105, G/F Pacific Tower 1472-1576, Quezon Avenue, Brgy. South Triangle, Quezon City. The Philippine Ulama Congress Organization, Inc. DO HEREBY CERTIFY that the following products are "Halal" in accordance with Islamic Law and Halal Standards, to wit:

FOOD SUPPLEMENTS

1. **LUXXE WHITE**
2. **LUXXE SLIM**
3. **LUXXE RENEW**
4. **LUXXE PROTECT**

xxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxx

This certification is issued in compliance with Halal Certification requirements.

This Halal Certificate is renewable every year provided that the raw materials identified and used in these products remain the same, and the Halal Assurance System is duly implemented. Violation hereof, shall render this certificate null and void.

This Halal Certificate is issued on the 5th day of June 2022 and valid until the 4th day of June 2023.

LUQMAN L. BIN USMAN IMAM, MA (Islamic Studies)
Shari'ah Counselor-at-law
President, PUCOI

Member, International Halal Integrity Alliance (IHIA)& Recognized by Thailand
Address: **31-B A. Lake St., Brgy. Balong-Bato, San Juan City, Metro Manila Philippines** Contact #: **+639057298166/+6329845323/+6325320126**
Bank Account: **3883-0661-88** Bank of the Philippine Islands, N. Domingo St., San Juan City, Metro Manila
Facebook Page: **facebook.com/philcongressorginc**
Email Address: **pucoinc@gmail.com**

منظمة مؤتمر العلماء بالفلبين
PHILIPPINE ULAMA CONGRESS ORGANIZATION, INC.
SEC. REG. NO. CN200520263
Excellence. Partnership. Service.

شهادة الحلال

HALAL CERTIFICATE

Control # 04/3NFP- 08082022

TO WHOM THESE PRESENTS MAY COME, GREETINGS:

After the thorough inspection, study and deliberation of the products distributed by FRONTROW ENTERPRISE PHILIPPINES INC., with office address at Unit 105, G/F Pacific Tower 1472-1576, Quezon Avenue, Brgy. South Triangle, Quezon City. The Philippine Ulama Congress Organization, Inc. DO HEREBY CERTIFY that the following products are "Halal" in accordance with Islamic Law and Halal Standards, to wit:

COSMETICS

1. FRONTROW SOAP 01 SKIN WHITENING BAR
2. FRONTROW SOAP 02 OATMEAL EXFOLIANT BAR
3. FRONTROW SOAP 03 ORANGE PAPAYA BAR

xxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxxx

This certification is issued in compliance with Halal Certification requirements.

This Halal Certificate is renewable every year provided that the raw materials identified and used in these products remain the same, and the Halal Assurance System is duly implemented. Violation hereof, shall render this certificate null and void.

This Halal Certificate is issued on the 8th day of August 2022 and valid until the 7th day of August 2023.

LUQMAN L. BIN USMAN IMAM, MA (Islamic Studies)
Shari'ah Counselor-at-law
President, PUCOI

Address: 31-B A. Lake St., Brgy. Balong-Bato, San Juan City, Metro Manila Philippines Contact #: +639279258482/+6329845323/+6329845332
Bank Account: 3883-0661-88 Bank of the Philippine Islands, N. Domingo St., San Juan City, Metro Manila
Facebook Page: facebook.com/philcongressorginc
Email Address: pucoinc@gmail.com

REPUBLIC OF THE PHILIPPINES
SECURITIES AND EXCHANGE COMMISSION
Ground Floor, Secretariat Building, PICC
City of Pasay, Metro Manila

COMPANY REG. NO. CS200903020

CERTIFICATE OF FILING
OF
AMENDED ARTICLES OF INCORPORATION

KNOW ALL PERSONS BY THESE PRESENTS:

THIS IS TO CERTIFY that the amended articles of incorporation of the

FRONTROW ENTERPRISE PHILS. INC.
(Amending Article VII thereof)

copy annexed, adopted on October 22, 2018 by a majority vote of the Board of Directors and by the vote of the stockholders owning or representing at least two-thirds of the outstanding capital stock, and certified under oath by the Secretary and a majority of the Board of Directors of the corporation was approved by the Commission on this date pursuant to the provision of Section 15 of the Revised Corporation Code of the Philippines, Republic Act No. 11232, which took effect on February 23, 2019 and copies thereof are filed with the Commission.

