

bone & tissue
regeneration

botiss
biomaterials

maxresorb[®] & maxresorb[®] inject

Innovative biphasic calcium phosphate

Scientific and clinical evidence

synthetic

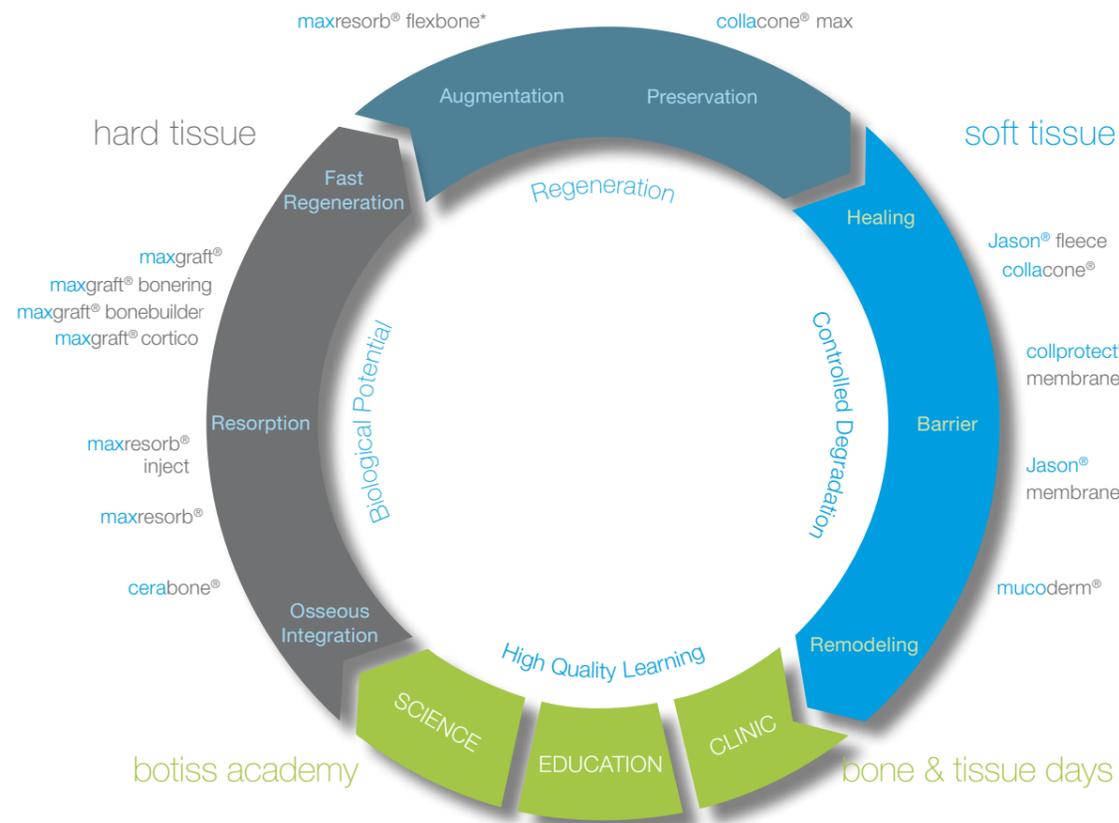
resorbable

safe

hard tissue



botiss regeneration system



Development / Production / Distribution

cerabone®	maxresorb®	maxresorb® inject	maxgraft® bonebuilder	maxgraft® bonering / maxgraft® cortico	maxgraft®
Natural bovine bone graft	Synthetic biphasic calcium phosphate	Synthetic injectable bone paste	Patient matched allogenic bone implant	Allogenic bone ring / Processed allogenic bone plate	Processed allogenic bone graft
maxresorb® flexbone*	collacone® max	Jason® fleece / collacone®	collprotect® membrane	Jason® membrane	mucoderm®
Flexible block (CaP / Collagen composite)	Alveolar cone (CaP / Collagen composite)	Collagenic hemostat (Sponge / Cone)	Native collagen membrane	Native pericardium GBR / GTR membrane	3D-stable soft tissue (Collagen) graft

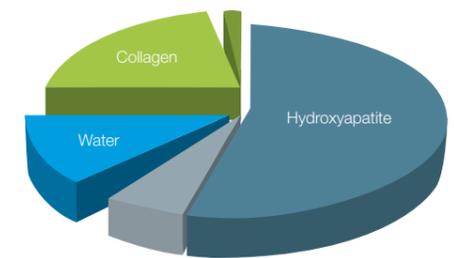
Bone

physical – chemical – biological

Bone is a highly specialized tissue with properties strongly adapted to its supporting and skeletal function. Bones are composed of ~65% inorganic matrix, the mineral phase, ~25% organic phase and ~10% water.

The main component of the mineral bone phase (~90%) is hydroxyapatite (biological apatite). This inorganic part is responsible for the high stability and compressive strength of the bone.

The organic phase mainly consists of collagen type I (~90%) being accountable for elasticity (tensile strength, ductility). Basis for bones' resistance to different mechanical forces is the interplay of collagen fibril, non-collagenous proteins and deposited mineral crystals.



Organic Substances	Inorganic substances
~90% Collagen	~90% Hydroxyapatite
~97% Collagen type I	~10% Magnesium
~3% Collagen type III	Sodium
~10% Amorphous basic substance	Iron
Proteins	Fluorine
Proteoglycans	Chlorine
Glycosaminoglycans	...
Lipids	



Femoral bone - outer cortical and inner cancellous bone clearly recognizable

Bone structure

Bones are constructed according to a lightweight principle; the structure enables a very high stability accompanied by a relatively low weight. The composition on the periphery is very compact (cortical bone, compacta), while the inner part is less densely structured with lattice-shaped bone trabeculae (cancellous bone).



