

# 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
<b>Residual Protein</b>	Max 0.3% or 0.1%	<b>&lt; 0.1%</b>
<b>Endotoxins</b>	< 0.5 or 0.05 IU/mg	<b>&lt; 0.04 IU/mg</b>
<b>Residual BDDE</b>	2 ppm Average	<b>&lt; 0.4 ppm (Non-detectable)</b>



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

<b>Meso</b>	<b>Light Lidocaine</b>	<b>Lidocaine</b>	<b>Deep</b>	<b>Deep Lidocaine</b>	<b>Volume Lidocaine</b>
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.



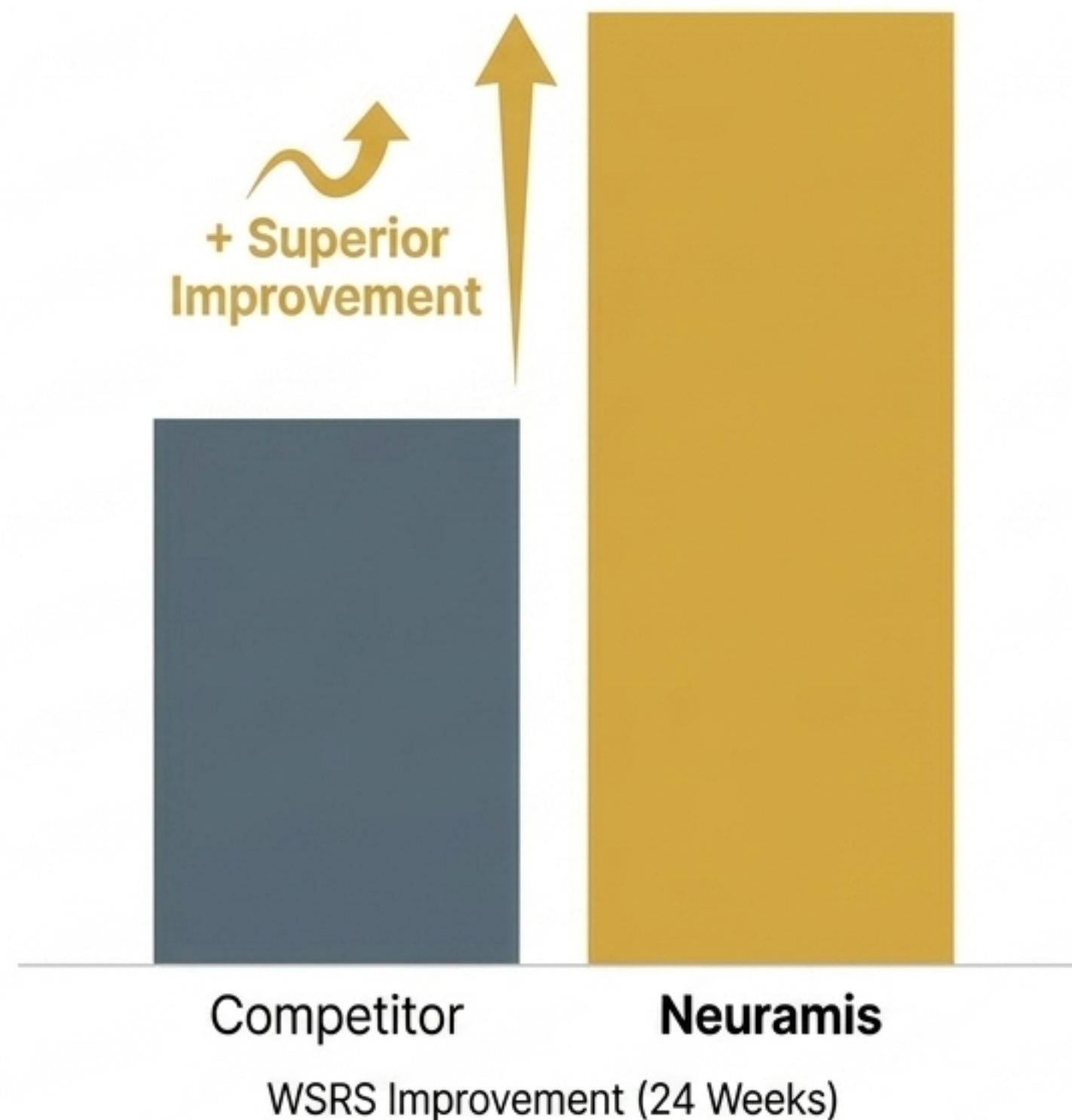
# Proven Clinical Efficacy

## 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

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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.**