





### Science • Technology • Innovation

At Collagen Matrix we are passionate about advancing the science of tissue repair and regeneration. That's why we're the driving force in the design, development and manufacturing of advanced collagen and mineral based medical devices that support the body's natural ability to regenerate.

Over our 20 years of proven performance, we have focused our proprietary technologies and innovative products to meet clinical needs through five key business units – Dental, Spine, Orthopaedic, Dural Repair and Nerve Repair.

### **Proven Performance**

Six Platform Technologies

We have developed six proprietary tissue engineering technologies to expand our broad line of collagen and mineral based medical device solutions.

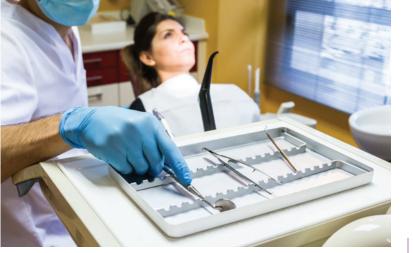
- T1 Reconstituted Collagen
- T2 Intact Collagen
- T3 Natural Carbonate Apatite Mineral
- T4 Collagen and Mineral Composites
- T5 Collagen Coatings
- T6 Crosslinking

\*

### **Evolution of Dental Products**

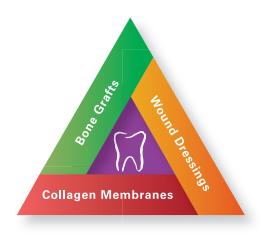






# 7.5 million

Our products have helped patients worldwide with over 7.5 million medical devices that have been produced across all five key business units.



### **Collagen Membranes**

Matrixflex<sup>™</sup>, MatrixDerm<sup>®</sup>, MatrixDerm<sup>®</sup> EXT,
MatrixDerm<sup>®</sup> Cap, and MatrixMem<sup>™</sup> membranes are
intended for use in oral surgical procedures as
resorbable materials for use in augmentation
around implants placed in immediate extraction
sockets, delayed extraction sockets, localized ridge
augmentation for later implantation, alveolar ridge
reconstruction for prosthetic treatment, filling of
bone defects, guided bone regeneration in
dehiscence defects and guided tissue regeneration
procedures in periodontal defects.

### **Wound Dressings**

Collatene™ Microfibrillar Collagen and MatrixDerm® Dental Wound Dressings are absorbent collagen matrices intended for the management of oral wounds and sores. They are indicated for use for denture sores, oral ulcers (non-infected or viral), periodontal surgical wounds, suture sites, burns, extraction sites, surgical wounds, and traumatic wounds.

### **Bone Grafts**

MatrixOss<sup>™</sup> Granules and MatrixOss<sup>™</sup> in Syringe, SynOss<sup>™</sup> Granules and SynOss<sup>™</sup> Putty deliver choices to clinicians by providing exceptionally high quality xenograft and synthetic alternatives to human bone graft materials.

Our products are indicated for use in bone repair such as augmentation or reconstructive treatment of the alveolar ridge and filling of periodontal defects. In addition, they are suitable for use in conjunction with products intended for Guided Tissue Regeneration and Guided Bone Regeneration and for filling defects after root resection, filling extraction sockets, and elevating the maxillary sinus floor.



# Membranes at-a-glance

#### Why use membranes?

- \* The membrane is used as a cell barrier to prevent soft tissue ingrowth in the defect site
- Contain bone graft material
- Support wound healing in guided tissue and bone regeneration procedures



#### Our collagen membranes are...

- Highly purified collagen derived from porcine tissue. Porcine animals are considered a non-TSE relevant species
- Highly biocompatible
- ❖ Programmed for different resorption times to accommodate various application
- ❖ Suturable with easy handling due to their structural integrity
- Not side specific, either side can be placed facing bone or soft tissue
- \* Easily repositionable to precise adjustment and placement. They do not stick to instruments
- \* Permeable to macromolecules and nutrients while creating an effective barrier to epithelial cells

Product	Material	Handling	Strength	Resorption
	Source	(conformability)	(suture pull avg)	Time
Matrix <i>flex</i> <sup>TM</sup>	Porcine peritoneum	High (very drapeable)	High 0.83 kg	3-4months
MatrixMem <sup>™</sup>	Porcine tendon	Moderate	Low 0.28 kg	4-6 months
MatrixDerm <sup>®</sup>	Porcine dermis	Moderate	Moderate 0.46 kg	6-9 months
MatrixDerm <sup>®</sup>	Porcine	Low	High	9-12 months
EXT	dermis	(space maintaining)	0.67 kg	
MatrixDerm <sup>®</sup>	Porcine	Low	Moderate	9-12 months
<i>Cap</i>	dermis	(space maintaining)	0.51 kg	