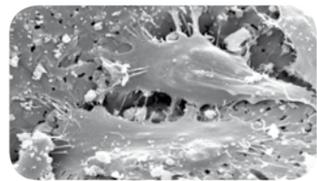
Human unicortical bone block

Bone biology and remodeling

communication of cells



Despite its high stability bone is in no way a rigid tissue, but is characterized by a high metabolism and is subject to constant *remodeling*. This dynamic is necessary to save the skeleton from degradation by the reparation of structural damages (micro fractures).



Active osteoblasts on bone substitute material

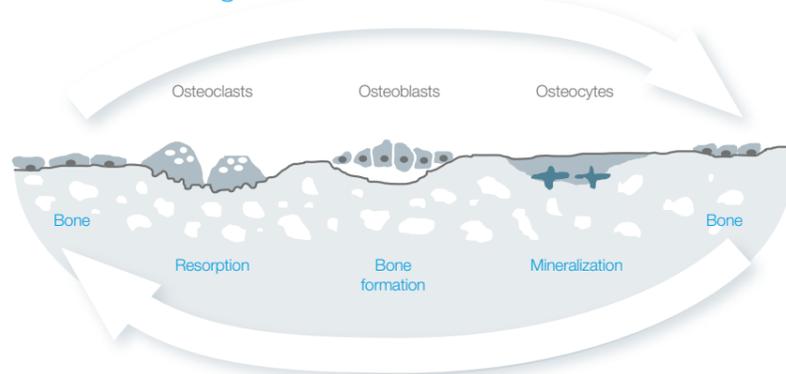
Furthermore, the continuous rebuilding serves to adapt the micro structure of the bone (direction and density of trabeculae) to changing loads. These adaptations are the reason for bone atrophy following missing load (e.g. atrophy of the jaw bone after tooth loss).



Wolff's law – bone density and structure adapt to changes in load

Three different types of bone cells contribute to bone *remodeling*. The degradation of old bone matrix is carried out by osteoclasts. In the course of this process so called resorption lacunae are built that afterwards are filled with new bone matrix by cells called osteoblasts. The osteoblasts are sealed by the mineralization of the extracellular matrix. These mature bone cells that are no longer able to produce osteoid are called osteocytes. Osteocytes are involved in the formation and restructuring of the bone and are therefore important for maintaining the bone matrix.

Bone remodeling



Balance between bone degradation by osteoclasts and bone formation by osteoblasts.

Bone and regeneration techniques



The use of bone graft materials

Bone graft materials are applied to replace and regenerate bone matrix lost by various reasons such as tooth extraction, cystectomy or bone atrophy following loss of teeth or inflammatory processes. For the filling of bone defects the patient's own (autologous) bone is considered the „gold standard“, because of its biological activity due to vital cells and growth factors¹. Nevertheless, the harvesting of autologous bone requires a second surgical site associated with an additional bony defect and potential donor site morbidity.

In addition, the quantity of autologous bone is limited. Today, due to a progressive development, bone graft materials provide a reliable and safe alternative to autologous bone grafts.

Clinicians can choose between a variety of different bone graft materials and augmentation techniques. Bone graft materials are classified by their origin into four groups.

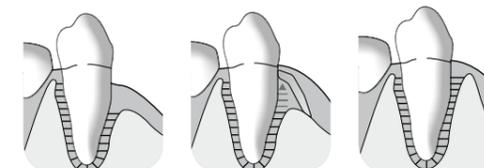
The GBR/GTR technique

The principle of Guided Bone Regeneration (GBR) or Guided Tissue Regeneration (GTR) is based on the separation of the grafted site from the surrounding soft tissue by application of a barrier membrane. Collagen membranes act as a resorbable matrix to avoid the ingrowth of the faster proliferating fibroblasts and/or epithelium into the

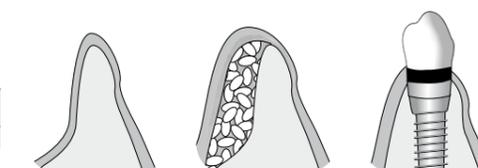
defect and to maintain the space for controlled regeneration of bone².

The application of bone graft material into the defect prevents the collapse of the collagen membrane, acting as a place holder for the regenerating bone and as an osteoconductive scaffold for the ingrowth of blood vessels and bone forming cells.

Guided Tissue Regeneration (GTR)



Guided Bone Regeneration (GBR)



¹ Illlich et al. (2011). Concise review: induced pluripotent stem cells and lineage reprogramming: prospects for bone regeneration. *Stem cells (Dayton, Ohio)* 29:555–563.

² Rothamel et al. (2012). Clinical aspects of novel types of collagen membranes and matrices - Current issues in soft- and hard-tissue augmentation. *European Journal for Dental Implantologists*.



maxresorb® 0.8 - 1.5 mm

maxresorb® 0.5 - 1.0 mm

Classification

Autologous:

- Patient's own bone, mostly harvested intraorally or from the iliac crest
- Intrinsic biologic activity

Allogenic:

- Bone from human donors (multi-organ donors or femoral heads of living donors)
- Natural bone composition and structure

Xenogenic:

- From other organisms, mainly bovine origin
- Long-term volume stability

Alloplastic:

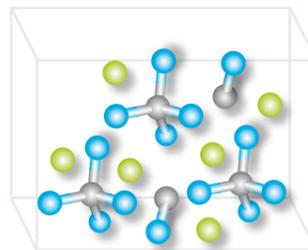
- Synthetically produced, preferably calcium phosphate ceramics
- No risk of disease transmission

For large defects a mixture of autologous or allogenic bone, which has excellent biological potential, and a bone graft material for volume stability of the grafting site, is recommended.

