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Implantology faces new challenges

Dear colleagues,

In recent years, implantology has been considered to be the driving force for innovations within dentistry. Companies and investors all over the world were impelled by the apparently infinite growth potential of this market. In the meantime, however, or so it seems, the mood has become much more reserved. Some implant systems have already been withdrawn from the market, and many companies do not act as aggressively as they used to. We have to find the answers to the following basic questions. Which are the real growth potentials of implantology, and how can implantology be placed on a changed market in the future? Contrary to the trend of the increase in gaining independence within our expert field, observable in the past, I believe that it will become more important to go back to the roots of implantology and consider it to be the real interface between periodontology, surgery, and prosthetics. The broad digitalization within dentistry by means of 3-D diagnosis combined with modern planning and navigation software, including CAD/CAM manufactured dentures, implies this interface function, also from the technological point of view. Due to the digitalization of dental practices, team spirit is fostered, the flow of information is improved, and cooperation between the partners is facilitated. Prosthetic dental planning and the prosthetic guidelines for surgery will be easier to communicate, ultimately contributing to the achievement of an optimal result, and thus efficiently meeting the desire of the patient. In this way, the old requirements and ideas for interdisciplinary cooperation can be realized in a much better manner. With this in mind, I would like to draw your attention to our 39th International Annual Congress which will take place from October 9th to 10th in Munich this year. Thanks to the high class speakers from Germany and abroad, our Annual Congress will again become a special highlight in advanced professional training. I look forward to welcoming you in Munich.

Yours,

Dr Friedhelm Heinemann
President of DGZI
Implantology faces new challenges
_Friedhelm Heinemann, Germany

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Bone quality, quantity and metabolism in terms of dental implantation

Abstract

Bone is the largest calcium storage, forms part of the supporting tissue and displays distinctive plasticity and adaptability. Thus, an adequate, differentiated composition and metabolism are required. The bone matrix consists of organic and inorganic structures. The cells, osteoblasts, osteoclasts, and osteocytes are responsible for bone formation, resorption and metabolism and, thus, for remodeling processes (formation and resorption) which permanently occur in bone tissue. Periosteum and endosteum form a functional unit with bone tissue itself and exercise protective, nutritive and growth functions.

The present paper provides an introduction to regular bone structure in the face area which is considered a precondition of successful implantation. The specific properties of the jaw bones have to be observed in this context.

Introduction

The orofacial or stomatognathic system with its components forms a functional circle representing a biocybernetic system. A key role is played by bone, particularly the jaw bones. The structure of the osseous viscerocranium is targeted to withstand and divert the chewing pressure. This requires a functional composition which is maintained even after osseointegration of bone graft substitutes (Fig. 1).

The periods of function and inactivity occurring in the jaw bones are more clearly noticeable than in any other bone of the body. For instance, the mandible and the maxilla display different atrophy processes. Continuous remodeling which guarantees adaptation to arising forces requires healthy bone.

Bone composition

Bone is a specific connective tissue which mainly consists of the extracellular substance/bone matrix, cells, blood vessels, and nerves.

Weight distribution (in kilograms) in a 70 kg weighing man shows that bone and bone marrow rank high, as shown by selected examples in table 1.

Table 1. Weight distribution (in kg) of tissues in a 70 kg weighing man in ranked order.

<table>
<thead>
<tr>
<th>Tissue</th>
<th>Weight (kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Muscles</td>
<td>30</td>
</tr>
<tr>
<td>fat</td>
<td>10</td>
</tr>
<tr>
<td>bone</td>
<td>7</td>
</tr>
<tr>
<td>bone marrow</td>
<td>3</td>
</tr>
<tr>
<td>blood</td>
<td>3</td>
</tr>
<tr>
<td>connective tissue</td>
<td>3</td>
</tr>
<tr>
<td>skin</td>
<td>2</td>
</tr>
</tbody>
</table>

Bone matrix

It consists of organic and inorganic material. The inorganic material which determines bone
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stiffness, constitutes about half of the matrix. The major part (60 per cent) is composed by calcium and phosphate in the form of hydroxyapatite crystals. Additionally, non-crystalline calcium phosphate, citrate, hydrogen carbonate, and furthermore, magnesium, potassium, and sodium salts are found. The apatite crystals are surrounded by matrix substance and lie along the collagen fibrilles. A so-called hydration cover facilitates ion exchange between crystal and body fluids. The water content of the bone matrix amounts to 10–20 per cent. The water-free matrix contains 70 per cent inorganic and 30 per cent organic substance.

The non-calcified organic matrix is referred to as osteoid and provides the bone with the required elasticity. It mainly consists of collagen.

Bone cells
Cells are indispensable for regular osseointegration processes. Osteoblasts: These mononuclear and nearly cubic shaped cells produce all essential organic matrix components of bone (osteoid). They are located exclusively at the surface of spongy trabecles, and synthesize and secrete collagen I, proteoglycans and glycoproteins. As they activity fades, the cells become increasingly flatter and form processes. The newly produced matrix proteins are released towards the surface of the existing bone matrix. The organic material enabling bone elasticity constitutes about 20 per cent. Finally, water amounts to about 10 per cent of the matrix substance.

The osteoblasts are activated by cytokines, growth factors and vitamin D3 and thereby enabled to induce osteoclast proliferation and differentiation.

Osteocytes develop from osteoblasts and represent metabolic centres of the bone. These cells predominate in bone. Their cell bodies are located in the lacunae. With their processes, they are interconnected by nexuses. The processes permit intercellular substance transport and enable the coordination of metabolic activities. An exchange of compounds among osteocytes, mineralized matrix and blood vessel also occurs in the gap system between cells and calcified bone matrix.

The osteocytes respond to mechanical stress exerted on the bone and sustain the extracellular matrix. After their death, the matrix undergoes resorption.

The jaw area is very likely to exhibit a population of bone cells different from cells from other regions in terms of their origin, their involvement in specific local developmental and bone formation processes and their specific adaptation to local biomechanics. At the moment, it may be merely speculated whether the activity of such neuroectodermally derived “jaw osteoblasts” essentially contribute to perfect dental implant treatment outcomes.