### **Membrane** characteristics

# The two key design parameters that we have identified in the development of the membranes are:

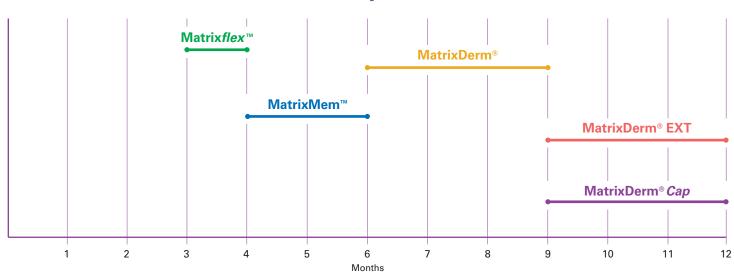
- Handling characteristics: Conformability vs. Stiffness
- Resorption time

This allows the clinician to select the most ideal membrane for the type of procedure.

### **Conformability**



### Resorption



#### Membranes can be used in the following types of dental surgeries:

- ❖ In augmentation around implants placed in immediate extraction sockets, delayed extraction sockets
- ❖ Localized ridge augmentation for later implantation
- \* Alveolar ridge reconstruction for prosthetic treatment
- Filling of bone defects
- Guided bone regeneration in dehiscence defects and guided tissue regeneration procedures in periodontal defects



# Matrix*flex*™



### REGENERATIVE COLLAGEN DENTAL MEMBRANE

Matrix*flex*™ regenerative collagen dental membrane is a strong, conformable collagen barrier membrane manufactured from purified porcine peritoneum tissue.

Matrix $flex^{\text{m}}$  is designed to provide a very drapable yet very strong membrane.

- \* Resorbable in 3-4 months
- High mechanical strength
- ❖ Highly purified intact porcine peritoneum tissue

CATALOG NO.	SIZE
CDMPP1520	15mm x 20mm
CDMPP2030	20mm x 30mm
CDMPP3040	30mm x 40mm











# **MatrixDerm®**



### REGENERATIVE COLLAGEN DENTAL MEMBRANE

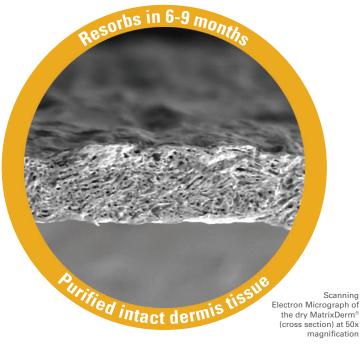
MatrixDerm® regenerative collagen dental membrane has enhanced characteristics to provide periodontal and dental surgeons with the ideal balance of properties to effectively address a host of clinical indications and surgical procedures. The product is manuafactured from porcine dermis tissue.

#### **Product Features:**

- \* Resorbable in 6-9 months
- Highly purified intact dermis tissue
- Balanced characteristics of handling, strength, and resorption time
- High mechanical strength

CATALOG NO.	SIZE
PDM1520	15mm x 20mm
PDM2030	20mm x 30mm
PDM3040	30mm x 40mm





\*





# **MatrixMem**<sup>™</sup>



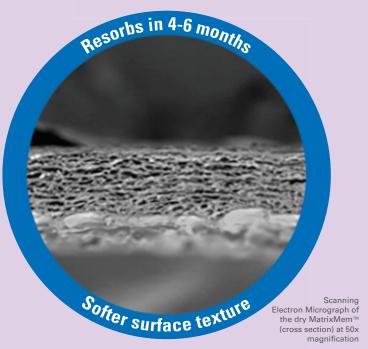
### REGENERATIVE COLLAGEN DENTAL MEMBRANE

MatrixMem<sup>™</sup> regenerative collagen dental membrane is a white, nonfriable, conformable membrane matrix engineered from highly purified type I collagen derived from porcine tendon.