Development of bone regeneration materials – usage of calcium phosphates

The benefit of calcium phosphate ceramics as bone regeneration materials was realized long ago, as they are the main component of bones and therefore provide an excellent biocompatibility without any foreign body reactions.

In contrast with the first solely bioinert biomaterials, the advantages of calcium phosphates are their bioactive properties as well as their resorbability. Calcium phosphates support the attachment and proliferation of bone cells and undergo a natural remodeling process that includes osteoblasts and osteoclasts and that is characterized by an initial integration of the material into the surrounding bone matrix and a gradual degradation. Among the calcium phosphates, hydroxyapatite (HA), alpha-tricalcium phosphate (α -TCP) and beta-tricalcium phosphate (β -TCP) or biphasic CaPs are most commonly used as bioceramics. Compared to all other calcium phosphates, hydroxyapatite shows the slowest solubility, therefore providing the highest stability. By contrast, the alkaline β -TCP demonstrates a higher solubility and thereby fast resorption kinetics.



Crystalline structure of maxresorb®

Hydroxyapatite (HA)
 $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$

Beta-tricalcium phosphate (β -TCP)
 $\text{Ca}_3(\text{PO}_4)_2$

An ideal bone regeneration material should be resorbed in pace with new bone matrix formation. The basic principle of the biphasic calcium phosphates is a balance between the stable hydroxyapatite, which can be found years after the implantation, and the fast resorbing β -TCP. Bone regeneration materials based on mixtures of HA and β -TCP have successfully been applied in dental regenerative surgery for more than 20 years.



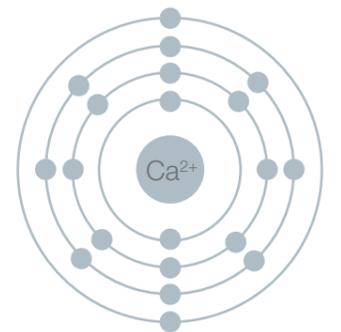
The ideal composition – biphasic calcium phosphates

The resorption properties of biphasic calcium phosphates can be changed by varying the mixing ratio of HA and β -TCP. A HA/ β -TCP ratio between 65:35 and 55:45 has been proven particularly suitable in many studies^{3,4} and offers a controlled resorption with parallel bone formation^{5,6}.

maxresorb® inject – injectable bone paste

Injectable calcium phosphates – cements and putties

Bone regeneration materials based on calcium phosphates are available as powders, granules and as porous blocks. The development of injectable bone regeneration materials started with the discovery of calcium cements in the 90's⁷. Cements result from the mixing of calcium phosphate powder with an aqueous solution. Following application, the hardening occurs *in vivo*. Cements create the possibility for several minimal invasive therapies of bony defects and offer an easier handling in many indications. The main disadvantage of the calcium phosphate cements is the reduced vascularization and natural remodeling experienced due to the stiffness and lack of interconnecting pores within the polymerised matrix. By mixing calcium phosphate granules with a water-based gel made of nano/micro hydroxyapatite granules (nano/micro HA) a moldable and non-hardening bone paste (putty) can be created. An example of such a non-hardening putty is maxresorb® inject.



Schematic drawing of a calcium atom

Calcium

Putties offer two significant advantages over cements: First, their increased porosity allow for the ingrowth of blood vessels and bone tissue, resulting in a fast and complete integration into new bone matrix and a rapid natural remodeling. Second, due to their large surface area, the nano/micro HA particles exhibit a high biologic activity resulting in an osteostimulative effect of these putties. Nano/micro HA particles support the adhesion of bone cells and thereby a fast formation of new bone as well as a fast particle degradation, offering additional space for the ingrowth of bone tissue.

- Alkaline earth metal
- One of the most common elements
- Essential mineral for humans
- Important for regulation of metabolism
- Besides phosphate, the main component of bone

³ Gauthier et al. (1999). Elaboration conditions influence physicochemical properties and *in vivo* bioactivity of macroporous biphasic calcium phosphate ceramics. *Journal of materials science. Materials in medicine* 10:199–204.

⁴ Schwartz et al. (1999). Biphasic synthetic bone substitute use in orthopaedic and trauma surgery: clinical, radiological and histological results. *Journal of materials science. Materials in medicine* 10:821–825.

⁵ Daculsi (1998). Biphasic calcium phosphate concept applied to artificial bone, implant coating and injectable bone substitute. *Biomaterials* 19:1473–1478.

⁶ Ducheyne et al. (1993). The effect of calcium phosphate ceramic composition and structure on *in vitro* behavior. I. Dissolution. *Journal of biomedical materials research* 27:25–34.

⁷ Brown and Chow (1985). Dental restorative cement pastes. US Patent 4'518'430, American Dental Association Health Foundation, USA.

maxresorb® – Innovative biphasic calcium phosphate



maxresorb® is an innovative, safe, reliable and synthetic bone graft substitute with improved controlled resorption properties and outstanding handling characteristics. The composition of 60% slowly resorbing hydroxyapatite (HA) and 40% beta-tricalcium phosphate (β -TCP) results in two mineral phases of activity:

maxresorb® supports the formation of new vital bone, maintains the volume and gives mechanical stability over a long time period.

The osteoconductivity of maxresorb® is achieved by a matrix of interconnecting pores and a very high porosity of approx. 80%, as well as pore sizes from ~200 to 800 μ m. The high macroporosity of maxresorb® is ideal for intense osteogenic cell growth and promotes the regeneration of vital bone. The high microporosity and surface roughness of maxresorb® facilitates diffusion of biological fluids and cell attachment.

The maxresorb® production process ensures a completely homogenous distribution of the two calcium phosphate phases; resulting in a high reproducibility. The unique maxresorb® production process leads to a highly nanostructured, bioactive rough surface for improved cell-adherence and hydrophilicity.