Periosteum and endosteum
Their main functions are protection and nutrition as well as the continuing supply of osteoblasts for

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**Fig. 2.** Osseous lamella.
**Fig. 3.** Functional composition of bone and dynamic equilibrium.
thickness growth and successful defect repair. The outer periosteum consists of fibroblasts, collagenous and elastic fibres (stratum fibrosum). The inner layer (stratum germinativum) is formed by divisible cells ("lining cells") which can differentiate into osteoblasts. The stem cells play an important role for bone growth and repair. Sharpey’s fibres are bundles of collagenous fibres connecting the periosteum with the bone substance.

The endosteum fills the inner cavities of bone, is thinner than the periosteum, consists of progenitor cells and comprises only a slight amount of connective tissue. The endosteal border cells are resting osteoblasts forming a continuous cell assembly on the inner cortical surface and the Haversian canal walls.

Bone types
Woven bone and lamellar bone are distinguished according to the arrangement of osteocytes and collagen fibres.

- **Woven bone** (also "primary bone") is found only during bone development and repair processes, and, therefore, also in osseointegration. Its mineral content (higher radiolucency!) is lower. The collagen fibrils run irregularly rather than in lamellar form. This type of bone is replaced by lamellar bone except for suture areas.

- **Lamellar bone** (Fig. 2): The collagen fibres and other matrix components form lamellae of 3–7 µm thickness which are arranged in concentric layers around a central channel (Haversian canal). This structure is referred to as Haversian system or osteon. Between the lamellae, the cell bodies of the osteocytes are located with processes running in respective channels (see above). Each channel harbors nutritive vessels, nerve fibres, and loose connective tissue. The canals communicate with the bone marrow cavity, the periosteum and with each other. Larger channels are Volkmann’s canals which also run outward. The entire channel system reflects a complex and delicate microcirculation.

Each osteon is surrounded by mineralized matrix with few collagen fibres (cementum). Immediately beneath the periosteum and around the marrow cavity lie more outer and fewer inner general lamellae. Between outer and inner general lamellae, the so-called osteons and the often irregularly shaped intermediate lamellae are located. The latter are residual lamellae of a Haversian system which was degraded during a remodeling process.

Principles of bone structure
- **Substantia compacta, Substantia spongiosa of the bones**

The outer compact bone layer is referred to as substantia compacta, while the inner layer which possesses numerous interconnected cavities is termed substantia spongiosa. The latter represents a spongy...
Bone shape and composition are adjusted to its mechanical function. According to its trajectorial architecture, structure conforms to tension and strain trajectories. Bone substance is arranged in such a way that best possible absorption and transmission is achieved with minimum expense of material in the loaded area. The trajectorial construction, e.g., of the mandible, permits selective material usage in the loaded area. This lightweight construction saves muscular strength for motor activity. Bone morphology is genetically determined and modified by external influences through differentiation. Bone is solid by virtue of its inorganic components and elastic due to its organic components. Thus, bone substance features a dynamic equilibrium of adaptation.2, 3, 8

Special properties of the jaw bones

The jaw bones as well as the tooth germs (except the enamel organ and enamel) develop from the neural crest cells which are preprogrammed for the facial region where to migrate.1 Development of cartilage, bone and teeth, however, requires numerous interactions between the mesectodermal cells, the ectoderm and the inner body surface. Almost all jaw bones (including the alveolar process) develop to woven bone by desmal osteogenesis. The osteoblast progenitor cells are derived from neural crest cells. Ossification of the alveolar processes is closely associated with tooth germ development. Desmal ossification is recapitulated in reparative processes. Similarly, osseointegration of dental implants follows this principle.2

Jaw atrophy

Jaw atrophy is a major field of implantology. Such inactivity atrophy results from a lack of strain stimuli and reflects complex atrophic and resorptive processes. Resorption of the (edentulous) alveolar processes is multifactorial and cannot be attributed to altered loading alone. Mechanical factors, biological and metabolic factors, and inflammatory causes are distinguished.

Mandibular involution is primarily vertical. The alveolar process is smaller than the basal arch which represents the constant part (Fig. 4). At old age, the mandible shows increased prominence.1 Vertical bone resorption is increased compared to the maxilla. These changes are even more marked in females. The mandibular basal arch is more extended than the maxillary one: by up to ca. 3 mm during the first three years after tooth loss, and by 0.3 to 0.4 mm p.a. during the following years. This difference is compensated by alveolar process inclination resulting in oral inclination of the mandibular alveolar process and vestibular inclination of the maxillary one.

Besides the basal arch which may also exhibit resorption during old age, there are structures that are not subject to atrophic changes, such as linea mylohyoidea, linea obliqua, spina mentalis, tori mandibulares, and trigona retromolaria.1

The basal arch as well as the alveolar walls show a trajectorially oriented arrangement.

Alveolar process atrophy also leads to an altered mandibular angle. This angle amounts to 150° in newborns and to 120° in adults. At old age, the angle tends to return to childhood values and may even exceed 160°. These angular changes also exert an effect on the trigonum retromolare sinking down in atrophic bone and, thus, assuming an almost topographical relation to the pars obliqua of the mandibular canal. In atrophic bone, the trigonum retromolare fails to serve as an implant abutment site.10

Maxillary atrophy primarily occurs horizontally and only slightly in vertical direction. The alveolar process which provides only little space for the tooth roots, is larger in relation to the corpus maxillae.7 Thus, the maxillary arch becomes more reduced. Due to resorption, its size decreases relative to the mandible. With increasing maxillary atrophy, the alveolar midcrest is displaced palatally (centripetally). Atrophy of the hard palate proceeds anteroposteriorly (Fig. 5) and may be of such an extent as to cause bone perforation and contact between oral and nasal cavity mucosa.

