

 Rhei Life

**Inspired by
Heritage.
Proven by
Science.**

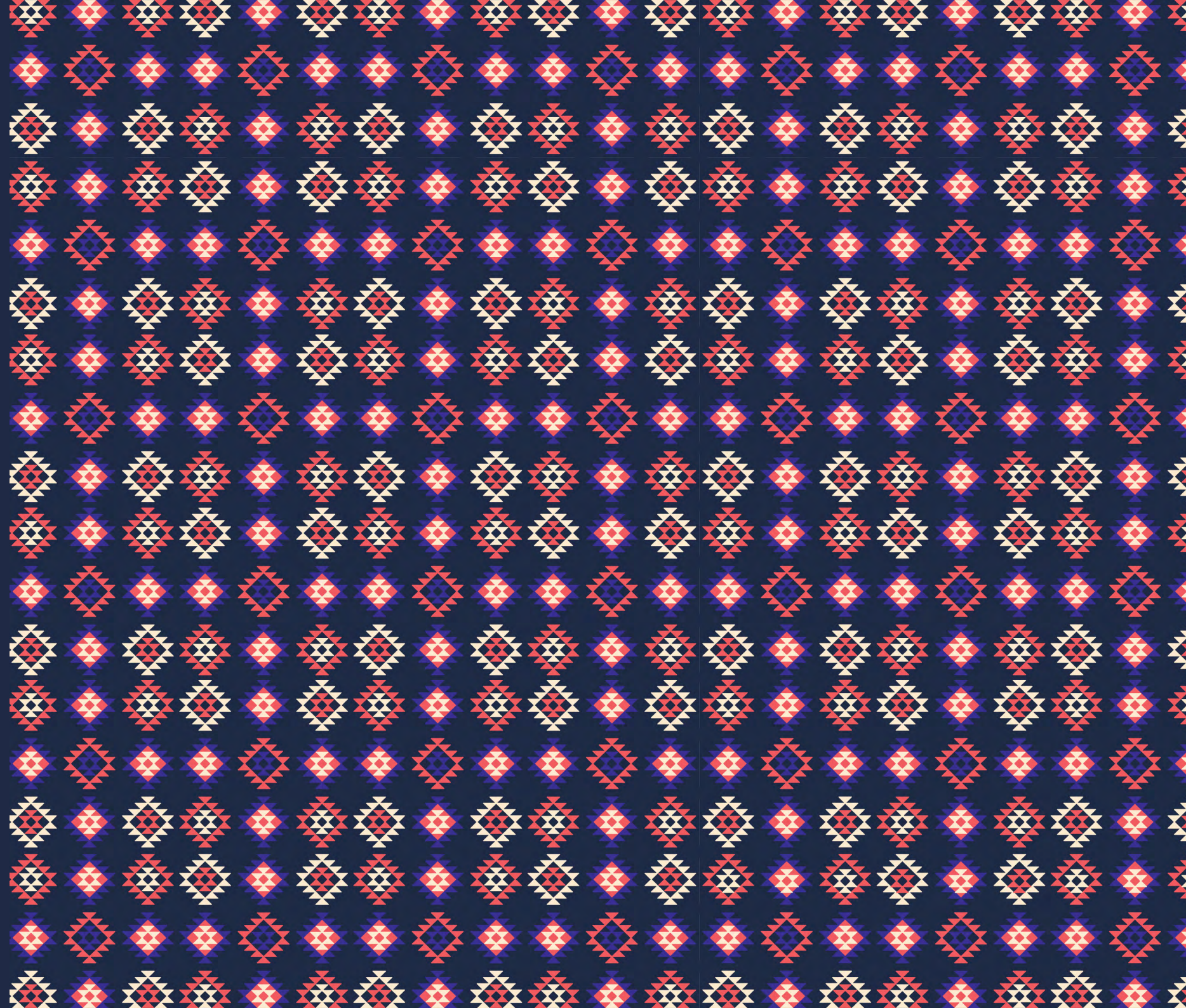


PORTFOLIO
Finished Products





Unique. Innovative.
Breakthrough.



WHO WE ARE

Founded in 2015 in Serbia and the United Arab Emirates, Rhei Life was established with a clear vision: to become a relevant and reliable stakeholder in the global healthcare landscape. The company initially focused on the Adriatic region, followed by systematic expansion across Europe. Today, Rhei Life is present and recognized across the global healthcare scene.

Our growth and achievements are built on two key pillars: a highly dedicated, multidisciplinary team of professionals and a carefully developed global network of partnerships. These long-term partnerships, nurtured with trust and commitment over the past decade, represent the most valuable capital of our company and an integral part of every success we have achieved.

TWO STRATEGIC DIVISIONS – ONE CLEAR VISION

Rhei Life operates through two clearly defined and complementary business divisions, both aligned with our long-term strategy and global ambitions.

FIRST DIVISION

Products Development for

GLOBAL MARKETS

Through continuous investments in research and development, Rhei Life positions itself as a global healthcare stakeholder by developing innovative products used by patients worldwide. Our portfolio includes medicines, food supplements, and medical devices, developed either in-house or through co-development models in collaboration with strategic partners.

Our products are commercialized globally through a trusted network of partners and customers who have placed their confidence in Rhei Life across all continents.

Today, Rhei Life products are present in **47 countries**, with:

- **80 partner companies worldwide**
- **Launched products or signed commercial agreements**

References

TRUSTED BY GLOBAL PHARMACEUTICAL COMPANIES

Rhei Life products are trusted by leading global pharmaceutical companies that have included our products in their portfolios across multiple therapeutic areas and markets worldwide.


These collaborations reflect long-term confidence in the quality, safety, scientific value, and commercial potential of Rhei Life products.

Recognize our products and Rhei Life as a reliable, long-term collaborator capable of delivering innovation, regulatory expertise, and commercial excellence.

Trusted by Global

HEALTHCARE LEADERS

With over 80 partners worldwide, we are proud to support leading pharmaceutical and healthcare companies with our products, including:



Teva · Stada · Zentiva · Sun Pharma · Dr. Reddy's · Walgreens Boots Alliance · Polpharma · Mega Labs · Duopharma Biotech Berhad · EL PHARMA (PHOENIX Group) · Al Safa United · Tamer Group · Lancasco · Unilab · Bivid

Special attention is given to **confirming product efficacy and safety through clinical studies conducted in accordance with international standards.**

In addition to in-house development, selected products are developed in close cooperation with partners through co-development models.

Our in-house developed products are manufactured in **11 different CMOs across Europe**, fulfilling **strict regulatory and quality requirements** for global markets. Manufacturing standards meet the requirements of national regulatory authorities as well as the stringent quality expectations of multinational pharmaceutical companies successfully promoting our products in some of the world's most regulated markets.

SECOND DIVISION

Representation and Distribution

SERVICES IN EUROPE

Through our European entity, as EU GDP license holder, Rhei Life acts as a Marketing Authorization Holder (MAH) for:

- **364 SKUs**
- **Generic medicines, OTC products, medical devices, and food supplements**
- **11 Eastern European markets**

Within this division, **Rhei Life represents 37 manufacturing companies from across the world**, providing comprehensive regulatory, distribution, and commercial services for European markets.

LEADERSHIP TEAM

Rhei Life is led by an experienced leadership team with strong expertise across pharmaceuticals, R&D, regulatory affairs, quality, supply chain management, and global business development.



MILOŠ STAKIĆ
Chief Executive Officer (CEO)



LUKA RADOJČIĆ
Chief Financial Officer (CFO)



JELENA GROZDANOVIĆ, MD
Head of Medical & Marketing Department



MARKO JEZDIĆ
Head of Adriatic Region



DARKO PETAKOV
Head of Global Business Development



JASMINA NENEZIĆ
Head of Supply Chain & Logistics



DORIS MILOŠEVIĆ
Head of In-Licensing



VLADIMIR JOVANOVIĆ
Legal & Compliance Manager



BOŠKO ČUBRA
Head of Tender Sales Department



VESNA PLAMENAC
Head of Quality Department



GLOBAL BUSINESS DEVELOPMENT TEAM

Our Global Business Development Team ensures strong international presence, deep local market understanding, and long-term partnership development across all regions.

OUT-LICENSING



SLOBODAN STEVIĆ
Senior Global BD Manager APAC



PREDRAG JOVANOVIĆ
Global BD Manager EU and CIS



DAVID STAKIĆ
Senior Global BD Manager MENA



PRACHI SABOO
Global BD Manager LATAM and Central America



DRAGANA JEREMIĆ
Senior Adriatic BD Manager



BRANKA JANKOVIĆ
Adriatic BD Manager



TEODORA MIKULIĆ
BD specialist



SANJA ŠKONDRIĆ
BD specialist



DAMLA UNLU PETROVIĆ
Global BD Manager MENA



MATEJA MARKOVIĆ
BD specialist



SARA POPADIĆ
Adriatic BD Manager



SLOBODAN GRAOVAC
Senior In-Licensing Manager



NEMANJA MIŠKOVIĆ
Senior In-Licensing Manager



SOFIJA STOJANOVIĆ
Senior In-Licensing Manager



JELENA STAMENKOVIĆ
In-Licensing Manager



IN-LICENSING



JOVANA JOKSIMOVIĆ
International Regulatory Affairs Manager



IVA GIBA
International Regulatory Affairs Manager



MARINA TOMAŠEVIĆ
International Regulatory Affairs Associate



NIKOLA PEŠIĆ
R&D Manager



BARBARA MILOVANČEVIĆ
Medical Affairs Specialist



ANA JELOVAC
Medical Affairs Specialist



STEFAN MASNIKOSA
Medical Affairs Specialist



REGULATORY & MEDICAL

PORTFOLIO

Finished Products

Book your meeting:
business.development@rhei.life

THERAPEUTICAL AREA	PRODUCT NAME	CATEGORY	FORM
		Food supplement	capsules
		Food supplement	dual sachets
		Medicine	film-coated tablets
		Medical device	hydrogel and liquid spray
		Medical device	fillers
		Cosmetics	cosmeceuticals
		Food supplement	capsules
		Food supplement	capsules
		Food supplement	capsules
		Medicine (OTC / Rx)	film-coated tablets
		Medicine (OTC / Rx)	film-coated tablets
		Food supplement	capsules
		Food supplement	capsules
		Food supplement	capsules
		Food supplement	capsules
		Food supplement	capsules
		Medical device	prefilled syringe
		Medical device	spray ear drops
		Medical device	hydrogel and liquid spray
	Kids gummies	Food supplement	gummies
		Food supplement	gummies