Unless this corporation obtains or already has obtained the appropriate Secondary License from this Commission, this Certificate does not authorize it to undertake business activities requiring a Secondary License from this Commission such as, but not limited to acting as: broker or dealer in securities, government securities eligible dealer (GSED), investment adviser of an investment company, close-end or open-end investment company, investment house, transfer agent, commodity/financial futures exchange/broker/merchant, financing company, pre-need plan issuer, general agent in pre-need plans and time shares/club shares/membership certificates issuers or selling agents thereof. Neither does this Certificate constitute as permit to undertake activities for which other government agencies require a license or permit.

IN WITNESS WHEREOF, I have hereunto set my hand and caused the seal of this Commission to be affixed to this Certificate at Pasay City, Metro Manila, Philippines, this 27th day of May, Twenty Nineteen.

GERARDO F. DEL ROSARIO
Director
Company Registration and Monitoring Department

AV/ioo

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

FDA Registration No.: FR-4000002546322

CERTIFICATE OF PRODUCT REGISTRATION
(High Risk Food Product)

Pursuant to the provisions of Republic Act No.3720, otherwise known as the Food, Drugs and Devices, and Cosmetic Act, as amended by Executive Order No. 175, and Republic Act No. 9711, otherwise known as the Food and Drug Administration Act of 2009, and other applicable laws, rules and regulations, the registration of the High Risk Food Product described hereunder is granted approval.

Product Name:	ENHANCED GLUTATHIONE (L-GLUTAMIC ACID,GLYCINE,L-CYSTEINE,NAC) FOOD SUPPLEMENT CAPSULE (Registered as Food Supplement with NO APPROVED THERAPEUTIC CLAIMS)
Brand Name:	LUXXE WHITE
Packaging:	WHITE OPAQUE HDPE BOTTLES (PACKED BY 60'S PER BOTTLE)
Company Name:	FRONTROW ENTERPRISE PHILS.INC.
Company Address:	UNIT 105,GROUND FLOOR,PACIFIC CENTURY TOWER,#1472-1476,QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON CITY
Company LTO:	CFRR-NCR-FW-4212
Manufacturer Name and Address:	AIE PHARMACEUTICALS,INC., 1845 S. VINEYARD AVE.#5 ONTARIO CA 91761

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each batch/lot of the product continues to meet all the legal requirements, and conforms to all the standards and specification of the product declared to the FDA, including compliance to the list of obligations enumerated at the reverse side of this document.

The authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

Issued on **19 February 2019** and valid until **19 February 2024** .

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV, Center for Food Regulation and Research

Remarks: Declare the complete list of Ingredients separate from Nutrition Information on the label.

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

FDA Registration No.: FR-4000002545286

CERTIFICATE OF PRODUCT REGISTRATION
(High Risk Food Product)

Pursuant to the provisions of Republic Act No.3720, otherwise known as the Food, Drugs and Devices, and Cosmetic Act, as amended by Executive Order No. 175, and Republic Act No. 9711, otherwise known as the Food and Drug Administration Act of 2009, and other applicable laws, rules and regulations, the registration of the High Risk Food Product described hereunder is granted approval.

Product Name:	BERRY EXTRACT HERBAL DIETARY SUPPLEMENT CAPSULE (Registered as Food Supplement with NO APPROVED THERAPEUTIC CLAIMS)
Brand Name:	LUXXE RENEW
Packaging:	HDPE WHITE OPAQUE BOTTLE
Company Name:	FRONTROW ENTERPRISE PHILS. INC.
Company Address:	UNIT 105,GROUND FLOOR,PACIFIC CENTURY TOWER, #1472-1476,QUEZON AVENUE, BRGY. SOUTH TRIANGLE,QUEZON CITY
Company LTO:	CFRR-NCR-FW-4212
Manufacturer Name and Address:	AIE,PHARMACEUTICALS,INC., 1845 VINEYARD AVENUE,SUITE #5 ONTARIO, CALIFORNIA U.S.A

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each batch/lot of the product continues to meet all the legal requirements, and conforms to all the standards and specification of the product declared to the FDA, including compliance to the list of obligations enumerated at the reverse side of this document.

The authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

Issued on **17 April 2019** and valid until **17 April 2024** .

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV, Center for Food Regulation and Research

Remarks: Declare nutrition information separately from ingredients on the label. Delete "Active" from ingredients.

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

FDA Registration No.: FR-4000002546191

CERTIFICATE OF PRODUCT REGISTRATION
(High Risk Food Product)

Pursuant to the provisions of Republic Act No.3720, otherwise known as the Food, Drugs and Devices, and Cosmetic Act, as amended by Executive Order No. 175, and Republic Act No. 9711, otherwise known as the Food and Drug Administration Act of 2009, and other applicable laws, rules and regulations, the registration of the High Risk Food Product described hereunder is granted approval.

Product Name:	(L-CARNITINE + GREEN TEA EXTRACT + ACAI BERRY EXTRACT) FOOD SUPPLEMENT CAPSULE (Registered as Food Supplement with NO APPROVED THERAPEUTIC CLAIMS)
Brand Name:	LUXXE SLIM
Packaging:	HDPE WHITE OPAQUE BOTTLE BY 60s CAPSULES
Company Name:	FRONTROW ENTERPRISE PHILS.INC.
Company Address:	UNIT 105,GROUND FLOOR,PACIFIC CENTURY TOWER,#1472-1476,QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON CITY
Company LTO:	CFRR-NCR-FW-4212
Manufacturer Name and Address:	AIE PHARMACEUTICALS,INC., 1845 S. VINEYARD AVE.#5 ONTARIO CA 91761

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each batch/lot of the product continues to meet all the legal requirements, and conforms to all the standards and specification of the product declared to the FDA, including compliance to the list of obligations enumerated at the reverse side of this document.

The authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

Issued on **16 August 2018** and valid until **28 September 2023** .

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV, Center for Food Regulation and Research

Remarks: LUXXE SLIM is approved as brand name and it should not be presented nor advertised as a claim.

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

FDA Registration No.: FR-4000007148493

CERTIFICATE OF PRODUCT REGISTRATION
(High Risk Food Product)

Pursuant to the provisions of Republic Act No.3720, otherwise known as the Food, Drugs and Devices, and Cosmetic Act, as amended by Executive Order No. 175, and Republic Act No. 9711, otherwise known as the Food and Drug Administration Act of 2009, and other applicable laws, rules and regulations, the registration of the High Risk Food Product described hereunder is granted approval.

Product Name:	HERBAL EXTRACTS WITH VITAMIN C, D AND ZINC FOOD SUPPLEMENT CAPSULE (Registered as Food Supplement with NO APPROVED THERAPEUTIC CLAIMS)
Brand Name:	LUXXE IMMUNPLUS
Packaging:	HDPE PLASTIC BOTTLE OF 60'S CAPSULE
Company Name:	FRONTROW ENTERPRISE PHILS. INC.
Company Address:	G/F PACIFIC CENTURY TOWER, 1472-1476 QUEZON AVENUE, BRGY. SOUTH TRIANGLE, QUEZON CITY, METRO MANILA
Company LTO:	LTO3000006069581
Manufacturer Name and Address:	HOMART PHARMAÇEUTICALS PTY LTD, 59 KIRBY ST. RAYDALMERE, NEW SOUTH WALES, 2116 AUSTRALIA

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each batch/lot of the product continues to meet all the legal requirements, and conforms to all the standards and specification of the product declared to the FDA, including compliance to the list of obligations enumerated at the reverse side of this document.

The authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

Issued on **29 December 2020** and valid until **29 December 2024** .