The alveolar process ends with the tuber retromolare behind the last molar. Before seven years of age, the tuber does not exist but as a rudiment. The alveolar canals for the nn.alveolares superiores posteriores develop from primordial alveolar sulci. Beyond 20 years of age, the tuber is fully differentiated, and alveolar canals are found. After 50 years of age, involution of the tuber occurs, and the alveolar canals reopen into alveolar sulci. These changes show that the tuber retromolare is not suitable as an implant abutment.
The anterior maxilla shows numerous channels harboring blood vessels and nerves which also present as sulci and open toward the maxillary sinus. The torus palatinus, the crista zygomatico, and the spina nasalis anterior do not reveal any atrophic alterations. The chewing pressure abutments of the maxilla are not subject to atrophic processes and, hence, preserved.

### Blood vessel supply
Blood vessel supply is a prerequisite of regular bone growth and metabolism. It is effected by means of the vasa nutricia which run without contortions through the corticalis into the medullary cavity (Fig. 2). Diaphyseal arteries branch out to marrow arteries. Several small arteries supply the metaphysis and epiphysis. The epiphyseal vessels display only few anastomoses with the metaphyseal and diaphyseal vessels. The corticalis is supplied from within and outside. The outer cortical sections are supplied directly by the periosteal vessels. Perfusion is increased in high-density bones compared to low-density bones. This should be considered for implantation.

Bone tissue lacks lymphatic vessels which occur only in the periosteum.

### Nerve supply
Nerves are indispensable for bone tissue viability as well. With the blood vessels, the nerves enter the bone, run in Volkmann's and Haversian canals (Fig. 2) and also reach the spongiosa. The nerve fibres are largely unmyelinated. The periosteum is innervated abundantly. There has been evidence of numerous transmitters (e.g., norepinephrine, neuropeptide Y).

### Bone function
Bone function is manifold and plays an important role for osseointegration.

### Supportive function
The bones make up the skeleton which determines skull weight and shape. Additionally, the skeleton provides the origin and attachment of the muscles which in turn influence cranial morphology. Owing to its biological plasticity, bone is capable of adaptation to stress.

### Protective function
The cranial bones protect the central nervous system, the sense organs, and the bone marrow. Development of masticatory pressure pillars in the viscerocranium serves for diversion of the chewing pressure and the trajectorial structure of the mandible (Fig. 1).

### Calcium storage
About 99 per cent of the body’s calcium is stored in the skeleton. The stored calcium is mobilized, when blood calcium concentration declines. Calcium is mainly released from hydroxyapatite crystals in the spongy substance. Acting as a calcium storage, bone tissue plays an important role in regulating calcium homeostasis.

### Hormone and vitamin metabolism
Bone is related to hormone and vitamin metabolism. The osteoblasts possess receptors for parathyroid hormone, vitamin D3, cytokines and growth factors. They produce factors that increase osteoclast proliferation and differentiation. Calcitonin exerts a receptor-mediated inhibiting effect on osteoclast activity. Both androgens and estrogens generally stimulate bone anabolism and accelerate epiphyseal gap and suture closure. They facilitate mineralization and bone formation. Opposite effects are exerted by cortisol.


The Literature list can be requested from the editorial office.

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Reconstruction of damaged fresh sockets by connective-bone sliver graft from the maxillary tuberosity, to enable immediate dentoalveolar restoration (IDR)—A clinical case

Abstract

This paper describes a procedure for immediate loading of an implant following tooth extraction, in a socket presenting severe damage to the vestibular bone plate and gingival recession in the region of the upper central incisors. The procedures of extraction of the tooth, immediate insertion of the implant, connective-bone graft from the maxillary tuberosity and immediate restoration were shown to be a predictable treatment alternative. These procedures led to restoration of the tooth, bone and gingival structures in a single surgical stage and to maintenance of the favorable esthetic and functional result 24 months afterwards.

Introduction

The situations in which immediate loading of implants following tooth extraction is indicated are expanding. Maintaining the integrity of the support tissues during a tooth extraction procedure is an indispensable requirement for indicating immediate loading. The finding of the phenomenon of integration between bone and titanium changed the course of dentistry. Thus, to promote osseointegration for a titanium fixture, it would be necessary to burry the implant in order to avoid premature loading, thereby requiring two surgical stages before inserting the prosthesis. The aim of this paper is to describe a technique for immediate loading of an implant following tooth extraction, in a socket presenting severe damage to the vestibular bone plate and gingival recession in the region of the upper central incisors.
It has been shown that certain forces are important for triggering a series of biological reactions that accelerate the bone repair process, thus encouraging the use of implants for immediate loading (Romano G 2001, Isidor F 2006).


Primary stability is the most important factor in indicating that an implant can be brought into immediate functioning. This is associated with the quantity and quality of the bone material, the geometry of the implant and the surgical technique used (Polizzi G 2000, Covani U 2003, Hamerle CH 2004, Covani U 2007, Canullo L 2007, Schropp L 2008).

Implants of conical design are the type most indicated for receiving immediate loading following tooth extraction. They promote greater lateral bone compaction, thereby increasing the bone density; they adapt better to the socket; and they have a greater contact surface with the bone. These characteristics increase the initial stability and enable better spreading of occlusal loads (Gomez-Roman G 2001, O'Sullivan D 2000, Abbou M 2003, Friberg B 2003, O'Sullivan D 2004, Glauser et al 2007).

The presence of local infection due to disease or trauma in the damaged tooth interferes directly with the quantity and quality of the soft and hard tissues. It may change the alveolar architecture at the potential site for implant insertion. Additional loss of these esthetically important tissues may occur while treating the local infection, thereby increasing the involvement of esthetics. One or more socket walls may be impaired, with or without gingival recession, and surgical techniques may be needed to restore their anatomy, thus contraindicating implants with immediate loading (Polizzi G 2000, Chen ST 2004, Lindeboom JA 2006, Schropp L 2008).

As a donor area, the maxillary tuberosity has a limited quantity of bone material available and presents low bone density and the difficulty of surgical access. Its advantages are the excellent postoperative recovery, better graft repair and the ease of harvesting the graft material and adapting it to the receptor region, because of the bone malleability (Rosa JCM 2008).

Anatomical and functional recovery of the hard and soft tissues surrounding the pos-extration immediate loading implant was achieved by means of the technique of connective-bone sliver graft, as a single specimen harvested from the maxillary tuberosity.

