facilitation of mucociliary regeneration PureRegen® gel Sinus for adhesion prevention and



Order information

Sterile gel (5cc) in glass syringe, delivery cannula	40-011-003	PureRegen® gel Sinus
Sterile gel (2cc) in glass syringe, delivery cannula	40-011-001	PureRegen® gel Sinus
Description	REF Code	Product name

Developed and manufactured by:

Distributed by:

Address and contact info:

M BIOREGEN

BioRegen Biomedical Co., Ltd.

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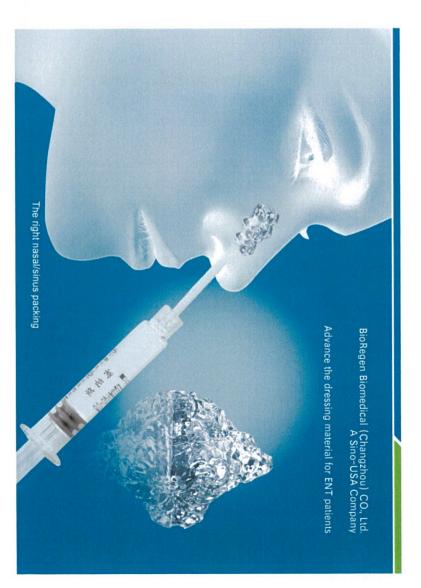
PureRegen® Gel SINUS

Self-crosslinked Hyaluronic Acid Gel Dressing Viscous and Controlled Degradation: Balanced to match the

tissue regeneration critical period

- Facilitates re-mucosalization acting as an adjunct
- Eliminates post painful removal
- Avoids secondary damage to the regenerated mucous
- Conforms to nasal cavity without dead space
- Prevents adhesion and scarring

Minimize bleeding and edema

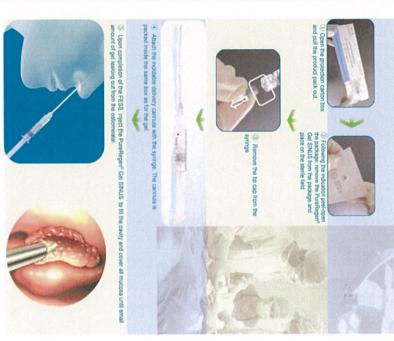






- proteins, excellent biocompatibility Proprietary self-crosslinking technique: no toxic residual and xenogenic
- natural hyaluronan. Gentle material modification method; preserved biological function of
- Adequate viscosity and hydrophilic property: creation of moist environ-
- Controlled degradation profile: physical barrier during the mucous regenment facilitating re-mucosalization
- eration critical period to prevent adhesion and scar formation.
- Terminal sterilization of the gel: maximized safety after application
- Dissolvable hydrogel: reduced discomfort to the patients and no need to

Instruction for Use:





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adhesion and facilitate mucociliary regeneration: Proven efficacy of PureRegen® Gel SINUS to prevent

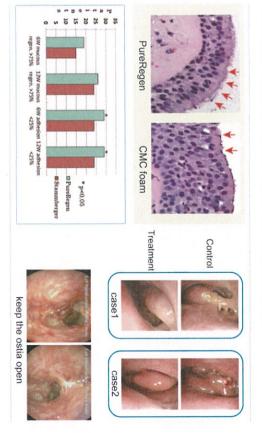
Controlled degradation profile:

removal and secondary damage to the fresh SINUS eliminates the need for painful packing complete dissolution of PureRegen® Gel mucosal surfaces and prevent adhesion. The providing effective barrier between the inflammatory processes after surgery, thereby matches the critical repair period, esp. the PureRegen® Gel SINUS degradation profile



reaction time during the mucous healing (1,2). to14 days, which matches the critical inflammatory PureRegen® Gel SINUS completely dissolves up

cosalization, and reduced edema and crust than the control group (3,4,5). USA demonstrated that PureRegen® Gel SINUS resulted in fewer adhesion, better re-mu-Prospective, randomized, controlled multiple center clinical studies (Level I) in China and



- 1. Watelet JB, et al: Wound healing of the nasal and paranasal mucosa: A review. Am. J. Rhinol. 16:77-84, 2002
- Endoscopic Surgery, Prof. Cornel Lancu (Ed.), ISBN: 978-953-307-717-8, InTech. Chapter 21; page 419-436, 2011 Pajic-Penavic I.: Endoscopic monitoring of postoperative sinonasal mucosa wounds healing. Advances in
- study. Am. J. Rhinol. Allergy. 27:71-76, 2013. Shi R, et al: The clinical outcomes of new hyaluronan nasal dressing: A prospective, randomized, controlled
- domized, controlled trial. Am J Rhinol Allergy 28, 508-513, 2014 Matheny KM, et al.: Self-crosslinked hyaluronic acid hydrogel (PureRegen® Gel Sinus) in ethmoidectomy: A ran-
- 5. Preventing adhesion and keeping the ostia open after functional endoscopic frontal sinus surgery by a novel crosslinked hyaluronan gel: Internal data. Manuscript 2015.

PureRegen® Gel Sinus

Novel crosslinked hyaluronan hydrogel

Advanced anti-adhesion barrier with the potential to regenerate mucous tissue

▲ Matheny KE et al. Self-cross-linked hyaluronic acid hydrogel in ethmoidectomy: A randomized, controlled trial. Am J. Rhino. Allergy, 29: 508, 2014.

▲ Shi R et al. The clinical outcomes of new crosslinked hyaluronan nasal dressing: Prospective, randomized, controlled trial. Am J. Rhino. Allergy, 27: 71, 2013.

- Stop minimal bleeding
- Prevent post-OP adhesion
- Promote functional tissue regeneration
- Cost effective
- No mixture needed, user-friendly
- Clinically proven efficacy
- Degradable and restorable without the need to remove
- Moldable delivery cannula to assist gel installation

