





REPARATIVE PHASE

HYALURONIC ACID SODIUM SALT 2,0% FOR INTRADERMAL INJECTION

INTRODUCTION

PROAGE is the latest generation of hyaluronic acid made to repair the extracellular matrix (ECM) by aging and restore the physiological function of the skin.

PROAGE reactivates the fundamental structure of the extracellular matrix in order that the cells involved in regeneration phase of new fibers can find the physiological functional relationships.

PROAGE increases the biosynthetic capacity of fibroblasts, stimulates cellular activity, and the synthesis of collagen, elastin and HA (hyaluronic acid), delays significantly the chronic skin photo-Aging.

PROAGE thanks to its effectiveness reorganize the ECM and bio-revitalizes dehydrated skin and damaged by the aging and sun exposure.

Can be injected before the use of fillers intradermic. It improves the density of the skin through the restructuring of the extracellular matrix.

PROAGE promotes the activity of the fibroblasts with formation of new collagen and elastin fibres.

DESCRIPTION

PROAGE is a sterile, biodegradable and isotonic intradermal gel. It consists of a medium molecular weight hyaluronic acid (1,5-2,0 x 10⁶ Dalton), obtained from Streptococcus Equi bacteria, formulated at a concentration of 20 mg/ml in a physiological buffer.

ACTIONS

- Deeply hydrates
- Breaks down free radicals
- · Limits the actinic keratosis
- Prolongs the neo angiogenesis

INDICATIONS

PROAGE is indicated in the physiological process of skin aging, the effects of which include the thickening of the stratum corneum and the alteration of the elastic fibers of the dermis in cases of cicatrisation resulting from superficial skin lesions (e.g., scars from acne and chicken pox).







TECHNICAL PRODUCT SHEET

PRODUCT DESCRIPTION

Sterile, biodegradable and isotonic intradermal gel, innovative Extracellular Matrix substitute, formulated at a concentration of 20 mg/ml in a physiological buffer. Pre-filled syringe containing 2 ml of non-pyrogenic gel, sterilized using moist heat.

	TECHNICAL C	CHARACTERISTICS	OF THE F	PRODUCT	
ACTIVE INGREDIENT		Sodium Hyaluronate 20 mg/ml with a molecular weight of 1,5 Million Daltons and 2 Million Daltons not crosslinked.			
EXCIPIENTS		Sodium monobasi chloride, water for		e, sodium dib	pasic phosphate, sodium
PRIMARY PACKAGING MATERIAL		Crystal syringe in b	tal syringe in blister VOLU		IE 2 ml
SECONDARY PAC	CKAGING MATERIAL	Cardboard			
COLOR		Transparent	ST	ERILITY	Sterile
LAW AND DIRECTIVES APPLIED		Medical Device Class III designed as Directive 92/43/CEE, manufactured in accordance with ISO 13485 requirements and regulation (EU) 2017/745.			
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PRODUCT PURP	POSE	skin, whose effec	ts include ecreased sk	thickening o	Il ageing process of the f the stratum corneum and the alteration of the
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