**Clinical case report**

The patient was a 50-year-old woman with a high smile line who came for treatment because of gingival recession in the region of the upper right central incisor.

At the clinical examination, periodontal probing of the compromised tooth showed severe damage to the vestibular bone plate, with gingival recession of 4 mm combined with vestibular bone loss to a depth of 10 mm, without apparent fistula.

Radiographic examination showed the presence of endodontic treatment, a short cast metal core, a metal/ceramic crown, bone height of 8 mm above the root apex and thickening of the hard layer in the region of the compromised tooth. Good bone material availability in the maxillary tuberosity was observed.

The treatment proposed consisted of extraction of the compromised tooth, curettage of the region, implant insertion, recovery of the socket defect by connective-bone sliver graft from the maxillary tuberosity and immediate temporary restoration. It was suggested to the patient that she might wish to replace the metal/ceramic crowns with metal-free crowns on the anterior teeth of the maxilla.

The following medications were prescribed:

- **Amoxicillin 500 mg**, taken as one capsule every eight hours, for seven days starting one hour before the procedure.
- **Dexamethasone 4 mg**, taken as one dose of 8 mg one hour before the procedure and then 4 mg per day for two more days.
- **Paracetamol 750 mg**, taken as one pill every six hours while in pain, beginning one hour before the procedure.

For better comprehension, the steps in the procedure will be described in detail in the following.
Sequence of procedures:

 Tooth extraction and implant insertion
1. Infiltrative anesthesia of 2% mepivacaine, with norepinephrine, at the base of the vestibule, in the palate and near the papillae adjacent to the compromised tooth.

2. Double incision at the level of the mesial papillae and distally to the compromised tooth, using a microblade (69 WS Swann-Morton®, England). The aim of this incision was to reposition the gingival tissue coronally, thereby minimizing the trauma to the tissues and promoting scar-free first-intention healing.

3. Incision in the sulcus, using the microblade, around the tooth that was to be extracted.

4. Non-traumatic extraction of the tooth using a periosteum, performing a pendular movement in the mesiodistal direction, with the aim of maintaining the integrity of the remaining bone walls.

5. Careful curettage of the socket, to remove the granulation tissue and the remains of the periodontal conjunctive tissue.

6. Probing of the socket walls to assess the degree of bone damage, in the apical-coronal and mesiodistal directions, and to confirm the anatomical shape of the defect.

7. Insertion of a NobelReplace™ Tapered TiUnite® (Nobel Biocare™, Göteborg, Sweden) implant, of dimensions 16.0 x 4.3 mm, with excellent apical stability and diameter compatible with the socket opening. By means of a surgical guide, the drill bits were directed towards preparing the bone bed, using the palatine wall to ensure adequate bone support and insert the implant in the ideal three-dimensional position. The implant bed preparation was started using the 2.0 cylindrical drill bit and continued using the 3.5 and 4.3 conical-shaped drill bits. The implant was inserted such that the ideal apical-coronal positioning was sought, independent of the abnormality of the gingival level. The implant platform was at a distance of 3 mm from the gingival margin of the homologous tooth. After insertion of the implant, the spirals remained exposed in the socket defect region.

8. Insertion of a temporary titanium abutment, adjustment of the occlusion and opacification of the metallic component by using photopolymerizable opaque resin (Amelogen® Plus OW, Ultradent Products, Inc., USA);

9. Construction of a provisional crown using aesthetic facet that was prepared earlier using photopolymerizable resin. The ideal emergence profile was established on the provisional crown, with free space to allow for accommodation of the soft tissues and to promote a thicker and more stable margin of gingival tissue;

10. Insertion of the provisional crown for adjustment of the occlusion in centric, habitual and excursive movements and for clinical confirmation of its adaptation, considering that the implant platform was above the gingival margin.

11. Removal of the provisional crown for finishing and polishing, and to perform the stage of reconstruction of the sockets bone defects.

 Graft harvesting
12. Infiltrative anesthesia in the donor area by means of 2% mepivacaine, with norepinephrine, at the base of the vestibule and in the palatine portion of the maxillary tuberosity.

13. Incision following the distal outline of the second molar, to the full depth of the soft tissue, followed by two relaxing incisions in the posterior direction, thus reproducing the shape of the defect in the receptor region.

14. Longitudinal incision dividing the gingival tissue as far as the most posterior portion of the relaxing incisions, while maintaining a thickness of around 1 to 2 mm of connective tissue covering the bone tissue.

15. With the use of a straight chisel, the bone was cut along the relaxing incisions, with the aim of defining the bone fracture line. The chisel was positioned on the incision line surrounding the distal part of the second molar and was driven in deeper, as far as the distal limit of the relaxing incisions, in order to obtain a uniform bone/gingiva sliver. An incision was made in the distal portion of the gingiva to remove the sliver, while taking care to maintain an epithelial pedicle, to ensure better nutrition for the flap that would be used to cover the donor region.

16. Harvesting of bone marrow from the donor region to fill possible spaces between the bone sliver and the exposed spirals of the implant.
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Suturing and stabilization of the connective–bone sliver

17. Harvesting of gingival tissue between the double incisions, in the region of the papillae.
18. Detachment of the gingival flap to make it possible to reposition it coronally, free from tension.
19. Manipulation of the connective-bone graft to reproduce the shape of the socket defect.
20. Careful insertion of the connective-bone sliver, as far as the level of the implant platform, while leaving the bone portion in contact with the exposed spirals of the implant and the connective tissue portion in contact with the internal portion of the gingival flap.
21. Stabilization of the graft by means of suturing the connective tissue portion of the graft on the internal face of the gingival flap, and passive accommodation of the flap in order to suture the papillae using simple stitches.
22. Compaction of the bone marrow, that was taken from the maxillary tuberosity, between the vestibular surface of the implant and the graft, to ensure the final stabilization of the connective-bone sliver.
23. Insertion of the provisional crown on top of the implant.
24. Torque of 20 N on the attachment screw of the provisional crown and sealing of the palatine orifice with temporary filling material (Fermit, Ivoclar North America, Amherst, NY, USA).
25. Suturing of the gingival flap in the donor region, using simple stitches.