GASTROENTEROLOGY
Food supplement

Hepalife® forte

Pharmaceutical form:

capsules

Packaging:

30 capsules

Recommended dosage:

1 capsule a day



INDICATED FOR

- Fatty liver therapy
- Chronic inflammatory liver disease (*hepatitis B and C*)
- Treatment of toxic liver damage (*alcohol, drugs and other harmful substances*)
- Ancillary therapy for liver cirrhosis
- Chronic damage from bile stone

COMPOSITION

- Siliphos® = silybin + phosphatidylcholine
- S-adenosyl-L-methionine (SAME)
- Choline L-bitartrate
- Vitamin E
- Vitamin B complex

SCIENTIFIC SUPPORT

Ongoing clinical trial

Efficacy and safety of Hepalife forte in treatment of patients with MASLD and MetALD

KEY ADVANTAGES

Hepalife® forte is a unique hepatoprotector that



Detoxifies



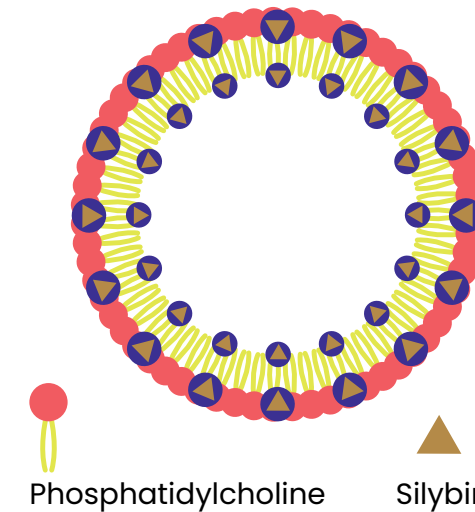
Regenerates



Protects the liver

Siliphos®

Trademark owned by Indena S.p.A., Italy.

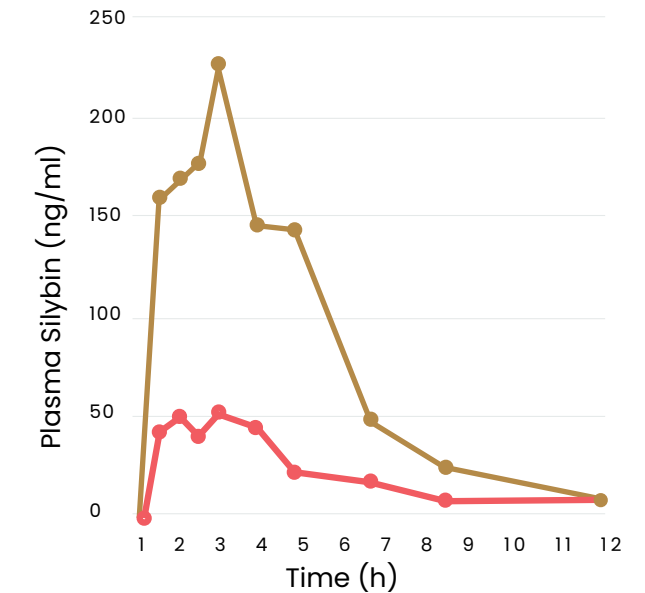


The absorption of Siliphos® is **4.6 times higher than silymarin.**¹

1. Kidd P, Head K. A review of the bioavailability and clinical efficacy of milk thistle phytosome: a silybin-phosphatidylcholine complex (Siliphos). *Altern Med Rev.* 2005 Sep;10(3):193-203. PMID: 16164374.

PATENTED

Siliphos® is a phytosome made up of **Silybin** and **Phosphatidylcholine**.



— Siliphos®
— Silymarin

CLICK **FOR MORE DETAILED PRESENTATION**



GASTROENTEROLOGY
Food supplement

DuoCare

Pharmaceutical form:

capsules

Packaging:

30 capsules

Recommended dosage:

1 capsule a day



INDICATED FOR

- Acute infectious diarrhea, regardless of the cause and presence of vomiting.
- Chronic watery diarrhea, regardless of etiology.
- Diarrhea caused by the use of antibiotics, laxatives, or other medications.
- Frequent vomiting.
- Elevated temperature associated with increased sweating.
- After sports activities and exercise.
- Traveler's diarrhea.

COMPOSITION

- Magnesium hydroxide
- Potassium chloride
- Sodium chloride
- *Lactobacillus rhamnosus GG*
- Sodium citrate
- Dextrose
- Maltodextrin

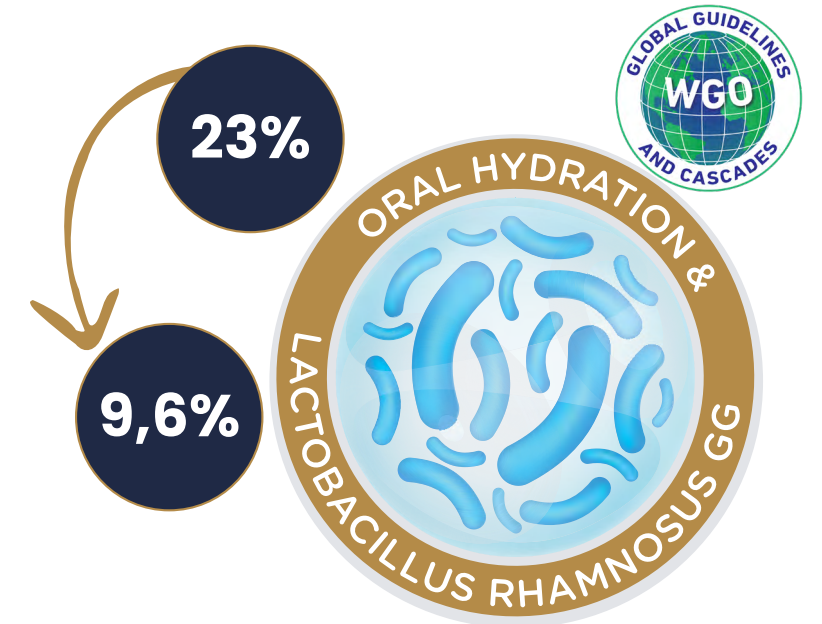
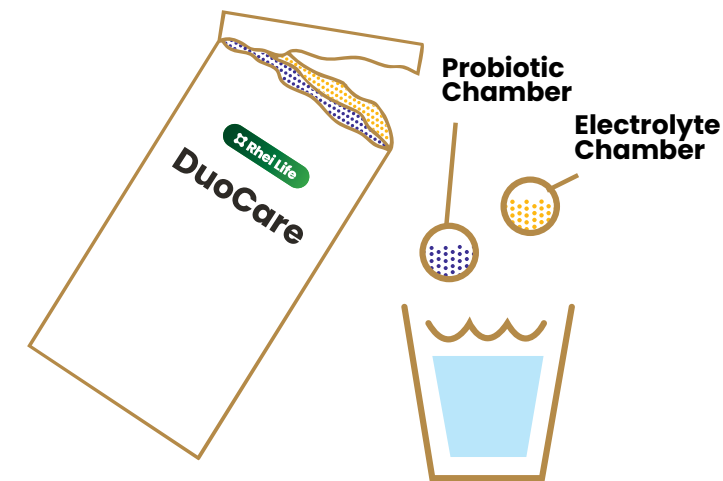
SCIENTIFIC SUPPORT

Lactobacillus Rhamnosus GG is one of the most clinically documented probiotic strains, supported by more than **250 scientific publications** and in use since 1990. According to the World Gastroenterology Organization (WGO) Global Guidelines (February 2023), LGG holds **Level 1 evidence**-the highest level of clinical evidence-for antibiotic-associated diarrhea, acute gastroenteritis, prevention of AAD, prevention of nosocomial diarrhea, and functional abdominal pain/IBS.

KEY ADVANTAGES

DuoCare contains a scientifically balanced blend of glucose, minerals and probiotic (*Lactobacillus Rhamnosus GG*) to replenish water and salt concentration and maintain gut flora as normal.

DUAL-CHAMBER SACHET



In a meta-analysis of 5 randomized clinical studies involving **445 children** from 2015, the use of LGG reduced the risk of AAD.

World Gastroenterology Organization Guide

February 2023

Disorder, condition	Probiotic strain	Level of evidence
Antibiotic associated diarrhea (AAD)	<i>Lactobacillus rhamnosus GG</i>	1
Acute gastroenteritis	<i>Lactobacillus rhamnosus GG</i>	1
AAD prevention	<i>Lactobacillus rhamnosus GG</i>	1
Prevention of nosocomial diarrhea	<i>Lactobacillus rhamnosus GG</i>	1
Functional abdominal pain / IBS	<i>Lactobacillus rhamnosus GG</i>	1

CLICK **FOR MORE DETAILED PRESENTATION**



GASTROENTEROLOGY
film-coated tablets

Prucalopride

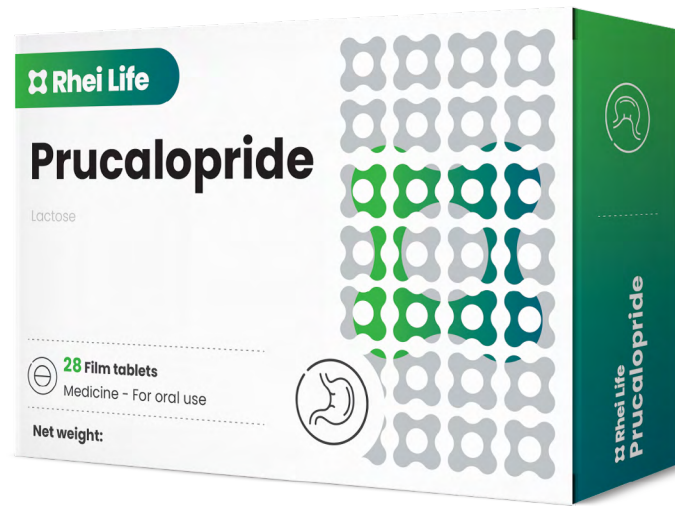
ATC code: A06AX05

INN: prucalopride succinate

Pharmaceutical form: Film-coated tablets

Dose: 1 or 2 mg

Pack size: 28



PHARMACODYNAMIC PROPERTIES

- Prucalopride has gastrointestinal prokinetic activity.
- Prucalopride is a selective agonist of high affinity for serotonin receptors (5-HT₄), which explains its prokinetic effect on gastric, intestinal and smooth muscles of colon.
- Prucalopride does not affect the function of glycoprotein P or cytochrome P450 and is not extensively metabolized in the body

SCIENTIFIC SUPPORT

European society of neurogastroenterology and motility guidelines on functional constipation in adults.