- \* Resorbable in 4-6 months
- Highly purified type I collagen derived from porcine Achilles tendon
- Moderately drapeable
- ❖ Softer surface texture

CATALOG NO.	SIZE
CMPT1520	15mm x 20mm
CMPT2030	20mm x 30mm
CMPT3040	30mm x 40mm











# **MatrixDerm® EXT**



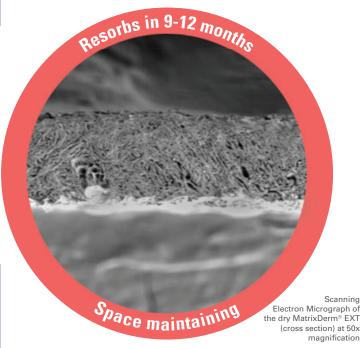
# REGENERATIVE COLLAGEN DENTAL MEMBRANE WITH EXTENDED RESORPTION TIME

MatrixDerm® EXT regenerative collagen dental membrane is a white, nonfriable membrane matrix engineered from highly purified porcine dermis.

- \* Resorbable in 9-12 months
- Highly purified intact thicker dermis tissue
- Longer in vivo stability
- Quick hydration time in seconds
- High mechanical strength

CATALOG NO.	SIZE
PDMX1520	15mm x 20mm
PDMX2030	20mm x 30mm
PDMX3040	30mm x 40mm











# MatrixDerm® Cap



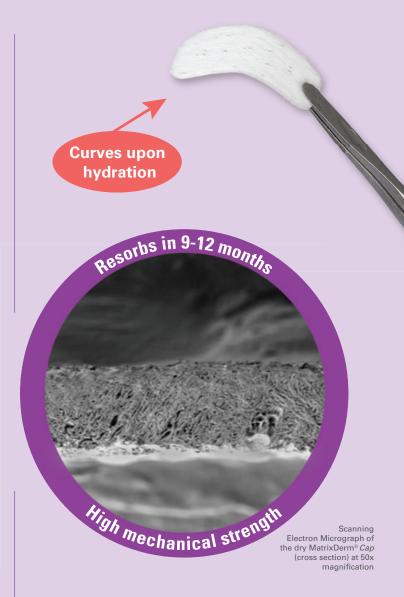
# REGENERATIVE COLLAGEN DENTAL MEMBRANE WITH EXTENDED RESORPTION TIME

MatrixDerm® *Cap* regenerative collagen dental membrane cap is a white, nonfriable membrane matrix engineered from highly purified porcine dermis. It is indicated for oral surgical procedures as a resorbable material for use in extraction sockets and small bone defects.

MatrixDerm® *Cap* was specifically designed to meet the needs for socket ridge preservation procedures, and it has a slight curve when hydrated to facilitate placement.

- \* Resorbable in 9-12 months
- Highly purified intact thicker dermis tissue
- Curves upon hydration
- Longer in vivo stability
- Quick hydration time in seconds
- High mechanical strength

CATALOG NO.	SIZE
PDMC1216	12mm x 16mm









# **Bone Grafts at-a-glance**

Collagen Matrix offers both a xenograft-derived bone graft matrix and a synthetic calcium phosphate bone graft matrix.





### Our bone grafts can be used for:

- \* Augmentation or reconstructive treatment of the alveolar ridge
- Filling of infrabony periodontal defects
- \* Filling of defects after root resection, apicoectomy, and cystectomy
- \* Filling of extraction sockets to enhance preservation of the alveolar ridge
- Elevation of the maxillary sinus floor
- ❖ Filling of periodontal defects in conjunction with products intended for Guided Tissue Regeneration (*GTR*) and Guided Bone Regeneration (*GBR*)
- ❖ Filling of peri-implant defects in conjunction with products intended for Guided Bone Regeneration (*GBR*)

Product	Material Source	Particle Size Range	Form/ Configuration	Volume Fill	Void Space (Porosity)
MatrixOss <sup>™</sup> Granules	Anorganic Porcine Bone Mineral (Cancellous)	0.25 - 1.0 mm	Granules	3 cc / gram	88%
MatrixOss™ Granules	Anorganic Porcine Bone Mineral (Cancellous)	1.0 - 2.0 mm	Granules	4 cc / gram	95%
SynOss™ Granules	Synthetic Carbonate Apatite	0.35 - 1.0 mm	Granules	2 cc / gram	81%
Syn0ss™ Putty	Synthetic Carbonate Apatite & Bovine Type I Collagen	0.35 - 1.0 mm	Putty	0.5cc, 1.0cc and 2.0cc	88%



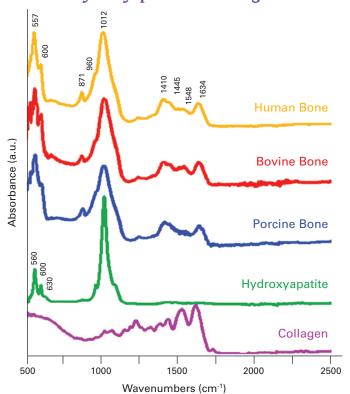
# Why Carbonate Apatite?



