

Norweijann "Owee" Cruz
PRESIDENT INFINITE VIDA GLOBAL CORP.

President Norweijann “Owee” Cruz brings a wealth of expertise in marketing, branding, and strategic business development, playing a pivotal role in shaping the vision and direction of Infinite Vida.

An alumna of the University of the Philippines, she has over 20 years of experience as a Marketing and PR Professional and is an Internationally Certified Color Analyst.

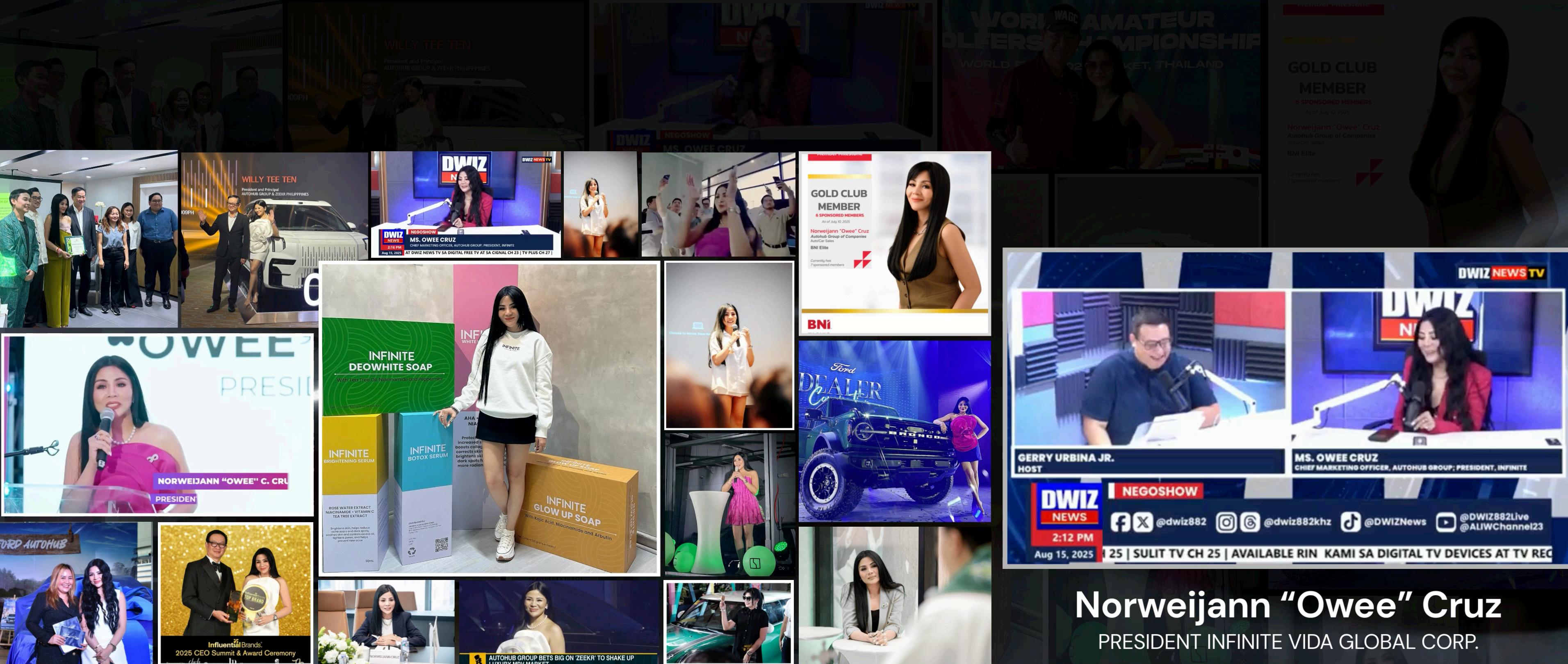
She is a recognized leader in the business community, serving as BNI Elite’s Network Leader and Gold Club Member. Her entrepreneurial and leadership roles include:

- Managing Director, ODV Creative Media Inc.
- President, Two Scents International Corporation
- General Manager, GAC’s Flagship Store
- Chief Marketing Officer & Head of Fleet Sales, Autohub Group of Companies — the exclusive Philippine distributor of Rolls-Royce, MINI Cooper, Lotus, Zeekr, Vespa, Triumph, Segway, and a major dealer of Ford, Nissan, Mitsubishi, Suzuki, Geely, Kia, GAC, among others.