The patient was instructed not to place any loading on the region and to make topical applications of 0.12% chlorhexidine, three times a day for seven days.

After a three-month period to allow osseointegration and restoration of the bone and gingival architecture, an abutment made of Procera zirconia was inserted using torque of 20 N. At that time, the anatomical shape of the emergence profile obtained for the provisional crown on the implant was checked.
Transfer molding was performed on the implant emergence profile and on the neighboring teeth. The dies were scanned using Procera Forte to construct copings made of Procera alumina. After testing the copings, transfer molding was performed to enable personalized application of porcelain. The esthetic finish was determined according to the adequacy of the gingival architecture and the crowns made of metal-free porcelain on the anterior teeth of the maxilla.

This technique was shown to be esthetically and functionally predictable: it minimized the surgical trauma and provided favorable results immediately and after 24 months of follow-up.

Discussion

In addition to the initial stability, the biological changes that occur when an implant is put into use at an early stage are of great importance in the bone repair process (Romanos G 2001, Isidor F 2006, Vandamme K 2007).

In cases of tooth loss together with loss of support structures, the esthetic risks are greater. In such cases, implant insertion associated with grafting or with the use of membranes is indicated, so that function is reestablished only after a healing period.

Many papers have reported the use of grafts from the maxillary tuberosity for correcting socket defects (ten Bruggenkate CM 1992, Misch CE 1999, Rosa JCM 2008). The maxillary tuberosity presents the disadvantages of the limited quantity of bone material available, the low bone quality and the difficulty of surgical access. Its advantages are the greater speed of graft repair, the ease of harvesting the graft material and adapting it to the receptor region because of the bone malleability, and the excellent postoperative recovery (Rosa JCM 2008).

The bone and periosteal cells of the maxillary tuberosity behave as osteoprogenitor cells (Cicconetti A 2007). Knowing that the vascularization...
pattern is vital for bone grafting success, the trabecular nature of grafts harvested from the maxillary tuberosity means that such grafts have high revascularization capacity and release growth factors to the receptor site. Hence, these grafts need to be transported rapidly, so that their fundamental properties are not lost (Prolo & Rodrigo 1985, Goldberg & Stevenson 1987).

With the aim of shorter time required for bone-implant integration, it needs to be emphasized that spongy bone presents greater vascularization and cellularization and is metabolically more active, thus favoring the initial osseointegration processes.

Stabilization of the bone graft in the receptor site is fundamental for avoiding micromovement and facilitates the graft neovascularization process. Graft revitalization and incorporation success depend on close contact between the graft and the host’s vascular bed (Burchardt H 1983/1987, Abrektsson T 1980/1980, Gوردh & Alberius 1999).

With the passage of time and with appropriate mechanical stimulation, the mechanical resistance of the grafted area will tend to increase. Early low-intensity stimulation increases the local blood flow and contact osteogenesis (Burchardt & Enneking 1978, Abrektsson T 1980/1980).

The use of a connective-bone sliver harvested from the maxillary tuberosity was indicated for restoration of the vestibular bone cortex of the damaged fresh socket and covering the exposed spirals of the implant, and it increased the thickness and quality of the gingiva. In combination with coronal repositioning of a gingival flap, correction of the gingival recession was enabled.

The connective-bone sliver graft from the maxillary tuberosity impeded cell competition between the hard and soft tissues, thereby promoting effective bone and gingival healing.

Immediate loading of the implant following tooth extraction, associated with grafting of a connective-bone sliver avoids the need to subject the patient to several surgical procedures, such as bone grafting, gingival tissue grafting, implant insertion, surgery to reopen the implant and conditioning stages for the soft tissue surrounding the implant. This also avoids aesthetic impairment through performing these procedures.

This technique promoted acceleration of the bone repair around the implant and in the graft, thereby providing effective connective-bone healing in a single procedure.

**Conclusion**

The connective-bone sliver graft from the maxillary tuberosity was indicated for recovery of the vestibular bone cortex of the damaged fresh socket and covering of the exposed spirals of the implant, and it increased the thickness and quality of the gingiva. In combination with coronal repositioning of a gingival flap, correction of the gingival recession was enabled.

The connective-bone sliver graft from the maxillary tuberosity impeded cell competition between the hard and soft tissues, thereby promoting effective bone and gingival healing.

Immediate loading of the implant in the damaged fresh socket associated with a connective-bone sliver graft from the maxillary tuberosity enabled dental alveolar restoration in a single procedure. Although the technique requires long-term follow-up, the result obtained was satisfactory and promising.

The Literature list can be requested from the editorial office.

**Contact**

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Clinical outcomes of flapless implant surgery using a handpiece driven stereographic surgical guide

**Material and methods**

Following the anamnesis, clinical and radiologic examinations of the patients were completed. Patient consent form was taken for the use of a stereolithographic surgical guide and flapless surgery. Five patients were included for this study. Scan prosthesis were fabricated using duplication method and digital volumetric tomography (DVT) were used for 3-D imaging of the patient’s jaw anatomy. Implants were planned on the software according to the teeth and alveolar anatomy visualized on DVT images. Surgical guide and handpiece attachment were fabricated with a fast prototyping machine. A triangular lock mechanism adapted on surgical guide and handpiece were used for guiding the osteotomy drills.

### Table 1

Distortion values (mm) in periapical, panoramic and tomographic X-rays (Sonick et al. 1994)

<table>
<thead>
<tr>
<th></th>
<th>Mean distortion</th>
<th>Maximum detected distortion</th>
</tr>
</thead>
<tbody>
<tr>
<td>Periapical</td>
<td>1.9</td>
<td>5.5</td>
</tr>
<tr>
<td>Panaromic</td>
<td>3.0</td>
<td>7.5</td>
</tr>
<tr>
<td>Computerised Tomography</td>
<td>0.2</td>
<td>0.5</td>
</tr>
</tbody>
</table>

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Computer aided planning and stereolithography allows clinicians to transfer the virtual implant positions to patient via surgical guide. Studies were presented using stereolithography recently, however most of were specific to an implant brand or type only. Hence, drill guidance and multiple numbers of guides inherits certain limitations and risks. A new type of surgical guide (Ötede®, Aytasarım, Ankara, Turkey) eliminating drill guidance was presented in this study.