Level of evidence

High

Level of agreement

100%

Recommendation

Strong

The serotonin (5-HT)-4 agonist prucalopride has prokinetic action in the entire gut and is effective in the management of chronic constipation, including conditions refractory to conventional laxatives.

POSODOLOGY

Adults: 2 mg once daily with or without food, at any time of the day.

If the intake of once daily prucalopride is not effective after 4 weeks of treatment, the patient should be re-examined and the benefit of continuing treatment reconsidered.

The efficacy of prucalopride has been established in double-blind, placebo-controlled studies for up to 3 months. Efficacy beyond three months has not been demonstrated in placebo-controlled studies.



INDICATED FOR

Symptomatic treatment of chronic constipation in adults in whom laxatives fail to provide adequate relief.



SKIN AND WOUND CARE

Medical device

Effigerm Wound Wash

Pharmaceutical form:

Hydrogel and liquid spray

Packaging:

60ml, 150ml, 250ml bottles with finger spray or nozzle or 500 ml or 1000 ml bottle with trigger spray or screw cap

Recommended dosage:

Apply as required



INDICATED FOR

- Cleansing and debriding at the wound and preventing infections.
- For the management of minor skin abrasions, minor lacerations, minor irritations, minor cuts, minor burns and intact skin.
- Wound irrigation solution.
- To irrigate and moisten.
- For sore irritated skin.
- Helps with the prevention of biofilm
- Can be used for moistening the wound dressing and moisturising the wound itself as well as for loosening crusted dressings.
- Flushing of irritants associated with wound malodour.

COMPOSITION

- Hypochlorous acid (HOCl) 0,016 %

SCIENTIFIC SUPPORT

HOCl has been widely evaluated by regulatory bodies, healthcare institutions, and academic organizations in over 50 countries, with more than 2,600 peer-reviewed publications supporting its antimicrobial efficacy and over 3,000 publications addressing its role in wound healing enhancement. There are **no reports of adverse reactions to topical applications** by these methods based on the US EPA's Toxicology Database DSSTox, the US CDC Toxic Substances and Disease Registry, nor at either the Development and Reproductive Toxicology Database, or the European Bioinformatics Institute of EMBL.

KEY ADVANTAGES

Effigerm is a Wound Irrigation Solution for cleansing, rinsing and moisturizing of wounds. For the management of minor skin **abrasions, minor lacerations, minor irritations**, minor cuts, minor burns and Intact skin - helps prevent biofilm.



Wound Irrigation



Skin-friendly pH



Hygienically Clean



Hypoallergenic



Non-irritant



Dermatologically Tested

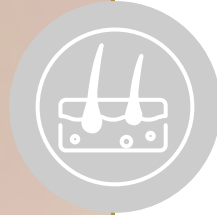


Helps prevent biofilm



Alcohol-Free

CLICK  FOR MORE DETAILED PRESENTATION



SKIN AND WOUND CARE
Medical device

Terra

PRODUCTS

Terra filler line consists of 5 different references, each one with a specific concentration- to be effective and perfectly integrated in different anatomical places.



FILLERS

Complete range of HA-based fillers:

- **BDDE cross-linked;**
- High elasticity;
- High level of skin compatibility and biocompatibility;
- Minimum risk of side effects;
- Natural and smooth feel in the tissue.

Fillers are enriched with **Proline** and **Glycine** that provide hydration and soften the skin tissue, acting as a humectant and emollient respectively.

These two amino acids are fundamental in stabilizing the structure of the hydrogel, especially in terms of viscoelastic properties and thermodynamic equilibrium and ensure better control of the swelling capacity of the hydrogel in the postimplant phase.



LIGHT

Concentration:

18 mg/ml of HA BDDE Cross-linked

FINE

Concentration:

22 mg/ml of HA BDDE Cross-linked

LIPS

Concentration:

24 mg/ml of HA BDDE Cross-linked

VOLUME

Concentration:

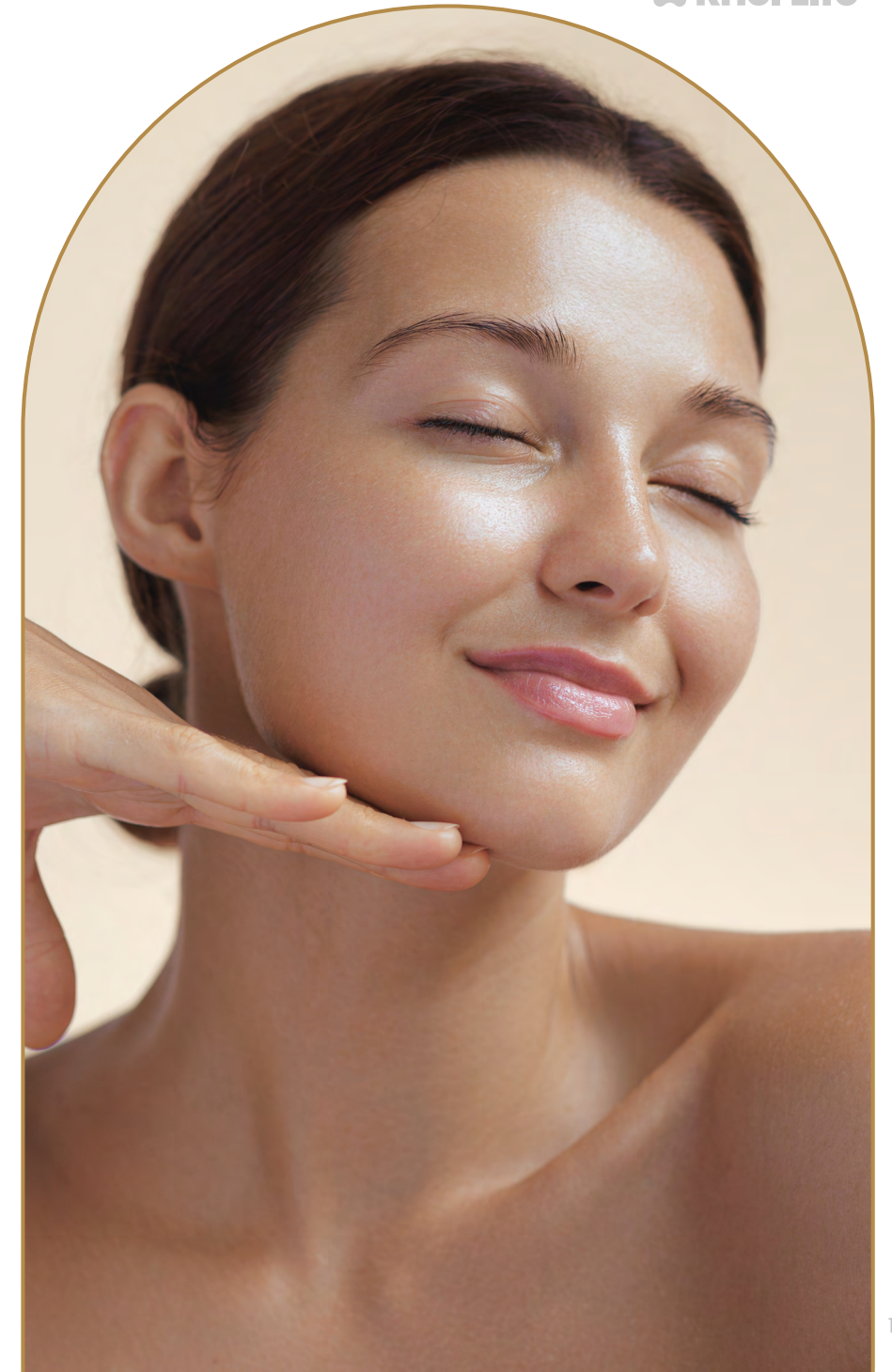
26 mg/ml of HA BDDE Cross-linked

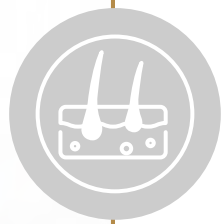
LIFT

Concentration:

28 mg/ml of HA BDDE Cross-linked

Class III medical devices regulated under the
EU MDR 2017/745





SKIN AND WOUND CARE
Cosmetics

Arista

Scientifically advanced skincare line that combines patented sonicated hyaluronic acid with high-quality active ingredients such as PDRN, growth factor mimetics, and ceramides.

All products are developed under strict GMP and ISO certifications, with sterile, single-dose formats that minimize contamination risk and reduce the need for traditional preservatives. The line offers targeted solutions for hydration, regeneration, anti-aging, and post aesthetic treatments, ensuring professional-level results through innovative technologies and delivery systems.



BEAUTY METHOD

ARISTA BEAUTY METHOD is a home-care line designed to extend and support professional aesthetic treatments.

Its advanced formulas combine patented sonicated hyaluronic acid with innovative active ingredients to address specific skin needs. The multi-molecular action provides deep hydration, smoothing, and revitalization while ensuring high bioavailability and skin tolerability. Suitable for daily and post-procedure use, it helps maintain skin glow and long-term results.

BEAUTY METHOD product range includes:

- Cleansing HA Gel
- Eye Contour Cream
- Perfect Brightness Cream
- Glow Up Cream
- Moisturizing HA Serum
- Retinol Serum
- Renewal Cream
- PDRN B-Mask
- Supreme Pure HA Serum
- Supreme Radiant SPF 50+
- Advanced Lift

REVITAL

ARISTA REVITAL is a professional post-procedure skincare line designed to support skin recovery and comfort after treatments such as chemical peels, lasers, and microneedling.

