

Neuramis® Dermal Fillers

The Smart Choice for
Facial Rejuvenation &
Structural Contouring.

EU MDR CE Certified • SHAPE™ Technology

Distributed globally by Yaksu Medi.



THE NEURAMIS ADVANTAGE



ENHANCED SAFETY

Manufactured with ultra-pure, non-animal Hyaluronic Acid from Japan (Shiseido). Raw materials are registered in the U.S. FDA Drug Master Files (DMF) and EDQM certified.



LONG-LASTING EFFECT

Patented SHAPE™ Technology employs a 2-step cross-linking process, enhancing structural cohesion and extending duration inside the dermis.



HIGH SATISFACTION

Outstanding skin integration and monophasic cohesiveness ensure natural-looking volume and immediate hydrobalance without artificial stiffening.

CORE INNOVATION: SHAPE™ TECHNOLOGY

Stabilized Hyaluronic Acid and
Purification Enhancement.



PREMIUM HA SOURCING

Utilizing microbial fermentation to produce high molecular weight, animal-free Hyaluronic Acid.

2-STEP CROSS-LINKING

Instead of standard single-step methods, SHAPE applies a uniform, dense network of BDDE bonds, resulting in a highly cohesive monophasic gel that resists early degradation.

ENHANCED PURIFICATION

A prolonged natural dialysis process washes the gel, radically minimizing uncross-linked (residual) BDDE to non-detectable levels.

THE ULTIMATE PURITY STANDARD

	INDUSTRY STANDARD (PH. EUR)	NEURAMIS STANDARD
Residual Protein	Max 0.3% or 0.1%	< 0.1%
Endotoxins	< 0.5 or 0.05 IU/mg	< 0.04 IU/mg
Residual BDDE	2 ppm Average	< 0.4 ppm (Non-detectable)



By exceeding European Pharmacopoeia standards and achieving EU MDR CE certification, Neuramis effectively mitigates the risk of delayed inflammatory reactions and edema.

THE NEURAMIS PORTFOLIO MATRIX

Global Spec: 20 mg/mL HA
Concentration across all variants

Superficial → Deep / Subcutaneous

Meso	Light Lidocaine	Lidocaine	Deep	Deep Lidocaine	Volume Lidocaine
Non-cross-linked + 0.9% Mannitol	0.3% Lido	0.3% Lido	No Lido	0.3% Lido	0.3% Lido
Depth: Superficial	Depth: Superficial/Mid	Depth: Mid Dermis	Depth: Mid/Deep Dermis	Depth: Mid/Deep Dermis	Depth: Deep/SubQ
Needle: 30G UTW	Needle: 30G UTW	Needle: 30G UTW	Needle: 27G UTW	Needle: 27G UTW	Needle: 27G UTW

*UTW = Ultra Thin Wall needles improve flow rates and lower extrusion force.

SUPERFICIAL TO MID-DERMIS SOLUTIONS



NEURAMIS MESO

Formula: 20mg/ml HA + 0.9% Mannitol
(antioxidant protection).

Indications: Skin hydrobalance, texture
improvement, deep hydration.



NEURAMIS LIGHT LIDOCAINE

Formula: 20mg/ml HA + 0.3% Lidocaine.

Indications: Correction of fine, superficial
wrinkles (undereyes, subtle lip hydration).



NEURAMIS LIDOCAINE

Formula: 20mg/ml HA + 0.3% Lidocaine.

Indications: Moderate wrinkles, glabellar
lines, forehead lines, and baseline lip
contouring.

MID TO DEEP DERMIS SOLUTIONS

Severe Wrinkles & Dynamic Contouring



NEURAMIS DEEP

Profile: Formulated without lidocaine for patients with local anesthetic hypersensitivity or allergies.

Delivery: 27G x 1/2" (13mm) UTW needle.



NEURAMIS DEEP LIDOCAINE

Profile: The most popular global variant, featuring 0.3% lidocaine for maximum patient comfort.

Shared Indications: Correction of severe wrinkles, pronounced nasolabial folds, marionette lines, and advanced lip volumization (e.g., Russian Lip technique).

Longevity: Clinically proven efficacy lasting 9-12 months.

Severe Volumization & Structuring

Neuramis Volume Lidocaine

Viscoelasticity: The highest viscosity, elasticity, and cross-linking density in the Medytox portfolio.

Injection Depth: Designed for deep dermis, supraperiosteal, or subcutaneous injection.

Indications: Non-surgical facelift techniques, restoring severe volume loss, cheek (malar) augmentation, chin/jawline contouring, and m-surgical rhinoplasty, chin/jawline contouring, and non-surgical rhinoplasty.