---

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**DDS, Phd, Assoc. Prof Dr, Istanbul University, Faculty of Dentistry, Department of Oral Implantology, Capa, Istanbul, Turkey
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during surgery. The guide is adapted on the mucosa in occlusion to the opposing dentition then secured with fixation screws. Following the removal of soft tissue with a punch drill, osteotomy was performed and total number of 72 implants was inserted by this method. Patients were screened following the surgery and a new DVT were obtained at the stage of loading for examining the accuracy of the implant positioning. Postoperative swelling was almost unnoticeable and the pain was reported only for the day of the surgery. All implants except 2 were osseointegrated at the end of 3 months. The method enables clinician to plan and place implants without the need of an exposed flap in selected cases however a large group of cases should be analyzed for the routine use of this guidance.

_Introduction_

Conventional implant planning mainly depends on panoramic X-rays. However certain drawbacks are present with panoramic and periapical X-rays such as distortion and two dimension restrictions. The difference in accuracy between different methods has been investigated and documented in a study by Sonick et al. (Table 1). Despite the acceptable mean distortion rate in panoramic and periapical X-rays maximum achieved distortion rates may lead to false proximity or intrusion to vital anatomic structures in rare occasions. Furthermore owing to the lack of anatomic references, visual planning and conventional implant placement is challenging especially in total edentulous cases. This type of a case is likely to end up with implants engaging the approximal regions with undesired emergence profile and screw access hole. As a result the case may be a failure especially in highly demanding patients with high lip line. Surgical guides can be used to overcome this problem to a certain degree however the exact 3-D positioning is unfortunately impossible with guides based on radiologic markers, impressions and intra-oral bone mapping techniques. Following the introduction of the computerized dental tomography, diagnosis and treatment planning can be visualized on three-dimensional (3-D) images. Further development and integration of DICOM data on the specific dental implant software enabled to construct a 3-D view of the alveolar bone and place implants in real dimension accuracy. By the introduction of volumetric tomography, dose concerns with conventional CT imaging were not an issue of concern. Because the dose with a DVT is equal or slightly higher than a conventional panoramic radiograph. The surgery can be virtually performed on the computer using realistic implant fixtures. In case of having a scan prosthesis the ideal implant position can be assigned in relation to bone height, width, angulation and emergence profile. Necessity of angled abutment and possible screw access points can also be planned in advance the fabrication of the final prosthesis. Stereolithography is a relatively new advancing technology in the dental field. Engineering part of this technology is still in demand of dental view and opinion. Most systems can only offer direction guidance with the surgical guides, however few of them offer depth control but only when certain brands of implants were used. The technique is time

Fig. 2. (a) Intraoral view of the maxilla, (b) duplication of the current prosthesis and preparation of the scan prosthesis with Br2S4, (c) scan prosthesis, (d) planning phase.
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consuming and costly at the moment and deviation rates is still an issue of concern. A new surgical guide (Ötede, Aytarasrim, Ankara, Turkey) is currently investigated in Istanbul University, Faculty of Dentistry, Department of Oral Implantology (Fig. 1). The main idea is to shift the guidance from drill to handpiece. A male-female triangular lock mechanism is placed to the guide. This enabled to drive the handpiece in the exact route and depth without drill dependency as seen in other systems like SimPlant, Aytasarim3D etc. The new system requires individual stereolitographic key for the handpiece to be used in the surgery (Fig. 2). The handpiece and drills were scanned before surgery to simulate depth control and drill guidance with the handpiece.

_Material and methods_

Following the anamnesis, clinical and oral examinations were performed for 5 patients with various edentoulism types. The patients were informed about the flapless option with guided surgery and written consent was taken. Patient’s current prosthesis was evaluated for the fit and positional accuracy. This prosthesis was duplicated in clinic with the negative impression technique. Cold polymerising acrylic was prepared with 10 per cent barium sulphate addition and mixed well then poured carefully into the drainage holes of the duplication mold. Following the setting, the mold was opened and the scan prosthesis is removed and trimmed and polished. The prosthesis is tried to ensure that the patient can wear it comfortably during tomographic scan. Any disturbing pieces or edges may lead patient to mis-position the prosthesis during scan causing error in the CT images (Fig. 2). The DICOM data is transferred to a personal computer. A 3-D implant image processing software (Implant3D, MediaLab, LaSpezia, Italy) was used to plan implants. Following the segmentation of the bone and prosthesis the implant placement is performed on the computer with guidance of the scan prosthesis.

Following the planning, the manufacturing of the surgical guide began. The system requires the scan of the handpiece head region. Drills to be used in the osteotomy were also required to be scanned for correct depth control. All components were tried before surgery to ensure the proper operation of the triangular lock mechanism.

_Surgery:_ The guide was tried once again for the accuracy of fit. Local anesthesia was administered with infiltration. Following the stabilization of the guide with fixation screws, handpiece lock was adapted to the handpiece. Mucotomy was performed with a punch drill. Routine osteotomy protocol was used using the triangular lock in the surgical guide. Depth control was also provided with the stop of the triangular lock. Following the insertion of the implants the guide was removed from the mouth and gingival formers are placed (Fig. 3). A total of 72 implants were placed in maxilla and mandible.

Postoperative follow-up: All patients were recalled after the surgery day and controlled everyday for a week. Then all patients were examined with one month intervals.

_Results:_ Two implants in a patient were lost in the post-op second week. Rests of the implants were osseointegrated at the end of healing period. No complications were observed during the healing period. Swelling was minimal due to the flapless surgery. Pain complication was also recorded to be minimal and mainly experienced in the day of surgery.