Properties of maxresorb®

- Synthetic
- Safe, reliable and sterile
- Biphasic homogenous composition
- Completely resorbable
- Very rough, hydrophilic surface
- Highly interconnected porosity



Incident light microscopy of maxresorb®

maxresorb® inject – Innovative synthetic injectable bone paste

maxresorb® inject is a unique and highly innovative, injectable bone graft paste, with controlled resorption properties.

The unique homogenous four-phasic composition of gel, active hydroxyapatite and granules of 60% HA/40% β -TCP supports the formation of new vital bone, maintains volume, and is gradually replaced by new mature bone.

The highly viscous maxresorb® inject paste allows perfect shaping, molding, fitting and complete bonding to the surrounding bone surface of the defect. maxresorb® inject is a non-hardening synthetic bone paste.



Easy handling and moldability of maxresorb® inject



maxresorb® inject syringe

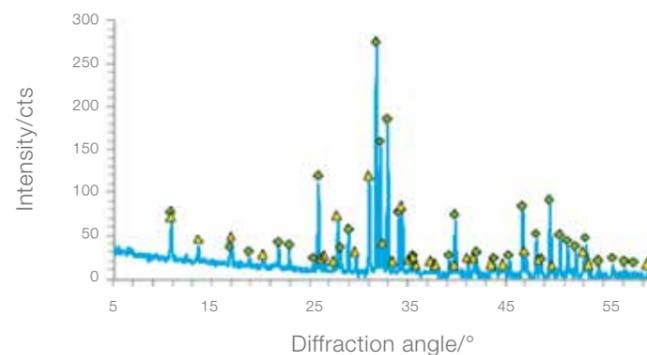
Properties of maxresorb® inject

- Injectable and easy handling
- Viscous and moldable
- Non-hardening
- Optimal adaptation to surface contours
- Active nano/micro HA particles

Production process



maxresorb® – absolute safety and phase purity



Safety by phase purity – x-ray spectroscopy of maxresorb®, Prof. Dr. C. Vogt, University of Hanover, all peaks can be assigned to HA (yellow) or β -TCP (green).

Indications:

Implantology,
Periodontology and
Oral and CMF Surgery

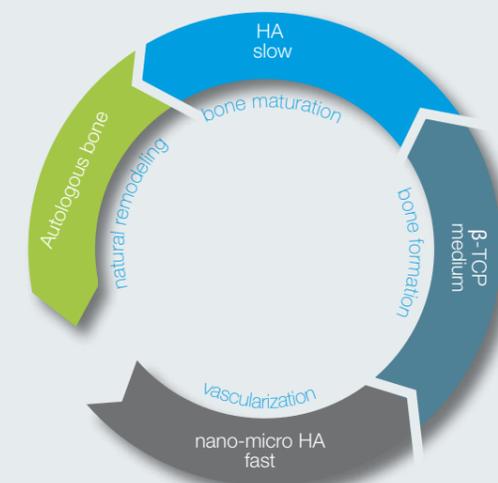
- Sinus lift
- Ridge augmentation
- Intraosseous defects
- Osseous defects
- Furcation defects
- Extraction sockets

Indications:

Implantology,
Periodontology and
Oral and CMF Surgery

- Sinus lift
- Intraosseous defects
- Extraction sockets
- Osseous defects
- Furcation defects

maxresorb® inject resorption profile Four-phasic activity



Product Specifications



maxresorb® granules

Art.-No.	Particle Size	Content
20005	0.5 - 1.0 mm (S)	1 x 0.5 ml
20010	0.5 - 1.0 mm (S)	1 x 1.0 ml
20105	0.8 - 1.5 mm (L)	1 x 0.5 ml
20120	0.8 - 1.5 mm (L)	1 x 2.0 ml

maxresorb®



maxresorb® blocks

Art.-No.	Dimension	Content
21211	20 x 10 x 10 mm	1 x block
21221	20 x 20 x 10 mm	1 x block

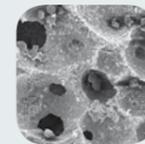
maxresorb® inject

Art.-No.	Unit	Volume
22005	1 x syringe	1 x 0.5 ml
22010	1 x syringe	1 x 1.0 ml
22025	1 x syringe	1 x 2.5 ml

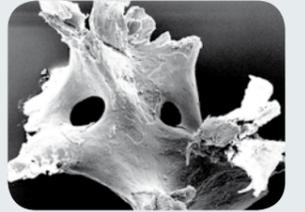


Biology as a model

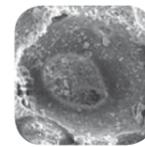
Interconnected porosity



The unique production process leads to porous ceramics, resembling the structure of human cancellous bone with fully interconnected pores.



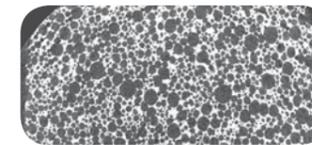
SEM image of human bone



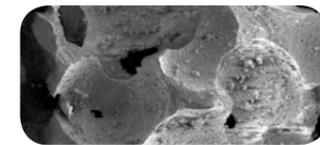
Interconnective porosity of maxresorb® inject

These interconnected pores are like tunnels within the material, providing access for fluids (blood) and also giving space and a surface for the ingrowth and migration of cells and blood vessels, thereby enabling the formation of new bone not only superficially but also inside the particles.