The preservative-free formulas have very low cytotoxicity and come in a dual-syringe system that is mixed before application to ensure a fresh, concentrated gel.

Based on a patented hyaluronic acid blend with targeted active ingredients, REVITAL soothes, hydrates, regenerates, brightens, and strengthens the skin barrier, reducing redness and discomfort while enhancing overall results.



CLICK  **FOR MORE DETAILED PRESENTATION**

VITARES

ARISTA VITARES is a professional line designed for advanced aesthetic and medical procedures. It includes ready-to-use, sterile solutions that support in-clinic treatments and enhance performance.

Packaged in single-dose, hermetically sealed glass vials, VITARES eliminates dilution and minimizes contamination risk. Made with medical-grade raw materials and produced in clean-room conditions with steam sterilization, it ensures safety, purity, and consistency.





NERVOUS SYSTEM Food supplement

Nerviplex

Pharmaceutical form:

capsules

Packaging:

20/30/60 capsules

Recommended dosage:

1 capsule a day

WATCH NOW

For **early stages of peripheral neuropathy symptoms with mild pain**



INDICATED FOR

- Painful syndrome in the lumbar region / Lumbago
- Diabetic peripheral neuropathy
- Sciatic pain
- Neck pain
- Painful syndrome in the shoulder and arm region
- Spinal nerve root compression various radiculopathies
- Disc herniation (before or postoperatively) with consequent neuralgia and polyneuropathy
- Pain due to osteoarthritis
- Postherpetic neuralgia

COMPOSITION

- UMP
- Niacin
- Vitamin B6
- Vitamin B1
- Folate
- Vitamin B12

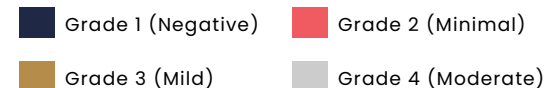
SCIENTIFIC SUPPORT

The efficacy of Nerviplex was confirmed in a randomized, double-blind, placebo-controlled clinical study in patients with carpal tunnel syndrome. The study demonstrated meaningful improvements in nerve function and functional recovery, along with reduced use of analgesics and a favorable safety profile, indicating true nerve recovery rather than only subjective pain relief.

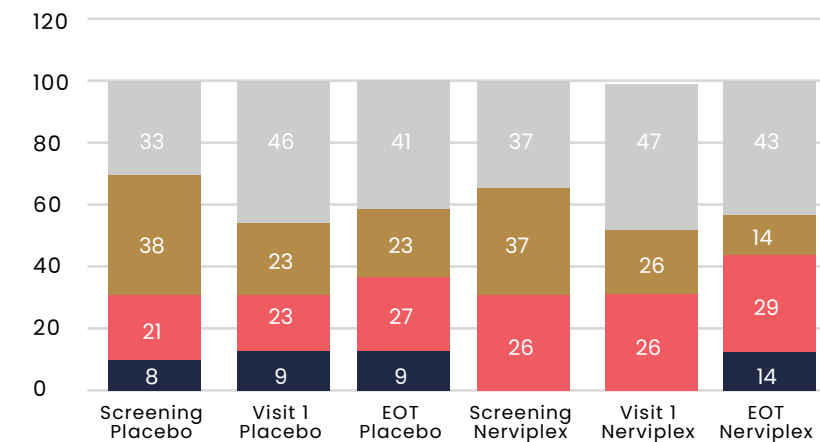
KEY ADVANTAGES

Is a nucleotide-based food supplement, with high concentration of nucleotides of pyrimidine bases and vitamins

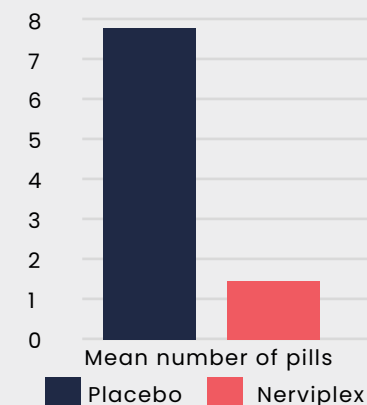
The proportion of subjects with **mild CTS (Grade 3)** decreased from **37 % at baseline to 14 %**, while the combined share of minimal and normal findings (**Grades 1-2**) increased from **26 % to 43 %**.



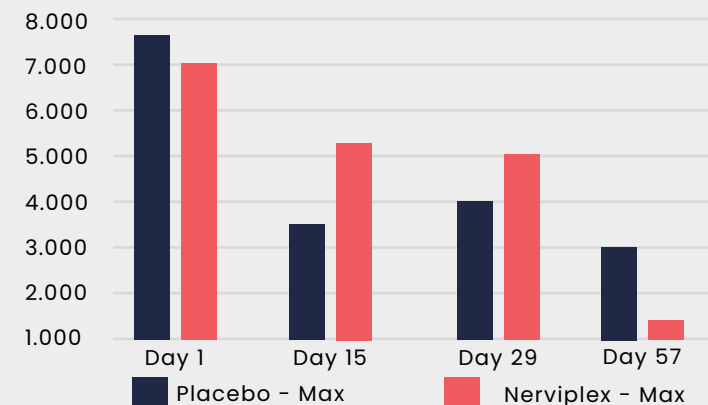
PADUA CLASSIFICATION changes



PAIN MEDICATION dose reduction

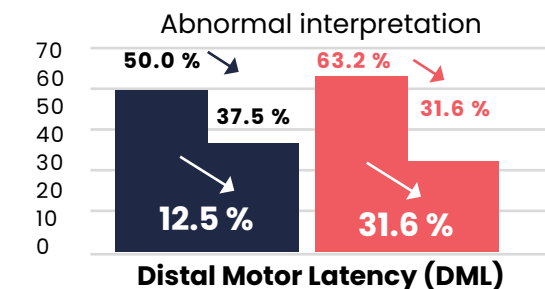
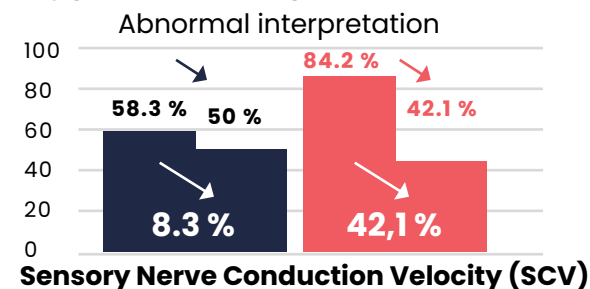


Cumulative Paracetamol or NSAIDs dose - Max



The Day 1 correction (7 000 mg) further supports the consistency of reduction over time **from 7 000 to 1 575 mg in Nerviplex, an almost 80% drop in maximum exposure.**

NCS PARAMETERS



CLICK FOR MORE DETAILED PRESENTATION



NERVOUS SYSTEM
Food supplement

Nerviplex forte
Pharmaceutical form:
capsules
Packaging:
20/30/60 capsules
Recommended dosage:
1 capsule a day

WATCH NOW ▶

For **moderate to severe** peripheral neuropathy



COMPOSITION

- UMP
- Vitamin B1
- CMP
- Vitamin B12

INDICATED FOR

- Painful syndrome in the lumbar region / Lumbago
- Diabetic peripheral neuropathy
- Sciatic pain
- Neck pain
- Painful syndrome in the shoulder and arm region
- Spinal nerve root compression various radiculopathies
- Disc herniation (before or postoperatively) with consequent neuralgia and polyneuropathy
- Pain due to osteoarthritis
- Postherpetic neuralgia