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV, Center for Food Regulation and Research

Remarks: Declare the approved product name prominently on the PDP, complete list of ingredients (separate from nutrition facts table) including component of the capsule used headed by the term "Ingredients", prescribed format of nutrition information with correct amount and % RENI equivalent of nutrients, precaution "For Adults use only. Not intended for children, pregnant and lactating women." and the phrase "NO APPROVED THERAPEUTIC CLAIMS" as per BC 2 s. 1999 on the label. Delete "Indications". To submit valid LTO as Food Importer once issued.

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

FDA Registration No.: FR-4000002223609

CERTIFICATE OF PRODUCT REGISTRATION

(Medium Risk Food Product)

Pursuant to the provisions of Republic Act No.3720, otherwise known as the Food, Drugs and Devices, and Cosmetic Act, as amended by Executive Order No. 175, and Republic Act No. 9711, otherwise known as the Food and Drug Administration Act of 2009, and other applicable laws, rules and regulations, the registration of the Medium Risk Food Product described hereunder is granted approval.

Product Name:	GRAPESEED EXTRACT OIL FOOD SUPPLEMENT CAPSULE (Registered as Food Supplement with NO APPROVED THERAPEUTIC CLAIMS)
Brand Name:	LUXXE PROTECT
Packaging:	HDPE WHITE OPAQUE PLASTIC BOTTLE OF 30'S AND 60'S
Company Name:	FRONTROW ENTERPRISE PHILS.INC.
Company Address:	UNIT 105,GROUND FLOOR,PACIFIC CENTURY TOWER,#1472-1476,QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON CITY
Company LTO:	CFRR-NCR-FW-4212
Manufacturer Name and Address:	AIE PHARMACEUTICALS,INC., 1845 S. VINEYARD AVE.#5 ONTARIO CA 91761

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each batch/lot of the product continues to meet all the legal requirements, and conforms to all the standards and specification of the product declared to the FDA, including compliance to the list of obligations enumerated at the reverse side of this document.

The authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

Issued on **21 April 2018** and valid until **21 April 2023** .

BY AUTHORITY OF THE DIRECTOR GENERAL

PILAR MARILYN M. PAGAYUNAN
Director IV, Center for Food Regulation and Research

Remarks: Declare the "Ingredients" followed by complete list of ingredients (delete the term "Other Ingredients" and separate list of ingredients from the Nutrition Facts), function of food additives used, nutrition information in prescribed format and expiry/expiration date/use-by-date/consume before date in prescribed format on the label.

Republic of the Philippines
Department of Health
Food And Drug Administration
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

Notification of Cosmetic Product

Pursuant to the existing rules and regulations of the Food and Drug Administration (FDA) for the notification of cosmetics, the product notification is hereby acknowledged,

Cosmetic Notification Number:	**NN-1000004323861**
Validity:	06 April 2025

Particulars of the Product

Brand Name:	FRONTROW
Product Name:	LUXXE WHITE REVEAL DYNAMIC DUO BB+CC HYBRID STICK
Variant/s:	01, 02, 03

Particulars of the Manufacturer

Name of the Manufacturer:	HWA SUNG COSMETICS
Address of the Manufacturer:	#161-5 DODANG-DONG WONMI-KU BUCHEN-SI KYUNGKI-DO KOREA 420-130
Country of Manufacture:	Korea, Republic of

Particulars of the Local Company Responsible for Placing the Product in the Market

Name of the Company:	FRONTROW ENTERPRISE PHILS., INC.
Address of the Company:	G/F PACIFIC CENTURY TOWER, 1472-1476 QUEZON AVE, BRGY. SOUTH TRIANGLE QUEZON CITY
LTO Number:	LTO-3000002878127

The product is allowed to be sold in the local market subject to compliance with the requirements of the ASEAN Cosmetic Directive and the FDA laws, and its implementing rules and regulations. This notification should not be taken as a guarantee and/or endorsement of the safety, quality, and claimed benefit of the product. Any subsequent changes to the information submitted in this notification will render this notification invalid and a new notification will have to be submitted.

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each consignment of the product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product declared to the FDA.

This authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

BY AUTHORITY OF THE DIRECTOR GENERAL

Engr. Ana Trinidad F. Rivera, MSc
Director IV, Center for Cosmetic Regulation and Research

Republic of the Philippines
Department of Health
Food And Drug Administration
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

Notification of Cosmetic Product

Pursuant to the existing rules and regulations of the Food and Drug Administration (FDA) for the notification of cosmetics, the product notification is hereby acknowledged,

Cosmetic Notification Number:	**NN-1000004318588**
Validity:	06 April 2025

Particulars of the Product

Brand Name:	FRONTROW
Product Name:	LUXXE WHITE REVEAL INSTABRIGHT BODY CREME SPF25
Variant/s:	NOT APPLICABLE

Particulars of the Manufacturer

Name of the Manufacturer:	SEOUL COSMETICS CO., LTD.
Address of the Manufacturer:	(143B-9L) NAMDONG IND, 718-8, GOJAN-DONG, NAMDONG-GU, INCHEON
Country of Manufacture:	Korea, Republic of

Particulars of the Local Company Responsible for Placing the Product in the Market

Name of the Company:	FRONTROW ENTERPRISE PHILS., INC.
Address of the Company:	G/F PACIFIC CENTURY TOWER, 1472-1476 QUEZON AVENUE, BRGY. SOUTH TRIANGLE, QUEZON CITY
LTO Number:	LTO-3000002878127

The product is allowed to be sold in the local market subject to compliance with the requirements of the ASEAN Cosmetic Directive and the FDA laws, and its implementing rules and regulations. This notification should not be taken as a guarantee and/or endorsement of the safety, quality, and claimed benefit of the product. Any subsequent changes to the information submitted in this notification will render this notification invalid and a new notification will have to be submitted.

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each consignment of the product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product declared to the FDA.

This authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

BY AUTHORITY OF THE DIRECTOR GENERAL

Engr. Ana Trinidad F. Rivera, MSc
Director IV, Center for Cosmetic Regulation and Research

Republic of the Philippines
Department of Health
Food And Drug Administration
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

Notification of Cosmetic Product

Pursuant to the existing rules and regulations of the Food and Drug Administration (FDA) for the notification of cosmetics, the product notification is hereby acknowledged,

Cosmetic Notification Number:	**NN-1000006388239**
Validity:	15 June 2023

Particulars of the Product

Brand Name:	FRONTROW 01
Product Name:	FRONTROW 01 SKIN WHITENING BAR WITH GLUTATHIONE +SKIN VITAMINS+KOJIC ACID
Variant/s:	NOT APPLICABLE

Particulars of the Manufacturer

Name of the Manufacturer:	CDJ SOAP MANUFACTURING
Address of the Manufacturer:	ZAPANTA COMPOUND,ROYAL SUBDIVISION,SAN JUAN,TAYTAY RIZAL
Country of Manufacture:	Philippines

Particulars of the Local Company Responsible for Placing the Product in the Market

Name of the Company:	FRONTROW ENTERPRISE PHILS.INC.
Address of the Company:	G/F PAIFIC CENTURY TOWER,1472-1476 QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON ITY
LTO Number:	3000002878127

The product is allowed to be sold in the local market subject to compliance with the requirements of the ASEAN Cosmetic Directive and the FDA laws, and its implementing rules and regulations. This notification should not be taken as a guarantee and/or endorsement of the safety, quality, and claimed benefit of the product. Any subsequent changes to the information submitted in this notification will render this notification invalid and a new notification will have to be submitted.

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each consignment of the product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product declared to the FDA.

This authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

BY AUTHORITY OF THE ACTING DIRECTOR GENERAL

Engr. Ana Trinidad F. Rivera, MSc
Director IV, Center for Cosmetic Regulation and Research

Republic of the Philippines
Department of Health
Food And Drug Administration
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

Notification of Cosmetic Product

Pursuant to the existing rules and regulations of the Food and Drug Administration (FDA) for the notification of cosmetics, the product notification is hereby acknowledged,

Cosmetic Notification Number:	**NN-1000006388271**
Validity:	10 June 2023

Particulars of the Product

Brand Name:	FRONTROW 02
Product Name:	FRONTROW 02 OATMEAL EXFOLIANT BAR
Variant/s:	NOT APPLICABLE

