OSSIX® VOLUMAX is a resorbable collagen membrane intended for use during the process of guided tissue and bone regeneration. It is produced by standardized and controlled manufacturing procedures.

The collagen is extracted from porcine tendons subjected to veterinarian inspection and is purified to prevent hypersensitivity reactions.

OSSIX® VOLUMAX is packaged in a sealed double blister contained in a cardboard box and is terminally sterilized by ethylene oxide (EtO).

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OSSIX® VOLUMAX is a resorbable collagen membrane intended for use during the process of guided bone regeneration (GBR) and guided tissue regeneration (GTR) as a biodegradable barrier for:

- soft tissue regeneration
- GTR
- Xenograft, osteoconductive and/or inductive bone substitute, or a mixture of these.

OSSIX® VOLUMAX does not dissolve or disintegrate when wet. An animal study has shown that OSSIX® VOLUMAX degradation is completed within approximately 6 months.

INDICATIONS
OSSIX® VOLUMAX is a resorbable collagen membrane intended for use during the process of guided bone regeneration (GBR) and guided tissue regeneration (GTR) as a biodegradable barrier for:

1. Ridge augmentation for later implant insertions.
2. Simultaneous ridge augmentation and implant insertions.
3. Ridge augmentation around implants inserted in delayed extraction sites.
4. Ridge augmentation around implants inserted in immediate extraction sites.
5. Alveolar ridge preservation consequent to tooth (teeth) extraction(s).
6. Over the window in lateral window sinus elevation procedures.
7. In implants with vertical bone loss due to infection, only in cases where satisfactory debridement and implant surface disinfection can be achieved.
8. In intra bony defects around teeth.
9. For treatment of recession defects, together with coronally positioned flap.
10. In furcation defects in multi rooted teeth.
11. Localized gingival augmentation.

CONTRAINDICATIONS
OSSIX® VOLUMAX must not be used in:

1. Patients with known collagen hypersensitivity.
2. Patients with sensitivity to porcine-derived materials.
3. Patients suffering from autoimmune diseases and connective tissue diseases, such as: systemic lupus erythematosus, dermatomyositis etc.

WARNINGS AND PRECAUTIONS
1. OSSIX® VOLUMAX is intended for a single use. Do not re-sterilize OSSIX® VOLUMAX.
2. Treatment of high risk patients, such as: smokers, patients suffering from uncontrolled diabetes mellitus, and uncontrolled periodontal disease may be impaired.
3. The safety of treatment with OSSIX® VOLUMAX in pregnant and nursing women and in children has not been yet established.
4. The outcome of regenerative procedures may be impaired in patients suffering from untreated periodontitis. Infection control and good oral hygiene should be achieved prior to surgical intervention.

ADVERSE EVENTS
1. Post marketing experience with OSSIX® PLUS, which is a thinner version of the membrane, reveals an excellent safety profile.
2. Adverse reactions with OSSIX® PLUS collagen membrane were not observed.
3. Yet, as the membrane is of a collagen origin, allergic reactions (e.g. erythema, swelling, induration and/or pruritus at treatment site) may not be entirely excluded.