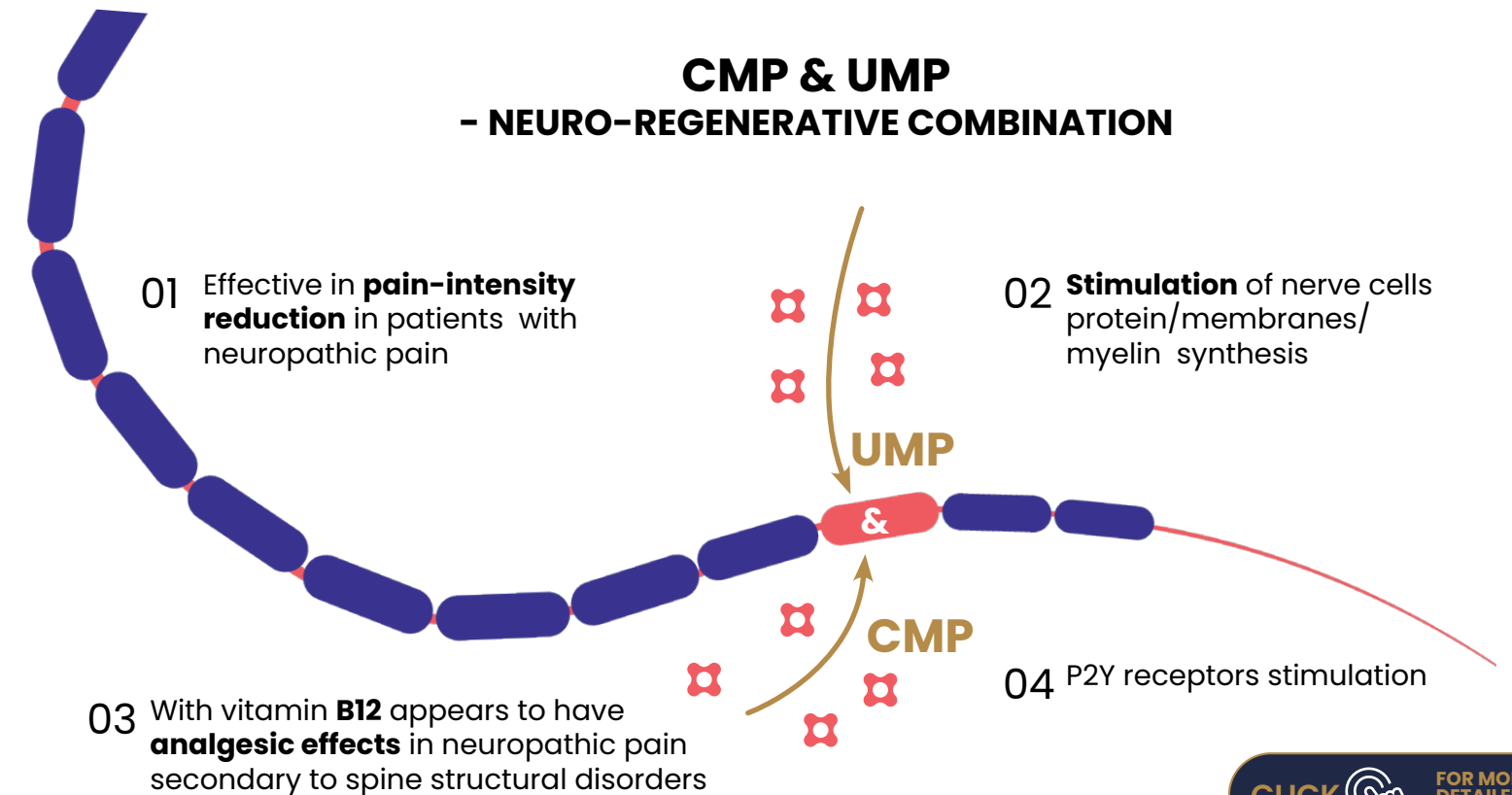
SCIENTIFIC SUPPORT

Nerviplex forte is currently being evaluated in the Safety and Efficacy of Nucleotides and B Vitamins for Treatment of Neuropathic Pain in Distal Symmetric Diabetic Polyneuropathy. By integrating subjective pain assessment with objective neurophysiological and biomarker endpoints, NUBVIDIA is designed to demonstrate not only symptomatic relief but also potential neuroregenerative and disease-modifying effects of the nucleotide-based formulation in diabetic neuropathy.

KEY ADVANTAGES

Is a nucleotide-based food supplement, with high concentration of nucleotides of pyrimidine bases and vitamins.

CMP & UMP
- NEURO-REGENERATIVE COMBINATION



CLICK  FOR MORE DETAILED PRESENTATION

RiboCARE[®] NEURO



RiboCARE is a registered trademark in the EU

INDICATED FOR

- Support brain health and nervous system function
- Helps improve cognitive performance, memory, and neuroprotection

COMPOSITION

- Cognizin[®] (Citicoline)
- RiboCARE[®] NEURO (nucleotide complex)
- Vitamin B12
- Vitamin B6
- Inulin

NERVOUS SYSTEM
Food supplement

Nucleocolin

Pharmaceutical form:

Capsules/ sachets

Packaging:

30 capsules/ 30 sachets

Recommended dosage:

2-4 capsules / 1-2 sachets daily



SCIENTIFIC SUPPORT

Cognizin[®] and RiboCARE[®] NEURO are clinically studied compounds known for their neuroprotective and cognitive benefits. Citicoline supports brain metabolism, neurotransmitter production, and neuronal repair, while RiboCARE[®] NEURO provides essential bioactive nutrients for neuronal communication, memory, and neuroprotection, particularly in cognitive decline disorders

KEY ADVANTAGES

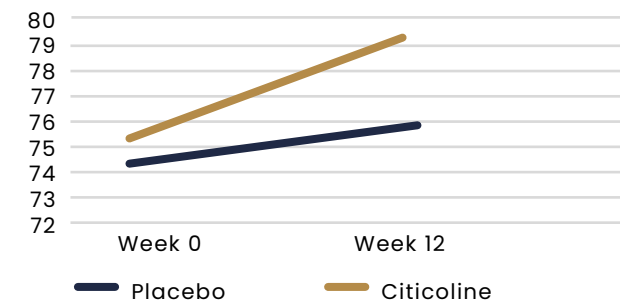
Nucleocolin is an advanced dietary supplement designed to support brain health and nervous system function. Featuring a synergistic blend of Citicoline (Cognizin[®]), nucleotides (RiboCARE[®] NEURO), and essential B vitamins, it helps improve cognitive performance, memory, and neuroprotection.

Cognizin[®]

For the evolution of your mind[®]

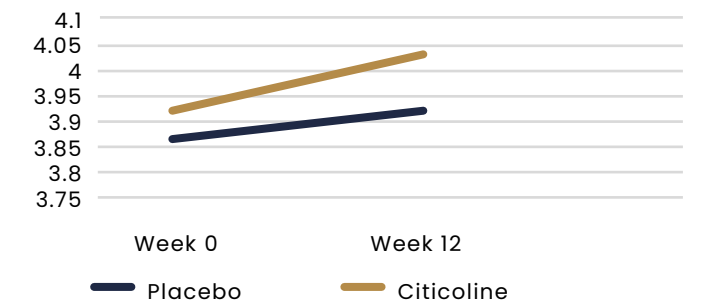
IMPROVING OVERALL MEMORY PERFORMANCE.

Significant improvement after the administration of citicoline



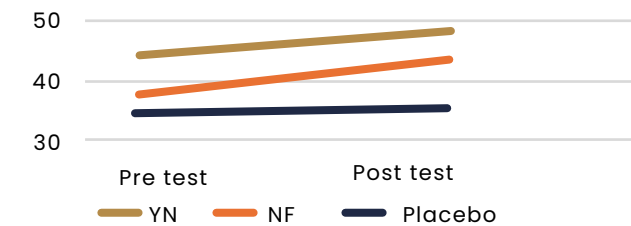
IMPROVEMENT OF EPISODIC MEMORY

Significant improvement in episodic memory of citicoline

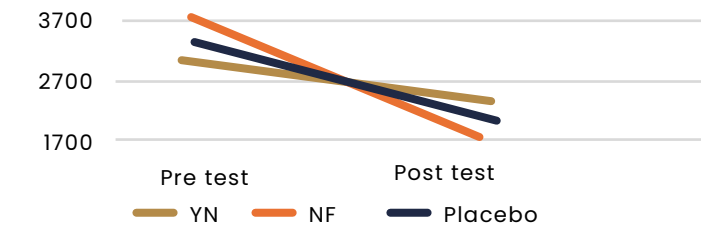


RiboCARE[®] NEURO

BRAIN-DERIVED NEUROTROPHIC FACTOR



F2-ISOPROSTANES



Cognizin[®] is a registered trademark of Kyowa Hakko Bio Co., Ltd
RiboCARE[®] is a Prosol S.r.l. brand registered in EU.

CLICK FOR MORE DETAILED PRESENTATION



NERVOUS SYSTEM
Medicine (OTC / Rx)

Bineuro
ATC code: A11EA
INN: Benfotiamine, Pyridoxine, Cyanocobalamin
Pharmaceutical form: Film-coated tablets
Route of administration: oral route
Dose: 150 mg / 50 mg / 0,2 mg
Pack size: 30

INDICATED FOR

Systemic neurological disease (peripheral neuropathy) with demonstrated deficiency of vitamins B1, B6 and B12.

KEY ADVANTAGES

- Benfotiamine, in contrast to most other thiamine derivatives, is a lipophilic compound (allowing to cross cell membranes easily) and, with a superior absorption, bioavailability, and effectiveness than thiamine.
- Maximum plasma levels of thiamine are about 5-fold higher after benfotiamine intake and the bioavailability is about 3.6 times as high as that of thiamine hydrochloride and better than that of other lipophilic thiamine derivatives.
- A 50 mg dose of vitamin B6 is safer for long-term use than higher concentrations, as it provides sufficient benefits without increasing the risk of nerve damage associated with high doses.

CLICK **FOR MORE DETAILED PRESENTATION**

NERVOUS SYSTEM
Medicine (OTC / Rx)

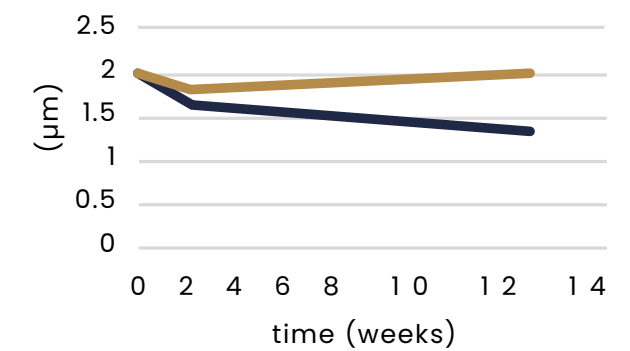
Bineuro forte
ATC code: A11EA
INN: Benfotiamine, Pyridoxine, Cyanocobalamin
Pharmaceutical form: Film-coated tablets
Route of administration: oral route
Dose: 150 mg / 50 mg / 1 mg
Pack size: 30



SCIENTIFIC SUPPORT

Benfotiamine + Pyridoxine + Cyanocobalamin

Improvement of vibration perception in diabetic polyneuropathy



30% reduction in vibration perception threshold

Changes in vibration perception threshold of the IInd metacarpal bone.

— B vitamins — Placebo

Stracke H, Lindemann A, Federlin K. A benfotiamine-vitamin B combination in treatment of diabetic polyneuropathy. *Exp Clin Endocrinol Diabetes*. 1996;104(4):311–316.



PAIN RELIEF
Food supplement

Rheipeptidase
Rheipeptidase forte
Rheipeptidase MAX

Pharmaceutical form:

capsules

Packaging:

15 capsules

Recommended dosage:

1-2 capsules a day depends on the type of product and condition.