#### Mineral structure similar to natural bone mineral

♦ Naturally occurring bone mineral is a calcium phosphate apatite with 2-8 wt % carbonate ion substitution. Unlike highly crystalline hydroxyapatite, carbonate apatite in bone has a lower degree of crystallinity, which allows the bone mineral to be remodeled in vivo.<sup>(1,2,3)</sup>

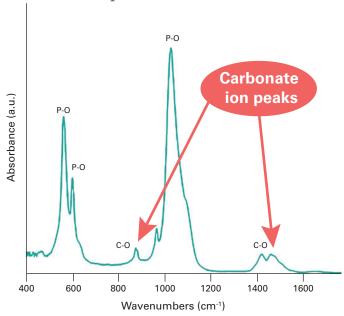
IR Spectra for human, bovine & porcine bones hydroxyapatite and collagen



❖ The figure above shows the IR spectra of the human, bovine and porcine bones, hydroxyapatite and collagen.<sup>(4)</sup> It reflects the identical chemical composition of human, bovine & porcine bones. They all show a carbonate ion peak at around 871 cm<sup>-1</sup> and 1410/1445 cm<sup>-1</sup> Our
proprietary method
to process bone tissue
preserves the natural porous
structure and maintains the
carbonate apatite
crystal structure. (5)

- ♦ After processing porcine bone tissue using our proprietary method, the IR of processed bone was similar to that of naturally occuring bone with carbonate band between 800 cm<sup>-1</sup> 900 cm<sup>-1</sup> and 1400 cm<sup>-1</sup> 1500 cm<sup>-1</sup>
- The material structure of carbonate apatite being similar to the mineral structure of natural bone, supports and facilitates remodeling of the new bone

IR Spectra for MatrixOss™



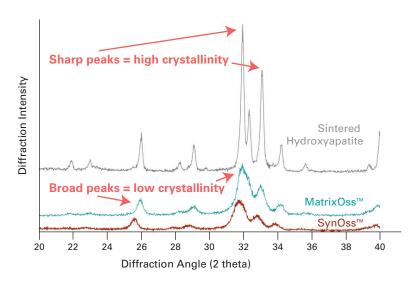


## 2

### Low crystallinity supports bioresorption

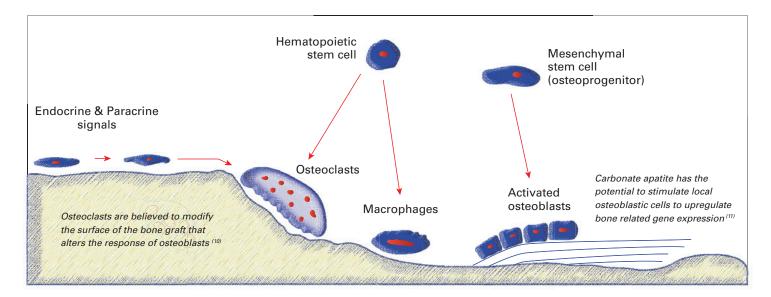
- The broad spectrum of the mineral indicates a smaller crystal size. Both diffraction patterns are close to mature native bone diffraction pattern that is consistent with bone mineral of apatite nature with low crystallinity. (6) Hyroxyapatite shows characteristically narrow spectrum with sharper peaks indicative of high crystallinity.
- Carbonate substitution leads to greater bioresorption by osteoclasts due to low crystallinity of the bone graft (3,7,8)

# X-ray diffraction for MatrixOss™, SynOss™ and Hydroxyapatite



\*

Carbonate apatite has been shown to increase the bone forming activities of **osteogenic** cells as well as enhance bioresorption of bone by **osteoclasts** (9)



- ❖ In-vivo the activities of osteoclasts and osteoblasts are closely related in a process known as coupling (10, 12, 13)
- ❖ The material structure and porosity (or void space) are two parameters that we have evaluated and optimized in the development of our bone graft matrices