DIRECTIONS FOR USE
1. Special instructions for use in periodontology
   A basic requirement for successful periodontal treatment includes eradicating the underlying bacterial infection as well as adequate oral hygiene. Therefore, prior to surgical intervention, patients must receive a hygiene phase of treatment, consisting of oral hygiene instructions, scaling and root planing, and occlusal adjustment when indicated. A postoperative maintenance phase can help to ensure long-term therapeutic success.
2. The bony defect should be exposed by full thickness mucoperiosteal flaps.
3. All soft tissues should be removed.
4. In GTR the root surface should be carefully debrided and planed. Root conditioning should be considered.
5. The cortical plate can be perforated in order to allow osteogenic tissues from the bone marrow to colonize the regenerating site.
6. By using sterile a-traumatic instruments and sterile gloves rinsed with sterile saline, OSSIX® VOLUMAX is removed aseptically from the package.
7. OSSIX® VOLUMAX should be immersed (the inner blister can be used) for 30 seconds in sterile saline, to allow for its expansion to its final dimensions (10x12.5, 15x25 mm, 25x30 mm and 10x40 mm). Initial trimming to the estimated final size may be performed prior to sterile saline immersion.
8. Trimming to the required dimensions: it is recommended that OSSIX® VOLUMAX extends 3-4 mm beyond the margins of the defect. One-two mm of uncovered bone to adjacent teeth must be allowed.
9. OSSIX® VOLUMAX is cut with sterile scissors over a sterile container; trimming to the required dimensions: it is recommended that OSSIX® VOLUMAX extends 3-4 mm beyond the margins of the defect. One-two mm of uncovered bone to adjacent teeth must be allowed.
10. The site to be augmented should be filled with a space-maintaining material. The user should follow the manufacturer’s instructions for the material used.
11. OSSIX® VOLUMAX must not be used in:
   1. Patients with known collagen hypersensitivity.
   2. Patients with sensitivity to porcine-derived materials.
   3. Patients suffering from autoimmune diseases and connective tissue diseases, such as: systemic lupus erythematosus, dermatomyositis etc.

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The outcome of regenerative procedures may be impaired in patients suffering from untreated periodontitis. Infection control and good oral hygiene should be achieved prior to surgical intervention.
12. The mucoperiosteal flaps are sutured while ensuring that the tissue is not under tension. Do not compromise blood supply to the defect area.
13. In GTR, the use of a periodontal dressing may be considered.

GUIDELINES FOR THE PATIENT

The success of any surgical treatment depends on fulfilling the directions for use along with guiding the patient, as follows:

1. Preoperative patient’s education regarding adequate oral hygiene and meticulous prophylaxis.
2. Postoperative patient’s care, e.g.:
   a. Soft diet, avoidance of contact with tongue, hard food or denture.
   b. Avoidance of contact with hot temperature food or liquids that may cause early disintegration of the collagen matrix.
   c. Following suture removal, rinsing with chlorhexidine for one minute twice a day or according to the manufacturer’s instructions.

POSTOPERATIVE REMINDERS

1. Clinical experience with OSSIX® PLUS, which is a thinner version of the membrane, reveals no inflammatory signs following accidental exposure. The membrane degrades slowly in the oral environment and the exposed area is covered by connective tissue and epithelium within a few weeks (see references a-h below).
2. Possible complications with any surgery in the oral and maxillofacial region include: infection, flap slough, perforation, abscess formation, bone loss, pain, soft tissue irregularities, and complications associated with the use of anesthesia.
3. Depending upon the type and severity of the complications, as judged by the dental surgeon, membrane removal may be indicated.

STORAGE AND HANDLING

1. OSSIX® VOLUMAX should be used by skilled, experienced and/or trained dental surgeons.
2. The material should be handled using sterile gloves or sterile atraumatic instruments.
3. Placement of OSSIX® VOLUMAX should be performed after membrane’s immersion in saline for 30 seconds.
4. Do not use the membrane if it is torn and/or damaged.
5. Do not use the membrane, in the event that the sterile packaging is opened and/or damaged.
6. Any remaining / unused membrane should be discarded according to local regulations.
7. OSSIX® VOLUMAX should be stored at temperatures between 15-30°C (59-86°F).
8. Do not use the membrane after the expiration date.

HOW SUPPLIED

1. OSSIX® VOLUMAX is supplied in a double blister pack, for single use only. Each pack contains one membrane.
2. OSSIX® VOLUMAX is available in four sizes: 10x12.5 mm, 15x25 mm, 25x30 mm and 10x 40 mm.

REFERENCES


Caution: Federal (USA) law restricts this device to sale by or on the order of a licensed health care professional.