INDICATED FOR

They can be used in all acute inflammatory conditions for the following indications:

- Surgery
- Otorhinolaryngology (ENT – ear, nose, throat)
- Obstetrics and Gynecology
- Orthopedics
- Pulmonology

COMPOSITION

- | | | |
|----------------------|----------------------------|--------------------------|
| RHEIPEPTIDASE | RHEIPEPTIDASE FORTE | RHEIPEPTIDASE MAX |
| • 60.000 SPU* | • 120.000 SPU* | • 250.000 SPU* |

* SPU = serrapeptase units

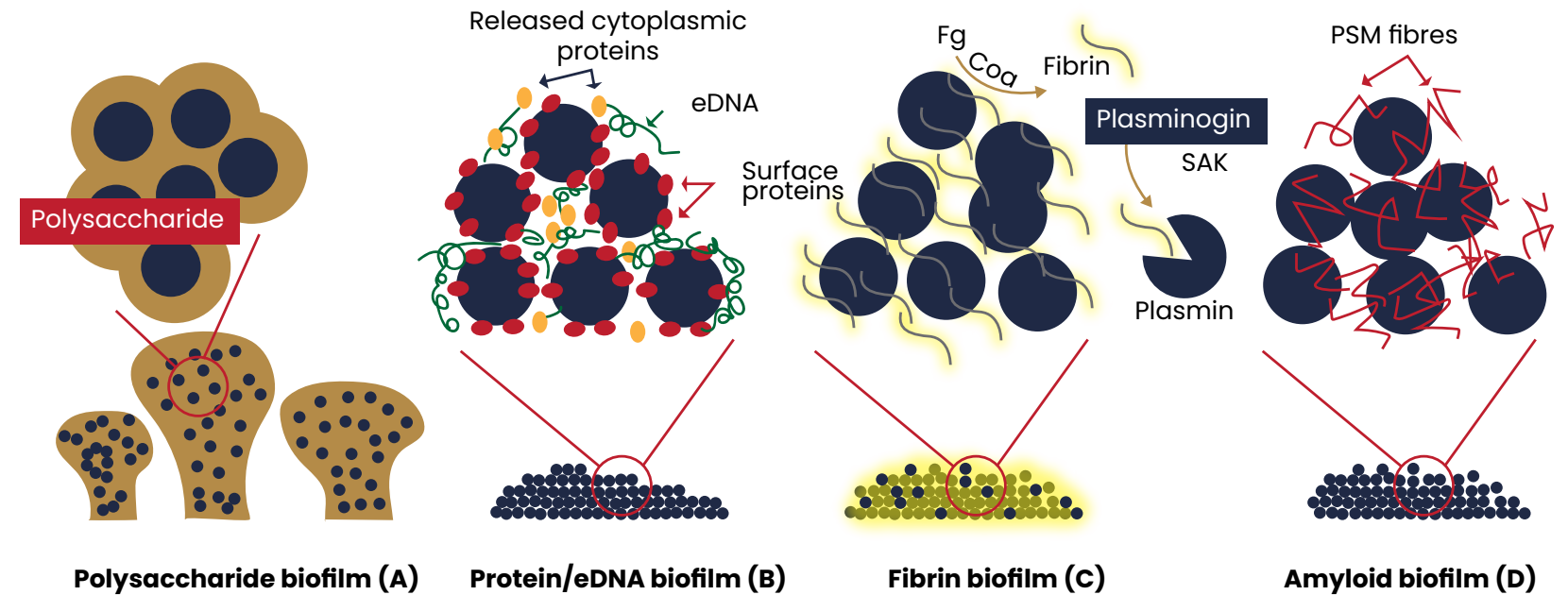
SCIENTIFIC SUPPORT

Experimental and clinical studies suggest anti-inflammatory, anti-edematous, mucolytic, and fibrinolytic effects, particularly in postoperative swelling, ENT disorders, musculoskeletal inflammation, and respiratory conditions with excessive mucus production. Additionally, in vitro data indicate that serratiopeptidase may disrupt bacterial biofilms, potentially enhancing antibiotic penetration when used concomitantly.

KEY ADVANTAGES

It is a proteolytic enzyme that has been used worldwide for more than thirty years. First isolated from *Serratia marcescens*. It has a highly pronounced enzymatic activity including a:

- potent anti-inflammatory
- antiedematous
- mucolytic
- fibrinolytic effect



1. Relieves inflammation, dilutes liquids formed after injury and facilitates their drainage
2. Serrapeptase blocks release of amines causing pain in the inflamed tissue
3. Breaks down fibrin and other non-living tissues without affecting the living tissue
4. Serrapeptase enzyme enhances the penetration of antibiotics when administered concomitantly by breaking down the biofilm created by bacteria

CLICK **FOR MORE DETAILED PRESENTATION**



PAIN RELIEF
Food supplement

Peamol

Pharmaceutical form:

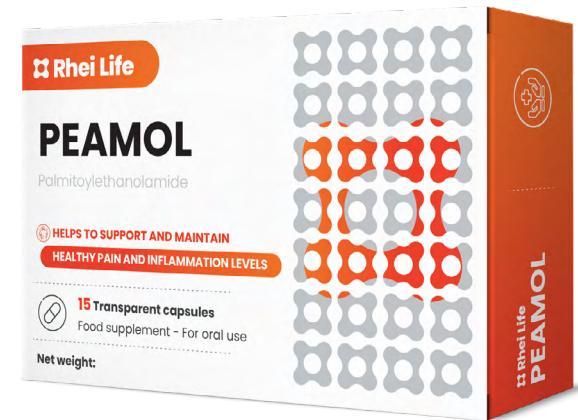
capsules

Packaging:

15 capsules

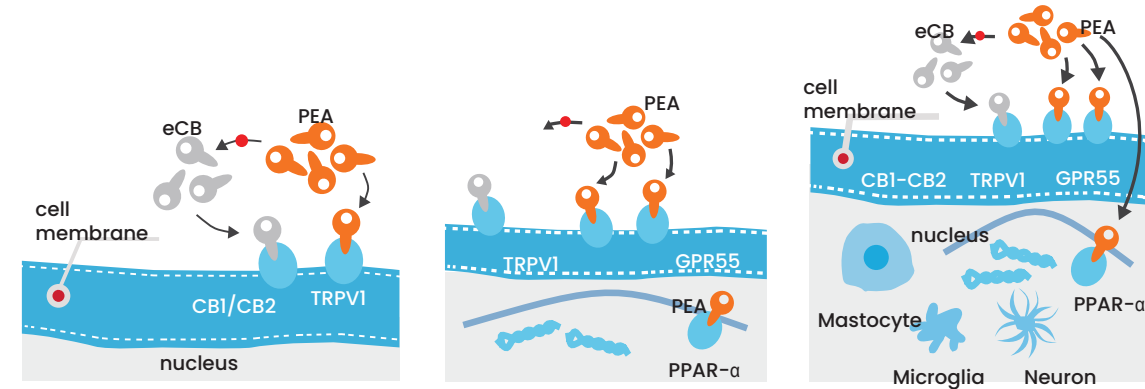
Recommended dosage:

1-2 capsules a day



KEY ADVANTAGES

PEAMOL is a dietary supplement containing palmitoylethanolamide (PEA) that has been shown to have neuroprotective, anti-inflammatory, and analgesic properties



Indirectly by activating of cannabinoid receptors CB1 and CB2 and TRPV1 ion channel

Directly through the PPAR- α and GPR55 receptors

By regulating the activity of mast cells and microglia

- Natural, non-opioid compound (no addictive potential and no pharmacological tolerance)
- Favorable safety profile – good tolerability with no acute or chronic toxicity reported
- No known drug interactions or contraindications
- Suitable for patients with impaired renal or hepatic function

INDICATED FOR

Chronic pain therapy in various etiopathogenesis

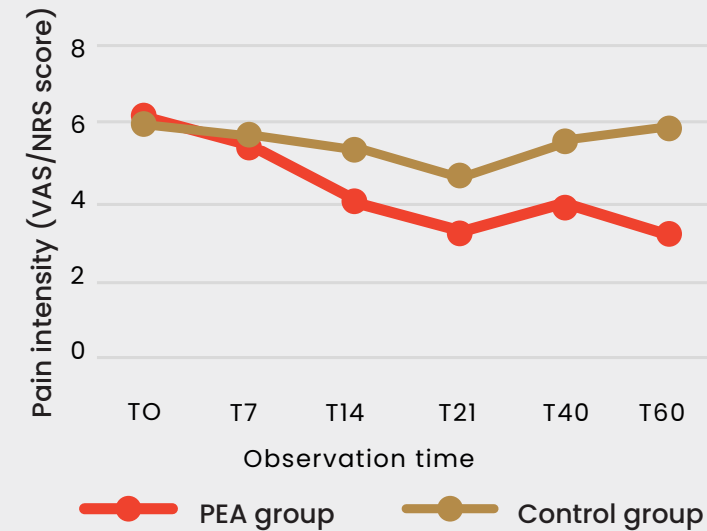
- Radiculopathy
- Osteoarthritis
- Herpes zoster
- Diabetic neuropathy
- Failed back surgery
- Oncological
- Other diseases with chronic pain

COMPOSITION

- Palmitoylethanolamide

SCIENTIFIC SUPPORT

In a large observational study involving 610 patients with various chronic pain etiologies, PEA significantly reduced mean pain intensity without serious adverse events. Furthermore, a pooled meta-analysis of 12 clinical studies demonstrated that PEA produced a progressive and statistically significant reduction in pain intensity, with effects approximately five times greater than control, and 81% of treated patients achieving a pain score ≤ 3 after 60 days of treatment. These findings support PEA as a safe and effective non-opioid option for chronic pain management.



Pain Physician 2016; 19:11-24 • ISSN 1533-3159

Palmitoylethanolamide, a Special Food for Medical Purposes, in the Treatment of Chronic Pain: A Pooled Data Meta-analysis

Antonella Paladini, MD, PhD1, Mariella Fusco, PhD2, Teresa Cenacchi, MD, PhD3, Carlo Schievano, PhD4, Alba Piroli, MD, PhD5, and Giustino Varrassi, MD, PhD6

Conclusion:

- PEA leads to a progressive reduction of pain intensity.
- PEA produced 5x higher effects compared to control group.