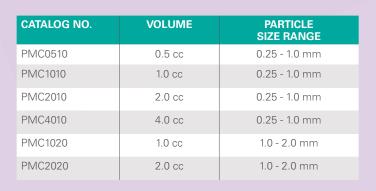
# **MatrixOss<sup>™</sup> Granules**

### ANORGANIC BONE GRAFT

MatrixOss™ Granules is an osteoconductive, porous, anorganic bone mineral with carbonate apatite structure derived from porcine cancellous bone

#### **Product Features:**

- Carbonate apatite structure similar to natural bone
- Highly porous
- Rough surface/faciliates cell adhesion
- ♦ High volume fill



#### Syringe

CATALOG NO.	VOLUME	PARTICLE SIZE RANGE
PMCS025	0.25 cc	0.25 - 1.0 mm
PMCS05	0.5 cc	0.25 - 1.0 mm



# Why **MatrixOss**™?

#### Safe

- Porcine animals are considered a non-TSE relevant species
- Bone tissue is subjected to several processing steps known to eliminate or inactivate viruses
- A rigorous process designed to effectively mitigate any risk of disease transmission and ensure safety for human implantation
- The bone graft is provided sterile and for single use only

#### **Carbonate apatite anorganic bone mineral**

- Carbonate apatite structures are better osteoconductive materials than hydroxyapatite 1,2,3
- Resorption and remodeling profiles are more similar to natural bone than those of synthetic materials, such as hydroxyapatite or tricalcium phosphate<sup>2</sup>



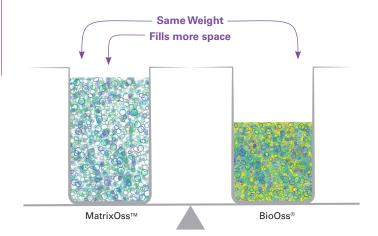
#### **Porous**

- Porosity permits vascularization of the defect site and enhances osteogenesis 14,15
- High porosity and large pores enhance bone ingrowth and osseointegration of the implant after surgery 15
- MatrixOss<sup>™</sup> macropores range between 0.1mm
   1.0mm <sup>16</sup>



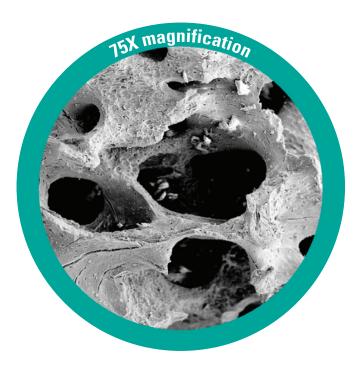
#### Volume Fill 16

- 1gram of small size particles fills approximately 34% more volume then Bio-Oss®
- ❖ 1gram of large size particles fills approximately 49% more volume then Bio-Oss®



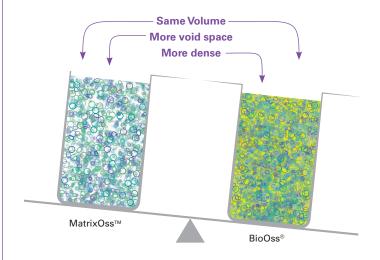
### **Surface Roughness** 17

Surface roughness affects cellular response, enhancing cell adhesion and proliferation and possibly other markers of expression of cell phenotype, like production of collagen type I, osteocalcin, extracellular matrix and mineralized material



### Void Space 16

- ❖ 88% void space for porcine mineral vs 78% void space of Bio-Oss® for small particles
- ♦ 95% void space for porcine mineral vs 88% void space of Bio-Oss® for large particles







# SynOss<sup>™</sup> Putty

### SYNTHETIC MINERAL-COLLAGEN COMPOSITE BONE GRAFT

SynOss™ Putty is a calcium phosphate-based mineral with a carbonate apatite structure similar to natural bone combined with type I collagen derived from bovine Achilles tendon. The mineral particles are dispersed within collagen fibers forming a 3-dimensional matrix. It is supplied dry and forms a moldable putty upon hydration. It is fully resorbed during the natural process of bone formation and remodeling.