Relevance of the structure of bone regeneration materials



Micro CT image of maxresorb®

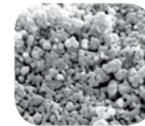


SEM image of maxresorb®

Macro - guidance
Rapid vascularization
Osteoconduction
Bone formation in pores

Rough surface – optimal condition for adhesion of cells and proteins

Beside safety, the advantage of synthetic materials lies in the reproducibility and ability to influence the structure. Due to a unique production process, maxresorb® has a very rough surface. This roughness is the basis for the osteostimulative effect often reported



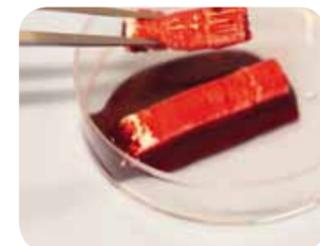
SEM image of maxresorb® showing very rough surface

for calcium phosphates. Proteins, such as growth factors, adhere to the surface and support the bony regeneration. Moreover, the nanostructured surface promotes the adhesion of cells and also their final differentiation. Likewise, the excellent hydrophilicity of maxresorb® is based on its surface roughness. Blood is very quickly absorbed, and proteins (e.g. growth factors) from the blood adhere to the inner and outer particle surface, promoting regeneration and integration.

Micro - communication
Ingrowth of cells
Blood uptake by capillary effects

Nano - nutrition
Adhesion of cells, proteins (growth factors) and nutrients

Excellent blood uptake of maxresorb® and maxresorb® inject



Blood uptake of maxresorb® (hydrophilic surface)



Hydrophobic material in contact with blood

In vitro research

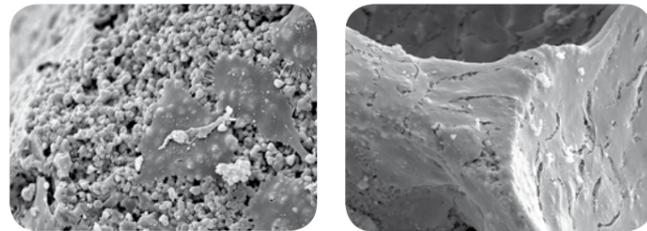
Proliferation of osteoblasts on maxresorb®
 Prof. Dr. Dr. D. Rothamel,
 University of Düsseldorf, Germany

The nanostructured surface of maxresorb® provides ideal conditions for the adhesion of osteoblasts. *In vitro* experiments demonstrated a fast proliferation of osteoblasts on maxresorb® granules.



Proliferation of SaOs-2 osteoblast-like cells on maxresorb®

After only seven days a dense colonization of cells was observed. The improved attachment and proliferation of osteoblasts promote the osseous regeneration, resulting in a fast integration of the particles into the newly formed bone matrix.



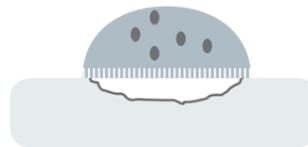
Osteoblasts on maxresorb® three and seven days after seeding

Osteoblasts:



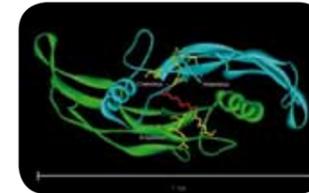
- Small, mononuclear cells, of embryonic mesenchymal cell origin
- Responsible for bone formation
- Settle on bone and release a collagenous basic substance (osteoid) into the intercellular space

Osteoclasts:



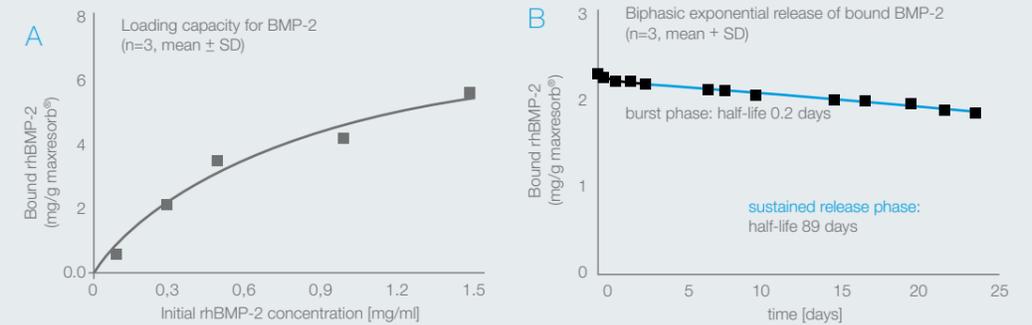
- Multi-nuclear giant cells formed by fusion of mononuclear progenitor cells of the bone marrow
- Main task is the resorption of bone substance by releasing protons (pH reduction) and proteolytic enzymes

Research with growth factors— Adsorption and release of BMP-2 from maxresorb®



In-vitro experiments show that maxresorb® can be loaded with up to 6 mg BMP-2/g (A). A two-stage, controlled exponential release of bound growth factors (B) indicates that maxresorb® is especially suitable to support the osseous integration®.

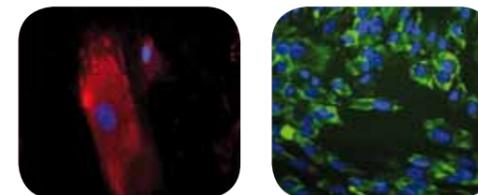
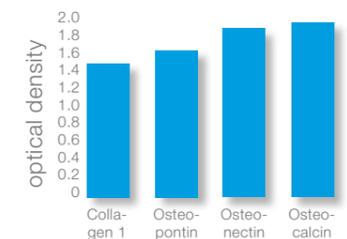
In-vitro experiments from Prof. Dr. H. Jennissen and Dr. M. Laub, University of Duisburg-Essen/MorphoPlant GmbH, Germany



Research with stem cells

maxresorb® supports the differentiation of stem cells
In-vitro experiments from Prof. Dr. B. Zavan and Prof. Dr. E. Bressan, University of Padova, Italy

Collagen, osteopontin, osteonectin and osteocalcin are proteins that are expressed from progenitor cells after they start to differentiate into osteoblasts. All of these marker proteins could be detected 14 days after seeding of stem cells on maxresorb® granules, indicating the correct differentiation of the stem cells®.