PAIN RELIEF
Food supplement

Onaceron® forte

Pharmaceutical form:

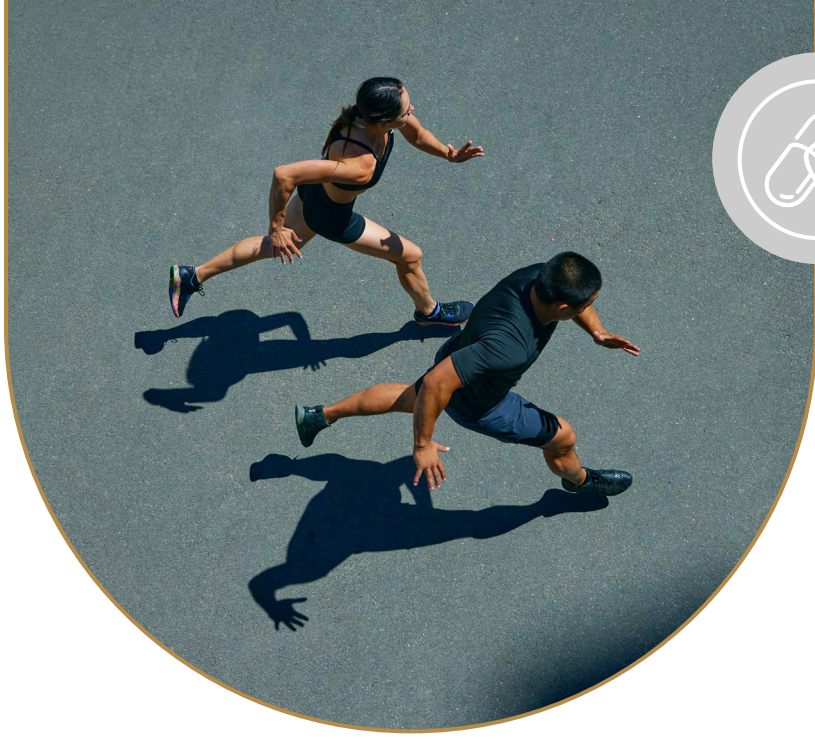
capsules

Packaging:

90 capsules

Recommended dosage:

3 capsules a day



INDICATED FOR

- Rheumatism
- Osteoarthritis
- Rheumatoid arthritis
- Sciatica and back pain
- Lumbago
- Swelling of the joints
- Inflammation and stretched ligaments
- Muscle contusions
- Sports injuries

COMPOSITION

- Glucosamine hydrochloride
- Chondroitin sulfate
- Mobilee® (standardized natural HA complex)
- Vitamin C

SCIENTIFIC SUPPORT

Onaceron Forte® demonstrated significant chondroprotective and anti-inflammatory effects in preclinical studies. In vitro, it reduced cartilage-degrading enzymes (MMP-1 and MMP-13) and increased glycosaminoglycan (GAG) synthesis in osteoarthritic chondrocytes, showing stronger effects than the standard glucosamine/chondroitin combination. In vivo, it reduced pro-inflammatory cytokines (IL-6 and TNF-α) and decreased arthritis severity in a collagen-induced arthritis model, supporting both anti-catabolic and anabolic activity in joint tissue.

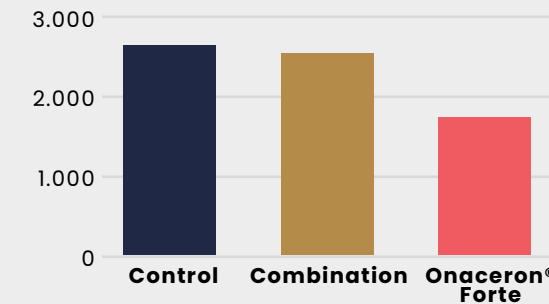
KEY ADVANTAGES

The synergistic combination of ingredients such as glucosamine hydrochloride, chondroitin sulphate, mobilee®, vitamin C, acts comprehensively on all joint structures, slowing down the progression of the disease.

Onaceron® Forte reduces cartilage-degrading enzymes

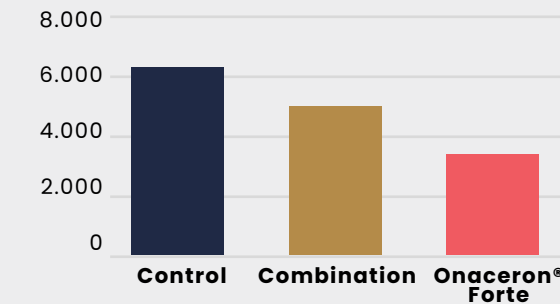
ONACERON FORTE® showed a significant reduction in the activity of MMP-1 and MMP-13 in OA where chondrocytes were stimulated by IL-1β.

MMP- 1 activity



MMP - 1 activity in the presence of IL 1B

MMP- 13 activity



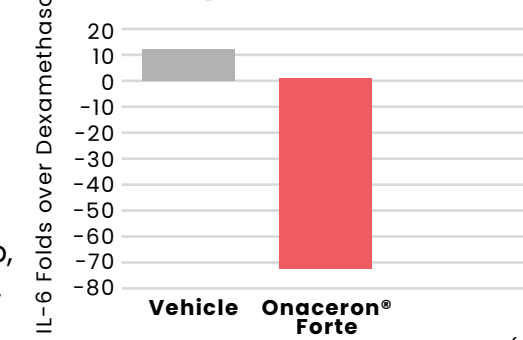
MMP - 13 activity in the presence of IL 1B

The effect was stronger compared to the standard combination (1500 mg glucosamine sulphate and 1200 mg chondroitin sulphate).

Onaceron Forte® reduces joint inflammation

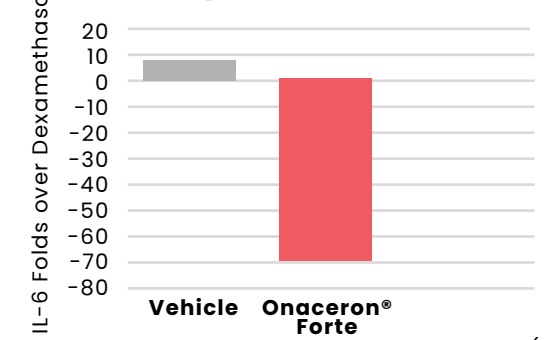
ONACERON FORTE® significantly reduced the production of pro-inflammatory cytokines IL-6 and TNFα compared to the control group, in an animal model of arthritis (rats, collagen-induced arthritis – CIA).

IL- 6 production



The results are shown in percentage (%)

TNFβ production



The results are shown in percentage (%)

CLICK **FOR MORE DETAILED PRESENTATION**



PAIN RELIEF
Medical Device

Rheial

Pharmaceutical form:

Prefilled syringe

Packaging:

1 prefilled syringe

Recommended dosage:

Once a month or a year depends on the type of product

WATCH NOW



INDICATED FOR

Substitute of synovial fluid into joints affected by degenerative or mechanical OA that gives pain or reduced mobility

COMPOSITION

- | | |
|-----------------------------|-----------------------------|
| HA (hyaluronic acid) | HA (hyaluronic acid) |
| • 20 mg / 2 ml | • 60 mg / 2 ml |
| • 32 mg / 2 ml | • 60 mg / 3 ml |
| • 40 mg / 2 ml | • 72 mg / 3 ml |
| • 48 mg / 2 ml | • 90 mg / 3 ml |

SCIENTIFIC SUPPORT

Retrospective single-center study included 60 patients with Kellgren–Lawrence Grade 2 and 3 gonarthrosis and compared the clinical outcomes of a single intra-articular injection of cross-linked hyaluronic acid with a standard three-dose regimen of linear hyaluronic acid. Evaluations were performed at baseline, and at three and six months post-injection using the WOMAC index and the Oxford Knee Score. Both treatment groups demonstrated statistically significant improvement compared to baseline.

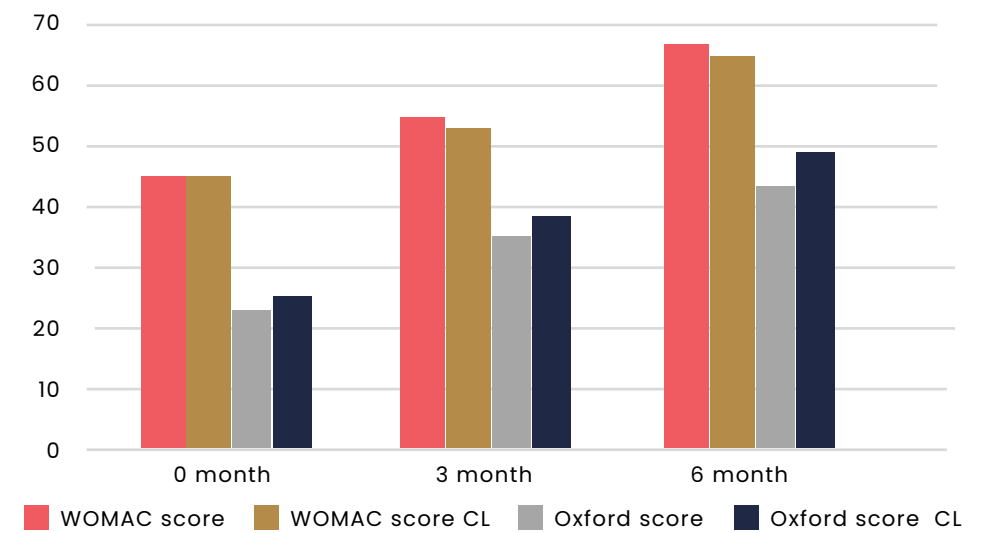
KEY ADVANTAGES

Rheial offers a wide range of HA concentrations and volumes for all types of joints and patients.

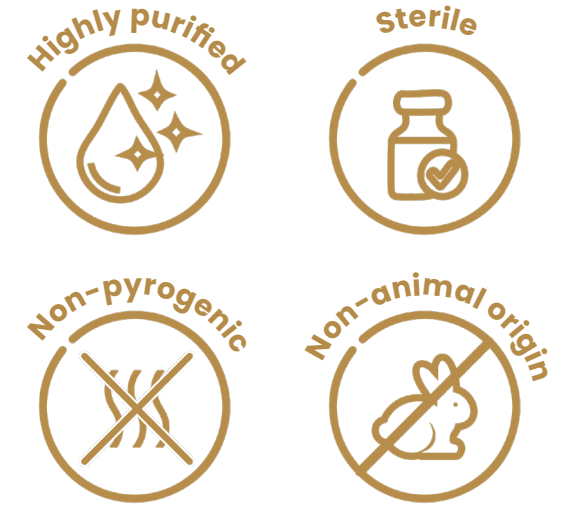
- **Linear Hyaluronic acid**
- **Cross-linked Hyaluronic acid**

Cross-linked hyaluronic acid is longer-lasting and more stable compared to non-cross-linked or natural hyaluronic acid.