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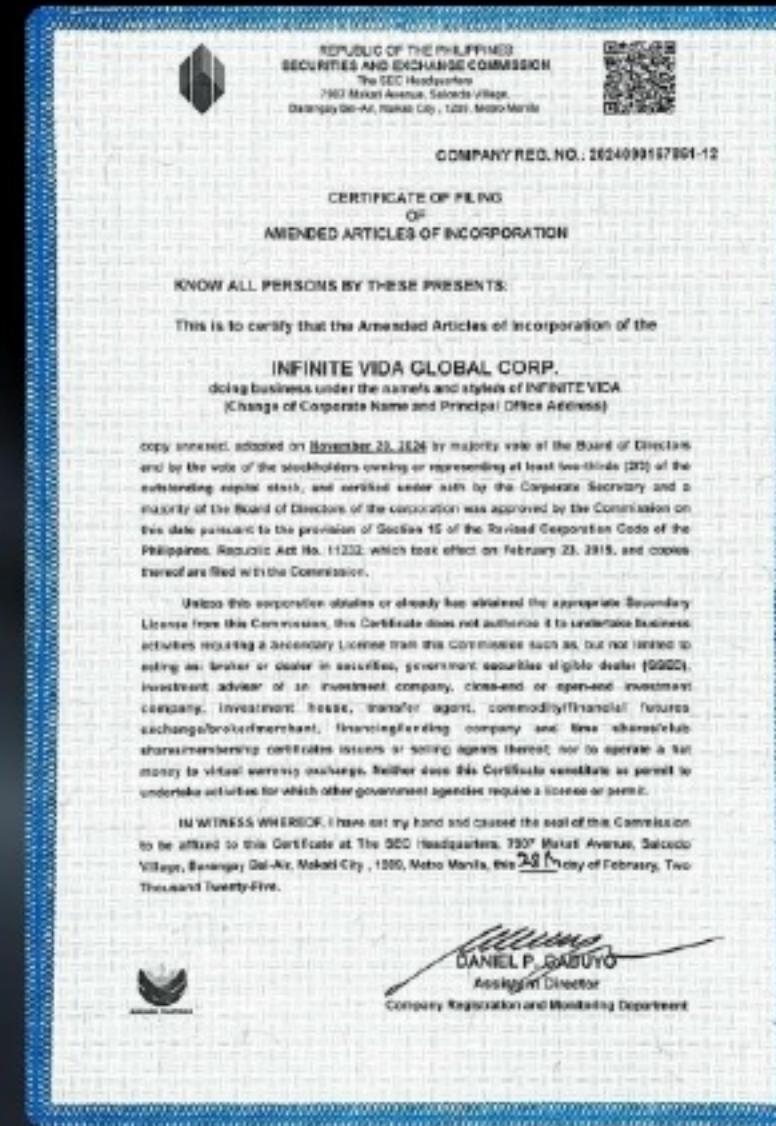
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THE BACKBONE OF OUR BUSINESS

**SEC REGISTERED
SEC REG. NO. 201309617861-12
PROOF WE ARE A LEGALLY RECOGNIZED CORPORATION.**

TIN REGISTERED
TIN No. 010-849-466-00000
WE PAY TAXES AND OPERATE WITH FULL TRANSPARENCY.

WE PAY TAXES AND OPERATE WITH FULL TRANSPARENCY.