Immunofluorescence staining of stem cells grown on maxresorb®; red – osteopontin, green – osteocalcin

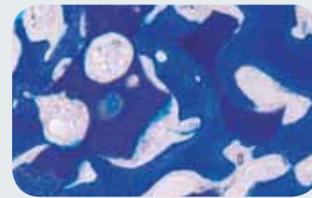
® Zurlinden et al. (2012). Immobilization and Controlled Release of Vascular (VEGF) and Bone Growth Factors (BMP-2) on Bone Replacement Materials. *Biomedical Engineering / Biomedizinische Technik* 57.

® Bressan et al. (2012). Donor age-related biological properties of human dental pulp stem cells change in nanostructured scaffolds. *PLoS One* 7:e49146.

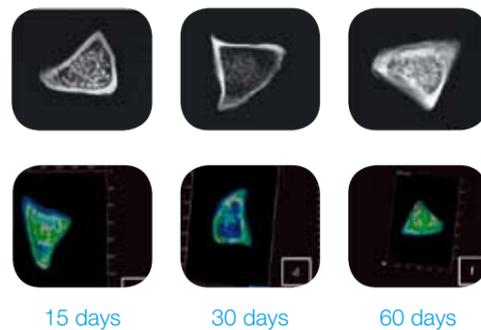
In-vivo pre-clinical testing

Enhanced bone formation and controlled resorption of maxresorb®

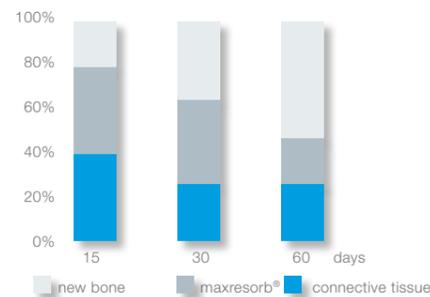
Histomorphometric and degradation study of maxresorb® in critical size defects in rabbits
Prof. Dr. J. L. Calvo-Guirado,
University of Murcia, Spain



Critical size defects were created in the tibia of rabbits and filled with maxresorb®. Nearly complete closure of the cortical defect after only 15 days. After 60 days, increase of medullary radiopacity, resembling cancellous bone¹⁰. The percentage of maxresorb® after 60 days is 27%.



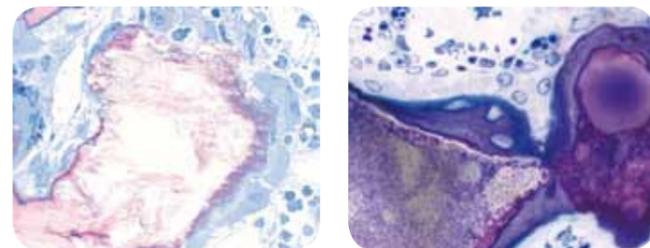
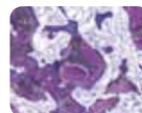
Radiographic image with corresponding thermal images showing the increase in radiopacity in the cortical and medullary zone



Histomorphometric results – percentages of new bone, maxresorb® and connective tissue

Fast integration and natural remodeling of maxresorb® inject
In vivo results of maxresorb® inject for filling of femoral defects in rats,
Prof. Dr. R. Schnettler, University of Gießen, Germany

Only three weeks after implantation, particles are covered by a layer of new bone matrix. A close contact between the newly formed bone and both components of the material (β -TCP and HA) can be seen.



Active osteoblasts (right picture) and osteoclasts (left picture) on the surface of the HA as well as the β -TCP component.

The presence of these cells is a sign for the natural remodeling of maxresorb® inject, with a degradation by osteoclasts and formation of new bone matrix by osteoblasts.

Predictable results for sinus floor elevation with maxresorb®

Results of a sinus lift study from Prof. Dr. Dr. D. Rothamel, University of Düsseldorf, Germany and Dr. D. Jelušić, Privat Clinic Opatja, Croatia¹¹

In a prospective randomized clinical study on 30 + 30 patients maxresorb® was compared to a pure beta-TCP for the indication of two-stage sinus floor elevation. Application of maxresorb® led to highly predictable bone regeneration with better volumen maintenance and radiological graft homogeneity compared to beta-TCP.

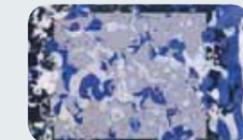
Histology of a trephine biopsy



Biopsy of trephine taken six months post-operatively



Detail of the histology



Computer-assisted histomorphometric analysis

Clinical case sinus lift, Dr. D. Jelušić

Following a healing phase of six months, biopsies from trephines taken at implant bed preparation demonstrated the osteoconductive properties of maxresorb®, supporting the formation of new bone matrix. Three-dimensional-radiological control images showed an excellent volume stability of the grafts, facilitating the insertion of the planned implants. No implant failures were observed in a first follow-up one year post-operative, emphasizing the safety and reliability of the biphasic material.



Elevation of mucoperiosteal flap



Preparation of lateral sinus window



Elevated Schneiderian membrane



Application of maxresorb®



Covering with Jason® membrane



Saliva-proof wound closure



Re-entry six months post-operatively

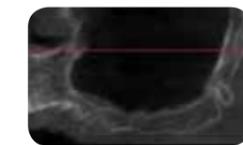


Implant uncovering

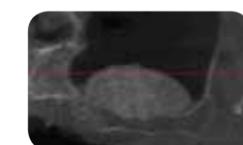


Inflammation-free soft tissue situation

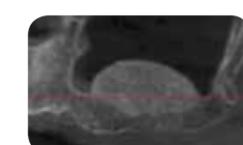
DVT control



Preoperative DVT: extended vertical bone defect



Situation post-operatively: large volume sinus lift without membrane perforation



Situation six months post-operatively: excellent volume stability and radiological homogeneity

¹⁰ Calvo-Guirad et al (2012). Histomorphometric and mineral degradation study of Osscerams: a novel biphasic B-tricalcium phosphate, in critical size defects in rabbits. Clinical Oral Implants Research, 2012, 23, 667-675.