_Discussion_

As a result of the advances in fast prototyping and 3-D printing field, many stereolitography based surgical guide systems became available on the market for providing accurate positioning of dental implants. The procedure can be outlined as a 3-D engineering and many variations are present amongst systems. As a result the benefits vary depending on the design of the guide. All systems provide horizontal guidance but depth control is provided only in few systems intended for a specific implant system. The present surgical guide may use any type of implant drill after it has scanned. The top of the female lock mechanism attached on the surgical guide serves as a stopper for providing depth control. Drill guidance is the main base of direction in static surgical guides. To maintain the directional accuracy, most system provides se-
Sequentially increasing diameter drill tubes either fixed or changeable. Inclusion of multiple numbers of drill guides during the surgery increases the surgical time, risk of deviation and complexity of the surgery. Some systems overcome the problem of multiple guides by using changeable tubes. However small tubes can be difficult to mount and remove in the oral environment. These small tubes bring the risk of inhalation, loss, stuck in place etc. Use of a single guide in this study significantly decreased the surgical time and relevant complications. Furthermore drill guidance requires the inclusion of metal tubes for the drill to be thoroughly operated. This may significantly reduce the irrigation and arise the risk of overheating in the osteotomy site owing to the friction and restricted liquid flow way through the metal tubes. The system used in this study does not contain any metal tubes in the drill pathway and has wide holes in the field of osteotomy allowing higher level of irrigation liquid to cool the osteotomy site. It should be always kept in mind that the risk of overheating can be a cause of failure in surgical guide systems and flapless surgery. The drill rpm should be below 750 rpm with maximum possible irrigation. Drill independency makes this system virtually suitable for any implant system which may be an advantage when compared to drill and implant system based guides.

Conclusion

The new surgical guide in this study has promising features and advantages compared to similar stereolithographic surgical guide systems. Handpiece orientation adds additional comfort and freedom for the surgeon. Further analytic studies are required to evaluate the positional accuracy and additional benefits of this system._

References

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5. Platforms: Standard 4.8mm
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* The 4.1mm and 4.8mm SwissPlant implants
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Implant Direct’s ratchet, insertion tool and
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3. **Surface Options - 17 year history:**
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4. **Diameter Options:**
   Six Diameters from 3.2mmD to 5.7mmD in 0.5mmD increments.

5. **Conical Connection:**
   Lead-in Bevel above Internal Hex - 23 year history (Niznick #4,960,381)
   Color Coded for easy Identification; Platform Switching on 3.2mmD, 4.2mmD and 5.2mmD Implants.

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Prosthetic compatibility with Screw-Vent, BioHorizons and MIS Implants
Extraction, immediate implant placement and guided bone regeneration using a flapless approach

author_Barry K. Bartee, DDS, MD, USA

Introduction

Esthetic requirements for implant supported restorations include the presence of natural soft tissue contours, and to the extent possible, the full presence of the interdental papillae. Regardless of the technique used for tooth extraction, some degree of soft and hard tissue loss is inevitable. The use of guided tissue regeneration membranes over sockets has been shown to be one method to preserve bone after extraction, but most techniques involve the use of large flaps and even vertical incisions to achieve primary closure. Because of the unique features of Cytoplast® dense PTFE membranes and the ability of the membranes to remain exposed in the oral cavity without risk of infection, the soft tissue architecture, keratinized tissue width and position of the mucogingival junction adjacent to the socket can be preserved. Using the minimally invasive tunneling technique described in this article facilitates minimally invasive socket reconstruction avoiding vertical incisions and incision of the interdental papillae. The technique described can be used for immediate implant placement, as in this case, or for socket preservation using particulate graft material if a staged approach is desired.
Case Presentation

This is a 60-year-old female who presented with a crown-root fracture of a non-vital maxillary right central incisor. The crown was temporarily stabilized with composite resin bonded to the adjacent teeth (Fig. 1). Extraction of the tooth and immediate implant placement was planned. To minimize soft and hard tissue recession, a flapless, minimally invasive extraction technique was employed (Fig. 2). The tooth root was extracted using only an intrasulcular incision. A #15 blade was used to sever the periodontal ligament and create space for root luxation and elevation (Fig. 3). Next, a subperiosteal pocket was created on the buccal and palatal aspect of the socket using a micro periosteal elevator (Fig. 4). Following luxation and initial elevation of the root with the micro elevator, the tooth was removed with forceps (Fig. 5). The interdental papillae were carefully undermined and elevated. This can be done with a small periosteal elevator or curette (Fig. 6). All remaining soft tissue was removed from the interior and margins of the socket with a sharp curette (Fig. 7). The implant osteotomy was done in the standard fashion, with the implant being placed against the palatal wall of the socket (Fig. 8). The gap between the facial aspect of the implant and the buccal wall was filled with a combination of autogenous bone chips harvested from the implant osteotomy combined with allograft bone (Fig. 9). A textured, high-density PTFE barrier membrane is placed. The membrane is trimmed, then placed into the superioistepal pocket on the palatal aspect (Fig. 10). The membrane is then tucked under the facial flap (Fig. 11). Next, the membrane is tucked under the interdental papillae, taking care to keep the edge of the material a minimum of 1.0 mm away from adjacent tooth roots (Fig. 12). A single 3-0 PTFE suture is placed to further stabilize the membrane. The membrane is intentionally left exposed, as primary closure is not required in this technique (Fig. 13).

Figure 14 shows the surgical site at three weeks. The exposed membrane is easily removed by grasping with a tissue forcep. Topical anesthesia may be used, but local anesthesia is not necessary.

The site at six weeks after implant placement (three weeks after membrane removal), reveals keratinized mucosa forming across the former extraction site (Fig. 15).

Figure 16 shows the clinical view following placement of the implant abutment and acrylic provisional restoration.

Summary

The flapless technique described provides a minimally invasive approach to extraction with socket grafting or immediate implant placement. Because the interdental papilla remains intact, there is less disruption of blood supply. As a result, there is a greater potential for maintenance of soft tissue volume. In addition, the use of a dense PTFE membrane improves the predictability of immediate implant placement, excluding the requirement for primary closure and resultant disruption of soft tissue architecture.

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Precautions for using zirconia implant abutments

The introduction of zirconia to the dental field opened the design and application limits of all-ceramic restorations. Thanks to its high strength and fracture toughness long span posterior restorations are now possible with high accuracy and success rate. Additionally, its white color allowed better reproduction of the required color especially in the anterior zone. These properties made zirconia an interesting material for the construction of implant abutments and superstructures.