Particulars of the Manufacturer

Name of the Manufacturer:	CDJ SOAP MANUFACTURING
Address of the Manufacturer:	ZAPANTA COMPOUND,ROYAL SUBDIVISION,SAN JUAN,TAYTAY,RIZAL
Country of Manufacture:	Philippines

Particulars of the Local Company Responsible for Placing the Product in the Market

Name of the Company:	FRONTROW ENTERPRISE PHILS.INC.
Address of the Company:	G/F PACIFIC CENTURY TOWER,1472-1476 QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON CITY
LTO Number:	3000002878127

The product is allowed to be sold in the local market subject to compliance with the requirements of the ASEAN Cosmetic Directive and the FDA laws, and its implementing rules and regulations. This notification should not be taken as a guarantee and/or endorsement of the safety, quality, and claimed benefit of the product. Any subsequent changes to the information submitted in this notification will render this notification invalid and a new notification will have to be submitted.

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each consignment of the product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product declared to the FDA.

This authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

BY AUTHORITY OF THE ACTING DIRECTOR GENERAL

Engr. Ana Trinidad F. Rivera, MSc
Director IV, Center for Cosmetic Regulation and Research

Republic of the Philippines
Department of Health
Food And Drug Administration
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

Notification of Cosmetic Product

Pursuant to the existing rules and regulations of the Food and Drug Administration (FDA) for the notification of cosmetics, the product notification is hereby acknowledged,

Cosmetic Notification Number:	**NN-1000006388213**
Validity:	15 June 2023

Particulars of the Product

Brand Name:	FRONTROW 03
Product Name:	FRONTROW 03 ORANGE PAPAYA BAR WITH GLUTATHIONE+SKIN VITAMINS+KOJIC ACID
Variant/s:	NOT APPLICABLE

Particulars of the Manufacturer

Name of the Manufacturer:	CDJ SOAP MANUFACTURING
Address of the Manufacturer:	ZAPANTA COMPOUND,ROYAL SUBDIVISION,SAN JUAN,TAYTAY,RIZAL
Country of Manufacture:	Philippines

Particulars of the Local Company Responsible for Placing the Product in the Market

Name of the Company:	FRONTROW ENTERPRISE PHILS.INC.
Address of the Company:	G/F PACIFIC CENTURY TOWER,1472-1476 QUEZON AVENUE,BRGY.SOUTH TRIANGLE,QUEZON ITY
LTO Number:	3000002878127

The product is allowed to be sold in the local market subject to compliance with the requirements of the ASEAN Cosmetic Directive and the FDA laws, and its implementing rules and regulations. This notification should not be taken as a guarantee and/or endorsement of the safety, quality, and claimed benefit of the product. Any subsequent changes to the information submitted in this notification will render this notification invalid and a new notification will have to be submitted.

The company hereby ensures that they shall respond and cooperate fully with the FDA with regard to any subsequent post-marketing activity initiated by the FDA. Further, the company shall be responsible for ensuring that each consignment of the product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product declared to the FDA.

This authorization is subject to suspension, cancellation, or recall should any violation of FDA laws, and its implementing rules and regulations, involving the product be committed.

BY AUTHORITY OF THE ACTING DIRECTOR GENERAL

Engr. Ana Trinidad F. Rivera, MSc
Director IV, Center for Cosmetic Regulation and Research

AIE PHARMACEUTICALS, INC.

CERTIFICATE OF ANALYSIS

PRODUCT NAME:	**Luxxe White - Enhanced Glutathione**	**PRODUCT CODE:**	**AIE#10254**
BATCH SIZE:	**2,400,000**	**LOT NUMBER:**	**UL101415**
MANUFACTURED DATE:	**06/2018**	**DATE OF EXPIRATION:**	**05/2021**
DELIVERY SYSTEM:	**Capsules**	**Size: 00**	**Color: White/White**

TEST	SPECIFICATION	RESULT
APPEARANCE:	**Off White Powder**	**CONFORM**
TYPE OF ANAYLSIS:	**USP27**	**CONFORM**
WEIGHT:	**925 mg**	**THEORETICALS**
DISINTEGRATION:	***USP XXIII / USP 2040**	**PASS**

* USP XXIII disintegration guidelines: 30 minutes for capsules and uncoated tablets, 45 minutes for coated tablets.