Standard / Cross-linked HA injection



IT CONTAINS HA FROM BIO FERMENTATIVE ORIGIN



A comparison of the functional outcomes was conducted resulting from two distinct injections:

a 2 mL dose of Rheial 2.4% 48 mg cross-linked HA vs Rheial 2 mL dose of 2.4% 48 mg linear HA knee injection.

CLICK FOR MORE DETAILED PRESENTATION



PEDIATRIC
Medical Device

Otines Plus

Pharmaceutical form:

Spray ear drops

Packaging:

7 ml bottle

Recommended dosage:

1-2 short sprays into the ear 3 times a day



INDICATED FOR

- Otitis media
- Otitis externa
- Ear pain
- Removing excess ear wax (cerumen)

COMPOSITION

- Olive oil (*Olea Europaea*)
- Lavender oil (*Lavandula angustifolia*)
- Vitamin E
- Rosemary extract (*Rosmarinus Officinalis*)
- St. John's Wort extract (*Hypericum Perforatum*)
- Mullein extract (*Verbascum Thapsus*)
- Pot marigold extract (*Calendula Officinalis*)
- Garlic oil (*Allium sativum*)

SCIENTIFIC SUPPORT

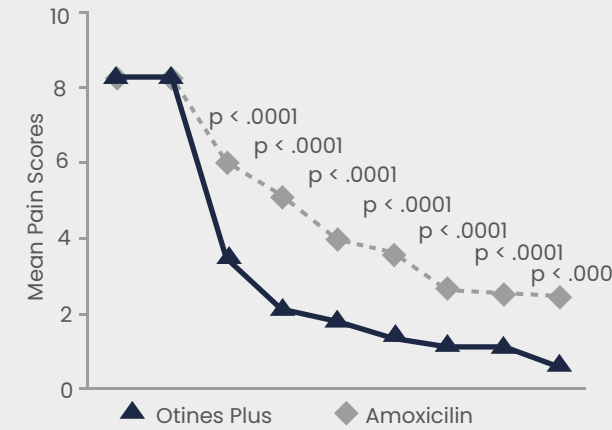
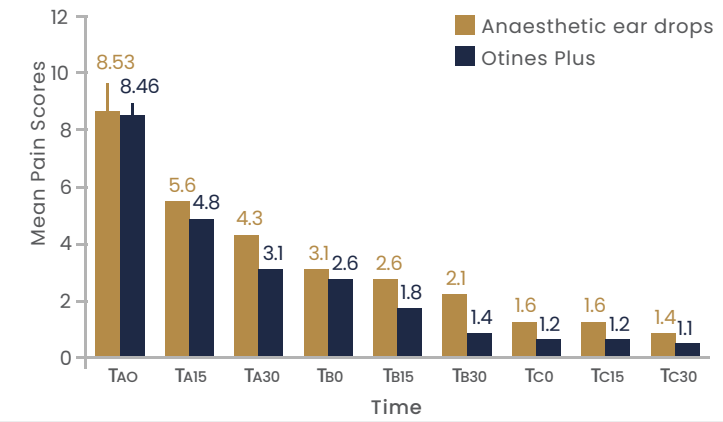
Otines Plus has been demonstrated to be effective in clinical trials in hospitals and in clinics in three studies conducted by pediatricians. **474 children participated in the three studies**, and the results were statistically significant. Otines Plus has been tested and found to be effective in children in comparison with an antibiotic as common as Amoxicillin, and also in comparison with anesthetic ear drops.

KEY ADVANTAGES

Otines plus is a class II medical device in the form of ear drops for spray application and contains natural ingredients used to relieve ear pain, remove excess ear wax (cerumen) and improve hearing.

Efficacy in the Management of Ear Pain Associated With Acute Otitis Media

Otines Plus, an ear drop formulation of naturopathic origin, is as effective as anaesthetic ear drops and was proven appropriate for the management of AOM-associated ear pain.



The conclusion of the study is that the efficacy of treatment of otitis media with natural ear drops is identical to the efficacy of antibiotic treatment of the otitis, and as regards reduction of otitis-caused pain, the efficacy of natural ear drops (Otines Plus) is significantly higher.

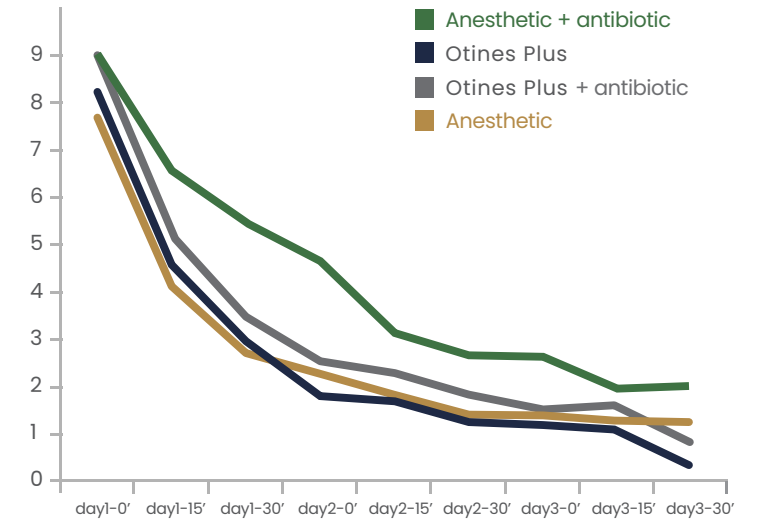
Treatment of acute otitis media: Efficacy compared to antibiotic therapy

CLICK **FOR MORE DETAILED PRESENTATION**

Treatment of ear pain in children

Patients treated with natural ear drops responded better than those using antibiotics.

The best results came from natural drops alone. The study concluded they are effective for AOM ear pain, with no added benefit from antibiotics.





PEDIATRIC
Medical Device

Effigerm Nappy Care

Pharmaceutical form:

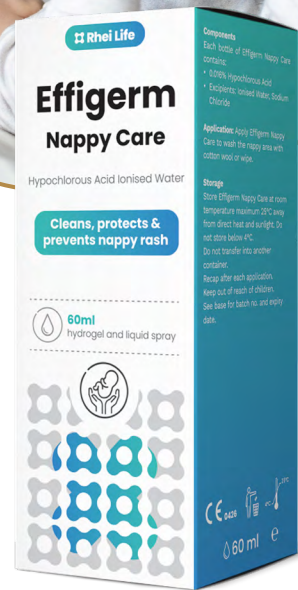
Hydrogel and liquid spray

Packaging:

60ml, 150ml, 250ml bottles with finger spray

Recommended dosage:

Apply as required.



COMPOSITION

- Hypochlorous acid (HOCl) 0,016 %

SCIENTIFIC SUPPORT

HOCl has been widely evaluated by regulatory bodies, healthcare institutions, and academic organizations in over 50 countries, with more than 2,600 peer-reviewed publications supporting its antimicrobial efficacy and over 3,000 publications addressing its role in wound healing enhancement. There are **no reports of adverse reactions to topical applications** by these methods based on the US EPA's Toxicology Database DSSTox, the US CDC Toxic Substances and Disease Registry, nor at either the Development and Reproductive Toxicology Database, or the European Bioinformatics Institute of EMBL.

KEY ADVANTAGES

Effigerm Nappy Care is a topical irrigation solution, its key mode of action is to modify the environment by mechanical removal of foreign bodies that may cause skin irritation and infection and to provide an optimal environment with regards to moisture, humidity, oxygen, and pH to assist in the healing process.

Liquid Formulation and Hydrogel Formulation – **2 STEP APPROACH:**

Step 1:

It is recommended to use the **liquid formulation** to wash the nappy area with cotton wool or wipe.



Step 2:

The **hydrogel formulation** is recommended to be used after cleansing and prior to reapplying the fresh nappy.



Both formulations allow the skin to breath and do not block pores or the nappy.



Hygienically Clean



Skin-friendly pH



Alcohol-Free



Dermatologically Tested



Hypoallergenic



Non-irritant



Suitable from day 1

CLICK FOR MORE DETAILED PRESENTATION



PEDIATRIC Food supplement

Kids gummies

Pharmaceutical form:

gummies

Packaging:

60 gummies

Recommended dosage:

2 gummies a day



KIDS GUMMIES

- Multivitamin
- Omega complete
- Fe+ iron support
- Vitamin D, Calcium
- Super immune booster
- Ultra omega complete
- Appetite booster
- K2+Ca+D3 bone booster
- Eye care
- Sleep aid

CLICK FOR MORE DETAILED PRESENTATION



HEALTH & WELLBEING Food supplement

Adult gummies

Pharmaceutical form:

gummies

Packaging:

60 gummies

Recommended dosage:

2 gummies a day



KEY ADVANTAGES

Gummy formulas are highly popular delivery methods for nutritional supplements, favored by children and adults as well. We produce carefully designed gummy formulas to meet all of our customer's nutritional supplement needs. In addition to high quality and attractive taste, our gummies are pectin based, manufactured halal, gelatin free, peanut free, and gluten free to satisfy our customers with dietary restrictions or sensitivities.

ADULT GUMMIES

- ### INDICATED FOR
- Super hair booster
 - Healthy joint support
 - Healthy bowel movement
 - Sore throat relief
 - Sleep well
 - Body cells & nerve functions

CLICK FOR MORE DETAILED PRESENTATION

CUSTOMER
REACH:

80 CUSTOMERS *from*
47 COUNTRIES



Europe (17)

Bosnia and Herzegovina
Croatia
Czech Republic
Latvia
Lithuania
Malta
Montenegro
Netherlands
North Macedonia
Poland
Romania
Russia
Serbia
Slovakia
Slovenia
Switzerland
Ukraine

Asia (13)

Azerbaijan
Iraq
Jordan
Kazakhstan
Kuwait
Malaysia
Oman
Philippines
Saudi Arabia
Sri Lanka
United Arab Emirates
Uzbekistan
Vietnam

Africa (3)

Egypt
Libya
South Africa

The Americas (14)

Bolivia
Colombia
Costa Rica
Dominican Republic
Ecuador
El Salvador
Guatemala
Honduras
Mexico
Nicaragua
Panama
Paraguay
Uruguay
Venezuela

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 Rhei Life

Our
Portfolio