- Carbonate apatite plus bovine type I collagen
- Moldable putty upon hydration
- Osteoconductive, 3 dimensional structure for forming cells
- Mineral particles are contained within the defect site

CATALOG NO.	VOLUME	SIZE
SMB050	0.5 cc	9.0mm x 8.0mm
SMB100	1.0 cc	11.0mm x 10.5mm
SMB200	2.0 cc	11.0mm x 21.0mm









# SynOss<sup>™</sup> Granules

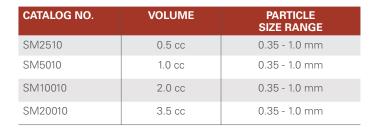
### SYNTHETIC MINERAL BONE GRAFT

SynOss™ Granules Synthetic Mineral Bone Graft is an osteoconductive calcium phosphate-based *(carbonate apatite)* bone graft material that provides clinicians and their patients with an ideal alternative to human allograft and animal origin bone graft material.

The similarity in structure between SynOss™ Granules and natural anorganic bone mineral allows the SynOss™ Granules in vivo resorption and remodeling profile to mimic that of natural bone.



- Synthetic alternative to allografts and xenografts
- Carbonate apatite structure similar to natural bone mineral
- Osteoconductive









# MatrixDerm® Tape, Foam and Plug

### RESORBABLE WOUND DRESSING

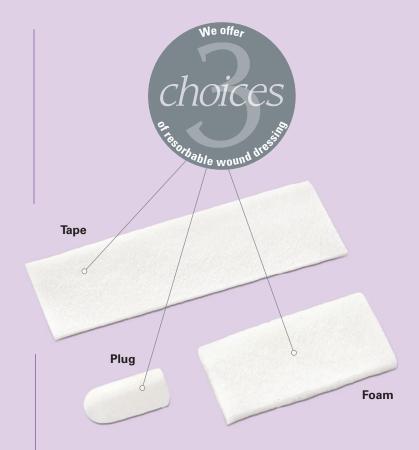
MatrixDerm® Resorbable Wound Dressings are absorbent, porous, collagen matrices engineered from purified collagen derived from bovine dermis tissue.

They are indicated for use for denture sores, oral ulcers (non-infected or viral), periodontal surgical wounds, suture sites, burns, extraction sites, surgical wounds and traumatic wounds.

- Applied directly to the wound and protect the wound and delicate new tissue
- Aids in wound healing
- Available in tape, foam and plug form
- Supplied sterile, non-pyrogenic and for single use only
- Essentially resorbs within 30 days of placement

CATALOG NO.	SIZE * Length x Width x Height
BDDWDT2575	2.5 cm x 7.5 cm x 1mm -Tape
BDDWDF2040	2.0 cm x 4.0 cm x 3.0 mm - Foam
BDDWDP1020	10 mm x 20 mm - Plug

<sup>\*</sup> Quantities of 10







# **Collatene**<sup>™</sup>

### MICROFIBRILLAR COLLAGEN WOUND DRESSING

Collatene™ Microfibrillar Collagen is an absorbent microfibrillar collagen matrix intended for the management of oral wounds and sores. It forms a gel when mixing with the wound's exudate and oral fluid to provide a moist healing environment.

It is indicated for use for denture sores, oral ulcers *(non-infected or viral),* periodontal surgical wounds, suture sites, burns, extraction sites, surgical wounds and traumatic wounds.

- Highly purified bovine type I collagen
- Applied directly to the wound and protects the wound bed and delicate new tissue
- Resorbable
- Collatene is supplied sterile, in individual vials and for single-use only
- Controls minor bleeding due to intrinsic hemostatic properties of intact collagen fibers



CATALOG NO.	VOLUME	QUANTITY
DWDG0010	0.1 gram	15 vials/box





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Matrixflex<sup>™</sup>, MatrixDerm<sup>®</sup>, MatrixDerm<sup>®</sup>, MatrixDerm<sup>®</sup> EXT MatrixDerm<sup>®</sup> Cap, MatrixOss<sup>™</sup> Granules, MatrixOss<sup>™</sup> Syringe, SynOss<sup>™</sup> Putty, SynOss<sup>™</sup> Granules, MatrixDerm<sup>®</sup> Tape, MatrixDerm<sup>®</sup> Foam, MatrixDerm<sup>®</sup> Plug and Collatene<sup>™</sup> are all trademarks of and manufactured by Collagen Matrix, Inc. Bio-Oss<sup>®</sup> is a registered trademark of Geistlich-Pharma, Switzerland. Please refer to the Instructions for Use for description, indications, contraindications, warnings, precautions and other important information. A surgeon must always rely on his or her own professional clinical judgement when deciding whether to use a particular product when treating a particular patient. Availability of these products might vary from one given

a particular patient. Availability of these products might vary from one given country or region to another as a result of specific local regulatory approval or clearance requirements for sale in such country or region.