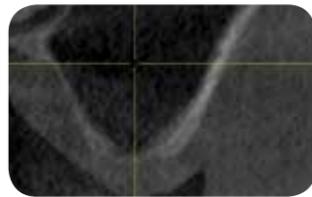
¹¹ Jelušić et al. Monphasic β -TCP vs. biphasic HA/ β -TCP in two-stage sinus floor augmentation procedures - a prospective randomized clinical trial. Clin Oral Impl Res. 00, 2016, 1-9 [Epub ahead of print]

Clinical case by Dr. Steffen Kistler,
Landsberg am Lech, Germany

Sinus lift with two-stage implantation



DVT control after sinusitis surgery, residual bone height one millimeter



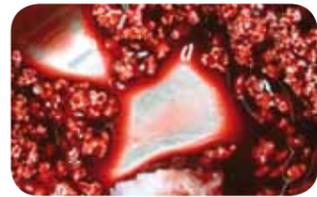
Transversal section to determine the depth of the sinus floor



Access to the sinus cavity by a lateral approach, minor perforation of the Schneiderian membrane



Covering of the perforation with Jason® fleece



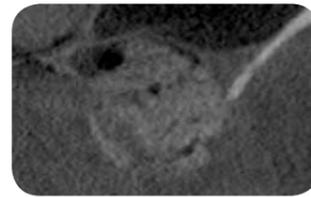
maxresorb® mixed with venous blood and collected bone chips



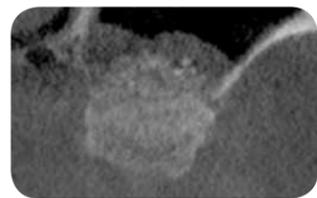
Augmentation of the sinus wall with a mixture of autologous bone and maxresorb®



Covering of the sinus window with collprotect® membrane fixed with two pins



Post-operative DVT control showing cavity between mucosa of the maxillary sinus and the membrane



Consolidation of graft material with minimal hyperplasia of sinus mucosa before implantation



Primary stable insertion of two implants after only eight weeks



OPG control of implant insertion



Uncovering of implants ten weeks post-operative



X-ray control after uncovering, showing dense regeneration of the graft material

Tip:

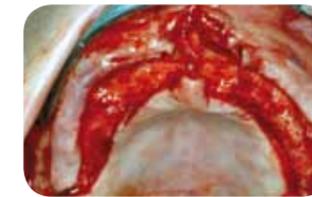
For easy application and optimal revascularization, the graft material should be mixed with blood collected from the defect, or with venous blood when larger volumes are needed.

Clinical case by PD Dr. Jörg Neugebauer,
Landsberg am Lech, Germany

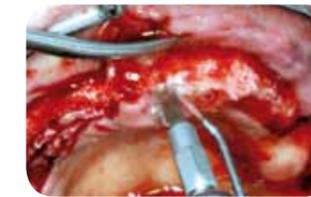
Circular bone splitting in the upper jaw



Three-dimensional implant planning with a radio-opaque scan template



Surgical presentation of the alveolar ridge with reduced amount of horizontal bone available



Deep bone splitting with oscillating saw in regio 15 to 25



Positioning of collprotect® membrane for application of bone graft material



Lateral deposition of maxresorb® to prevent resorption of the vestibular wall



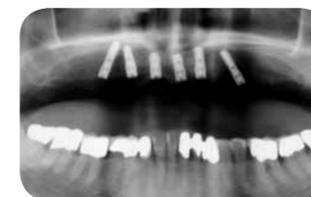
Covering of the augmentation site with the initially inserted membrane



Tight wound closure with a continuous seam following the periosteal splitting



Complication-free healing of the augmented ridge



OPG control of inserted implants along the anterior sinus floor



Re-entry surgery in combination with vestibuloplasty to form the vestibulum



Soft tissue situation after healing with inserted abutments



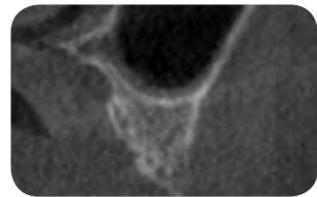
Inserted bridge with terminally screwed and anteriorly cemented implants

Tip:

To achieve an even contour when stabilizing bone splitting in lateral augmentations, the smaller granules (0.5 - 1.0 mm) should be used.

Clinical case by Dr. Frank Kistler,
Landsberg am Lech, Germany

Sinus floor elevation with simultaneous bone splitting
and implantation



DVT image demonstrating horizontal and vertical amount of bone available



Reduced amount of bone on both sides of the upper jaw



Surgical presentation of the ridge with mobilization of the sinus mucosa through a lateral window



Splitting of the ridge after crestal osteotomy with bone condenser



Augmentation of the sinus cavity and fixation of the lateral wall with maxresorb®



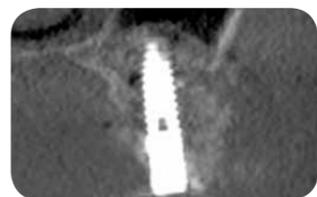
Lateral augmentation with maxresorb® and osteotomy site with Jason® fleece



Covering of augmentation site with collprotect® membrane



Single sutures for tight wound closure after periost splitting



DVT image to control the inserted graft material



Control three months after augmentation of the alveolar ridge



Good consolidation of the bone graft material with wide alveolar ridge



Reduction of mucosal situation at re-entry surgery



Crestally stable bone level at re-entry

Tip:

For stabilization of bone splitting, a combined application of graft material and membrane shows the best long-term results.