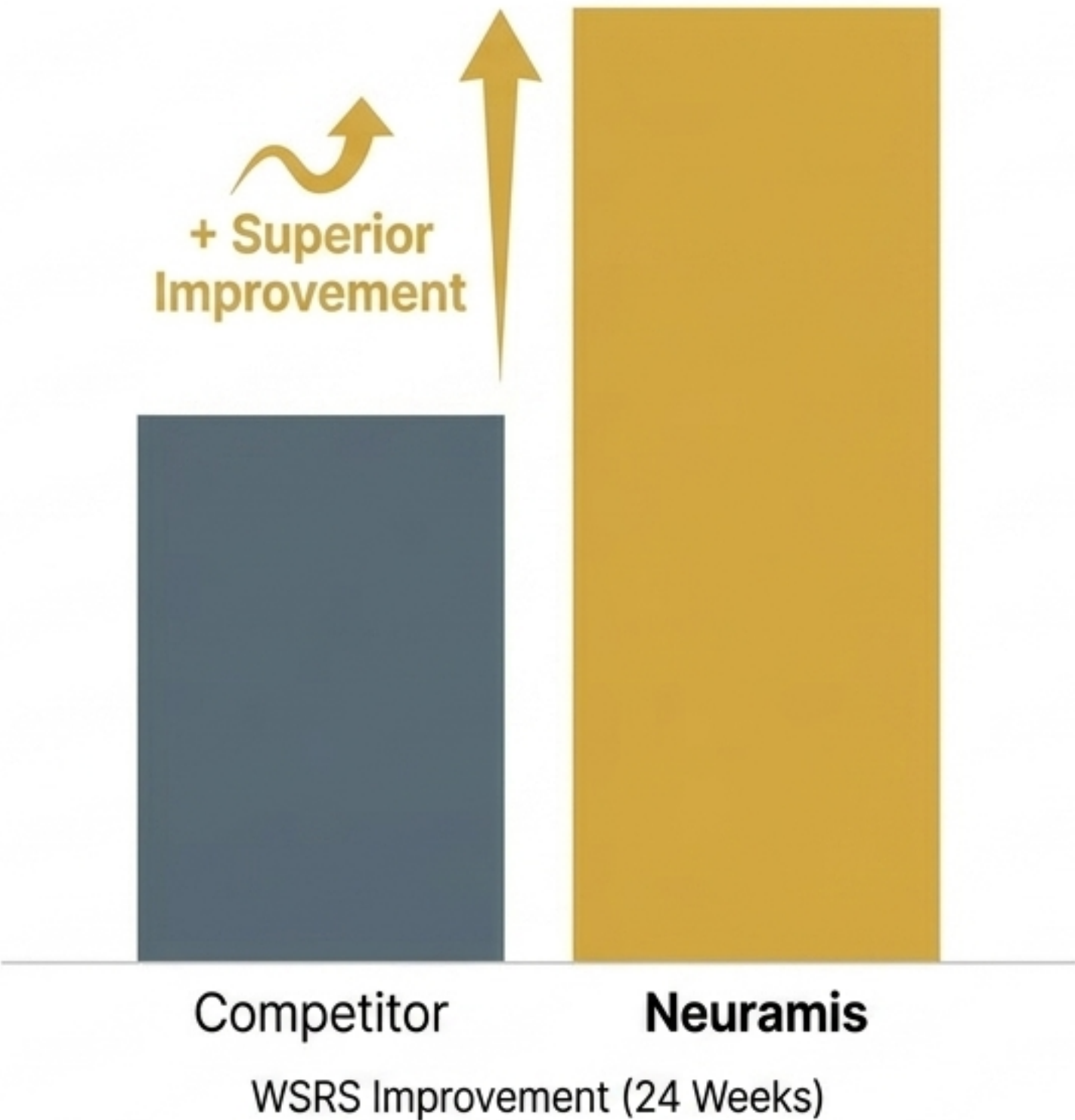
Hardware: 27G x 1/2" (13mm) UTW needle for high mechanical lifting capacity.

Longevity: 12-18 months of sustained structural support.



Proven Clinical Efficacy

Phase III Clinical Trial Highlights
Study Design: Phase III, randomized, multi-center, double-blind, active-controlled trial (Neuramis Deep Lidocaine vs. Product A/Perlane-L) for nasolabial folds.
Results (WSRS): Demonstrated non-inferiority and superior Wrinkle Severity Rating Scale (WSRS) improvement at 24 weeks.
Safety Profile: Statistically comparable or lower rates of treatment-related adverse events.



EU MDR CE Certification

Classifying Neuramis as a highly verified Class III medical device meeting the strictest European technical and pharmacological assessments.



Procure Authentic Neuramis® Today

Supplier Information Block

Company: Yaksu Medi

Website: yaksumedi.com

Whatsapp: +82 10 7240 1990



Ensure Patient Safety. Always source authentic Neuramis products. Look for the genuine Medytox protective hologram and exact packaging color-coding to guarantee EU-compliant purity and performance.

Contact Yaksu Medi today for global B2B wholesale pricing, clinical supply, and verified logistics.