**BUSINESS PERMIT
PERMIT No. 15266**

AUTHORIZED BY THE LOCAL GOVERNMENT TO OPERATE LEGALLY.

FDA LICENSED (FOOD & COSMETICS)
CFRR-NCR-FT-1186508 | LTO-3000014251701
YOUR HEALTH AND SAFETY ARE NON-NEGOTIABLE — OUR PRODUCTS ARE FDA-APPROVED.

 	Bureau of the Prescription Drugs and Biologics FDA - U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES Notification of Cosmetic Product Reason for Issuing: Safety and Registration of the Food and Drug Administration (FDA) or the certification of compliance. The product contains no health hazard(s). Product Notification Number: PMA-2000-012345 Date: 10 September 2001 Particulars of the Product: Name of the Company: ABC Corporation Address of the Company: 1234 Main Street, Anytown, USA 12345 USA Name: ABC Corporation Description: A topical pharmaceutical product containing 10% hydrocortisone. Chemical Name: Hydrocortisone Other Information: This certificate is issued under the authority of the Food and Drug Administration. It certifies that the health and chemical health of the product has not changed since the information contained in this certificate was made. Reissuance of this certificate will require a new application and a fee. <p>The company, firmly convinced that they shall acquire any legal rights FDA confers by issuing this certificate, hereby declares that it will not commence the sale or distribution of the product until it has received a copy of the certificate and that it will not commence the sale or distribution of the product described in this certificate until such time as it has received a copy of the certificate or any confirmation of its issuance.</p> <p>This information is subject to inspection, prosecution, removal, recall or revocation of FDA license and/or suspension of all or any part of the product described by the manufacturer.</p> <p style="text-align: center;">DET. AUTHORITY OF THE MANUFACTURER/IMPORTER</p> <div style="text-align: center;">  <i>John Doe, President</i> <i>ABC Corporation</i> </div>
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 	Regulations of the National Institutes of Health NIH—NIH Grants, NIH Extramural Activities	
Notification of Extramural Project		
<p>Project under way involving one or more institutions of the Food and Drug Administration (FDA) to further the public health mission of the agency (e.g., Analytical Chemistry)</p>		
<p>Current Notification Period: View previous form 510(k)</p>		
<p>Validity: 24 September 2012</p>		
<p>Identification of the Project:</p>		
<p>Product Name: PAPYRUS PAPYRUS PAPYRUS PAPYRUS PAPYRUS</p>		
<p>Product Type: ARTICLE ARTICLE ARTICLE ARTICLE ARTICLE</p>		
<p>Category: NOTIFICATION</p>		
<p>Name of the Company: PAPYRUS, INC.</p>		
<p>Address of the Company: 11575 UNIVERSITY DR STE 100, ALEXANDRIA, VA 22314-2345, USA</p>		
<p>Office Number: 11575 UNIVERSITY DR STE 100, ALEXANDRIA, VA 22314-2345</p>		
<p>The producer is attempting to or is about to sell the listed medical device in compliance with the requirements of the PMA (510(k)) notification and the FDA has issued a preliminary ruling (PR) or equivalent. This notification should not be taken as a guarantee of future enforcement of the safety profile and performance of the product. The FDA may take action at any time if it determines that the product does not meet its established safety and performance standards, even though no enforcement action has been taken.</p>		
<p>The company hereby certifies that they shall support any company listed with the FDA with regard to any enforcement investigation activity initiated by the FDA. Further, the company shall be responsible for ensuring the safe design, manufacture, or use of the medical device in view of the legal requirements, and conformeity to all the conditions and specifications of the premarket clearance or the PMA.</p>		
<p>This submission is subject to termination, modification, or recall should any subsequent FDA inspection, and/or enforcement action, determine the product to be contraindicated.</p>		
<p>BY AUTHORITY OF THE DIRECTOR OF THE CENTER</p>		
		
<p>View All Notifications View My Notifications</p>		
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GLOBAL RECOGNITION WITH TRUSTED ACCREDITATIONS