Clinical case by Dr. Georg Bayer,
Landsberg am Lech, Germany

Lateral augmentation



DVT image showing the reduced amount of bone available in the area of the mental foramen



Lateral bone defect following lateral bone defect following root tip resection



After preparation of the implant bed the thin vestibular wall is visible



Insertion of implant in the reduced bone amount



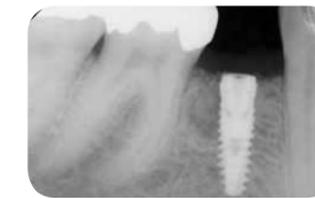
Lateral augmentation with maxresorb® and application of a dry collprotect® membrane



Complete covering of augmentation site and implant with the membrane



Wound closure by soft tissue expansion without vertical releasing incisions



Post-operative x-ray



Stable keratinized gingiva after insertion of healing abutment at re-entry



X-ray control at re-entry

Tip:

For lateral augmentation with minimally invasive surgery, initial placement of a membrane and subsequent application of a graft material is advantageous.

Clinical case by Prof. Dr. Dr. Daniel Rothamel,
University of Düsseldorf, Germany
Sinus lift with two-stage implantation



Clinical situation before sinus lift



Clinical situation before sinus lift, occlusal view



Surgical presentation of the buccal wall



Preparation of a lateral sinus window



Introduction of Jason® membrane into the sinus cavity



Jason® membrane in the sinus cavity to protect the Schneiderian membrane



Filling of the sinus cavity with maxresorb®



maxresorb® in the sinus cavity



Additional lateral augmentation with maxresorb®



Covering of the augmentation area with Jason® membrane



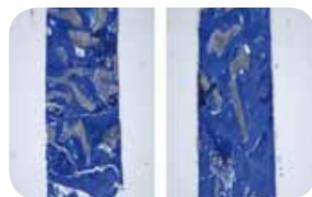
Tension-free wound closure with single button sutures



Good osseous integration of the maxresorb® particles without soft tissue ingrowth six months post-operatively at re-entry



Stable insertion of two implants into sufficient bone matrix



Histology of biopsy taken at implantation



Detailed image of histology showing complete integration of maxresorb® particles within new bone matrix



Post-operative radiograph

Clinical case by Dr. Frank Kistler,
Landsberg am Lech, Germany
Internal sinus lift



Endodontically treated tooth 26 with apical cyst formation



X-ray control before implantation with partially regenerated extraction socket



Presentation of the soft tissue situation before implantation



Preparation of the implant bed for internal sinus lift with bone condenser



The maxresorb® inject paste is brought to instrument for application



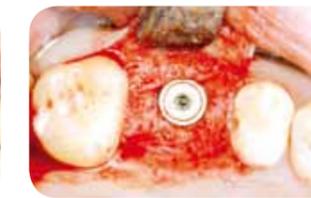
Insertion of maxresorb® inject for internal sinus lift



Augmentation of the sinus floor by a crestal approach



Insertion of maxresorb® inject with bone condenser



Inserted implant before wound closure

Tip:

For sinus floor elevation, the large maxresorb® granules (particle size 0.8 - 1.5 mm) are especially suitable to gain sufficient space for osteogenesis and revascularization, even when larger volumes of the bone graft material are applied.



X-ray control clearly showing the inserted maxresorb® inject

Tip:

For internal sinus lift, the moldable graft material maxresorb® inject is ideally applied by a lateral approach as no further mixing with blood is needed.

Clinical application of maxresorb® inject

Clinical case from Dr. Damir Jelušić,
Opatija, Croatia

Immediate implant installation



Extraction of tooth 14 and 15



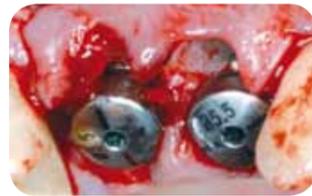
Buccal dehiscence of the bone wall of tooth 14



Osteotome technique with insertion of maxresorb® inject (transalveolar) at tooth 15



Immediate implant insertion in extraction sockets of tooth 14 and 15



Placement of the healing abutments



Placement of Jason® membrane at the buccal bone wall



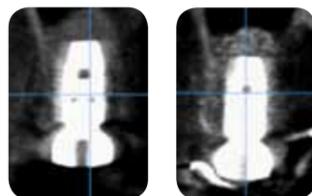
maxresorb® inject placed at buccal wall and protected by Jason® membrane



Wound closure and suturing



Tissue situation after five months of healing



3D CBCT four months post-operative



Situation after removal of healing abutments



Clinical view at control one year after surgery

Clinical application of maxresorb® inject

Clinical case from Dr. Damir Jelušić,
Opatija, Croatia

Ridge preservation and augmentation



Pre-operative x-ray



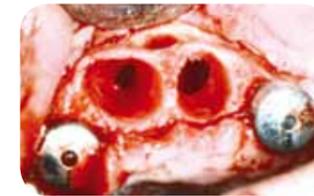
Clinical situation before surgery



Implant insertion in regio 12, 11 and 23



Insertion of healing caps



Extraction of tooth 21 and 22, defects of the buccal wall visible



Filling of the extraction sockets with maxresorb® inject



Covering of socket and buccal wall with Jason® membrane



Wound closure and suturing



Situation after six months healing time



Situation after removal of healing caps



Final prosthetic restoration



X-ray control eight months after implantation and extraction

bone & tissue
regeneration

botiss
biomaterials

Innovation.
Regeneration.
Aesthetics.

soft tissue

education

hard tissue

botiss biomaterials GmbH
Hauptstr. 28
15806 Zossen / Germany

Tel.: +49 33769 / 88 41 985
Fax: +49 33769 / 88 41 986

contact@botiss.com
www.botiss.com
www.facebook.com/botissdental