CAPSULE INGREDIENTS ANALYSIS

TEST	SPECIFICATION	RESULT
Each Capsule contains:		
L-Glutamic Acid	180.0 mg	180.0 mg
L- Glycine	200.0 mg	200.0 mg
L-Cysteine	120.0 mg	120.0 mg
NAC	100.0 mg	100.0 mg
Vitamin C (Ascorbic Acid)	100.0 mg	100.0 mg
Alpha Lipoic Acid	50.0 mg	50.0 mg
Grapeseed Extract	25.0 mg	25.0 mg

Inactive Ingredients:
Magnesium Stearate, Silicon Dioxide, Gelatin and Water.

MICROBIAL TESTING / USP FULL MICROS

TEST	SPECIFICATION	RESULT
TOTAL BACTERIAL COUNT:	≤10,000 cfu/g	<10,000 cfu/g
SALMONELLA SPECIES:	ABSENT	ABSENT
ESCHERICHIA COLI:	ABSENT	ABSENT
STAPHYLOCOCCUS AUREUS:	ABSENT	ABSENT
YEAST AND MOLDS:	≤1,000 cfu/g	<1,000 cfu/g

HEAVY METAL TESTING:

TEST	SPECIFICATION	RESULT
HEAVY METALS:	≤ 10 ppm	< 10 ppm
ARSENIC:	≤ 3 ppm	< 1 ppm
CADMIUM:	≤ 1 ppm	< 1 ppm
MERCURY:	≤ 1 ppm	< 1 ppm
LEAD:	≤ 5 ppm	< 5 ppm

QC Manager: _____ Date: 11/25/19

1845 S. VINEYARD AVENUE, SUITE 5, ONTARIO, CALIFORNIA 91761, USA.
TEL: 909.947.9898 / FAX: 909.947.9813

I undertake to respond to and cooperate fully with Food and Drug Administration (hereafter referred to as "THE AUTHORITY") with regard to any subsequent post-marketing activity initiated by the authority.

I undertake to ensure that the product's technical and safety information is made readily available to the authority concerned and to keep records of the distribution of the products for product recall purposes, and other purposes as provided in existing laws, rules and regulations.

I undertake to notify the Authority of any adverse event, fatal or life threatening serious adverse event as soon as possible by telephone, facsimile transmission, email or in writing, and in any case, no later than 48 hours after first knowledge.

I undertake to act immediately on potential food safety concerns should my product source or origin declare/announce/notify a product recall order or any actions taken involving safety issues, and I am responsible to stop distribution or remove product from the outlets or take appropriate actions and inform the Authority on risk management actions undertaken and/or to be undertaken.

I declare that the particulars given in this product registration are true, all data, and information of relevance in relation to the registration have been supplied and that the documents enclosed are authentic or true copies.,

I understand that I shall be responsible for ensuring that each consignment of my product continues to meet all the legal requirements, and conforms to all the standards and specifications of the product that I have declared to the Authority.

I understand that I cannot place reliance on the acceptance of my product registration by the authority in any legal proceedings concerning my product, in the event that my product has failed to conform to any of the standards or specifications that I had previously declared to the Authority.

I understand that I will need to comply with all the labeling requirements as stipulated by Administrative Order No. 2014-030 and other pertinent laws and regulations associated with labeling.

I undertake to declare truthful product information and shall not cause the dissemination of any false, deceptive or misleading advertisement by print, radio, television, outdoor advertisement or other medium for the purpose of inducing or which is likely to induce directly or indirectly the purchase of the product.

I understand that any change or variation in the formulation of registered product will require new registration to the Authority and the subject shall be treated as new product.

I hereby agree and affirm full responsibility for the safety of my product/s and agree to indemnify and/or hold FDA free and harmless against any issues that may arise in the manufacture, import, export, distribute, transfer, promote, advertise, sponsor, sell, offer for sale, and where appropriate the use and testing, and marketing of my food product/s