The fabrication of zirconia implant abutments utilizes state of the art CAD/CAM technology which uses patient’s models for the production of an individual customized abutment. Moreover, the CAD phase allows accurate positioning and angulation of the zirconia abutment ensuring obtaining the best esthetics.

The fabrication of zirconia implant abutments is complicated by the problem of providing adequate fixation to the implant body. For titanium abutments, the fixation screw exerts direct pressure on the abutment which in turn is provided with external or internal hex to provide connection with the implant body. On the other hand, zirconia is a brittle material and friction between the fixation screw and the internal surface of the ceramic abutment could produce high internal stresses which could lead to unexpected fracture.

This problem is solved by insertion of a friction fit internal metallic nut (Procera zirconia abutment for Strauman implants) which is equipped with an external hex for establishment of proper contact with the implant body (Fig. 1). Additionally, the fixation screw interlocks with the metallic nut during tightening procedure (Fig. 2). This design is associated with the problem of generation of internal stresses which could lead to unexpected fracture of the zirconia implant abutment. This article evaluated the fracture causes of several broken zirconia implant abutments. Fractographic analysis of the broken segments allowed recognition of the location and site of the critical fracture.

Case presentations

Case 1

A broken zirconia implant abutment was sent by the treating dentist for fracture analysis. Patients

Fig. 1 SEM image, 27x. Internal metallic nut which depends on friction fit inside the zirconia implant abutment.
Fig. 2 SEM image, 10x. The metallic fixation screw used to press on the metallic nut for providing adequate retention with the implant body.
Fig. 2a A broken maxillary premolar was extracted and implant was immediately inserted.
records indicated that the patient complained of loosening of the implant supported zirconia veneered crown (tooth 21). The treating dentist also complained that the internal metallic nut lost friction contact with the zirconia abutment and that he had to re-assemble the components before screw fixation. The abutment was fractured after two incidence of screw loosening. The same problem led to fracture of the second abutment after which the dentist decided to insert a titanium abutment. Scanning electron microscopic examination indicated that the abutment was fractured due to pressure from the metallic screw nut.

**The solution**

Once the metallic nut has lost friction fit with the zirconia abutment it can not be correctly reinserted inside the abutment and areas of friction contact between the improperly assembled components could lead to generation of high internal stresses causing fracture of the zirconia implant abutment as was reported for the two examined abutments. A new abutment should be used in such case.

**Case 2**

A broken zirconia implant abutment was examined. SEM analysis revealed that it was an angled abutment which corrected the tilt of an implant replacing a maxillary lateral incisor. The entire buccal wall was fractured beneath the temporarily cemented zirconia veneered crown.

**The solution**

Zirconia is a brittle ceramic material that must be used in adequate thickness to gain full potential of its high strength. A minimal wall thickness (0.5-0.7mm) is required in the entire structure of the zirconia implant abutment. This thickness must be increased in areas of high stresses to avoid unexpected fracture. Tilt correction—resulted in over reduction of the buccal wall (0.3 mm thickness) which resulted in fracture of the weakened segment. To reduce possibility of fracture, it is recommend to use a metallic abutment for correction of angle of insertion.

**Case 3**

A broken veneer porcelain from a Procera zirconia superstructure. This new design combines both the implant abutment and the framework of the restoration in one single structure thus reducing the number of components the dentist uses during the prosthetic phase.

This single component zirconia structure does not utilize an internal metallic nut for achieving contact with the implant body. On the contrary, this single component super structure utilizes directly the fixation screw to obtain direct fixation to the implant body.

**The solution**

Using single component superstructures has several advantages as it simplify the handling procedure, does not require anti-rotation feature, and reduces the num-
On the other hand, using single component zirconia implant superstructure which is composed of zirconia abutment and the framework as one component could facilitate easier handling and simplify insertion procedure due to reduction of the components used. Moreover, careful design consideration of the requirements of both the abutment and the zirconia framework is mandatory to ensure good function of each element. Lack of adequate support beneath the veneer ceramic or over reduction of the axial walls of the zirconia abutment could lead to unexpected fracture.

**References**


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Resorption mechanism of an injectable calcium phosphate bone regeneration cement—Radiographic and histological monitoring

author_Sérgio Alexandre Gehrke, Brazil

Introduction

Calcium phosphate cements have been extensively studied since the 80’s for their application as bone substitutes in various indications, mainly in the orthopaedic and odontology fields. They are well suited for osseous repair and reconstruction, because they proved to be biocompatible, osteoconductive and resorbable. However, their main interest lies in the fact that they are easy to apply in situ, can be shaped, and finally harden, thus providing a stability which is beneficial to the reconstruction site and the whole bone formation process.

Calcium phosphate bone cements are often known as products exhibiting very slow resorption. It was even shown that a particular apatite cement presents extremely limited resorption and osteoconduction after 12 months in a sheep model for craniofacial repair (Zins 2008). This slow resorption characterizing some calcium phosphate cements is partly due to the absence of macroporosity in the hardened cement. Nevertheless, it is important, when dealing with bone cements, to look into their composition once hardened. Most of the calcium phosphate bone cements consist of apatite after setting (hydroxyapatite or carbonated apatite), which is the most stable phase in vivo. Another type of calcium phosphate cement is the brushite cement. Brushite is the less stable calcium phosphate compound in vivo, and the only one to be soluble in blood serum. This means that the chemical nature of brushite cements (quickly resorbed) theoretically compensates their lack of macroporosity (slowing down the resorption rate). This difference in resorption process between apatite and brushite cement was clearly evidenced by Apelt and co-authors in a sheep model study (Apelt 2003).

As far as we know, most of the brushite cements are still in the development phase and only two of them are already on the market: one for orthopedic surgery (Jectos®, Kasios®, France), and one for dental applications. The latter is called PD VitalOs Cement® (Produits Dentaires SA, Switzerland). It has been specifically designed for dental surgery and offers the following practical advantages:

- It can be easily injected in cavities offering limited access;
- It is ready to inject: no preparation of the cement is needed during surgery;

Fig. 1 Pre-operative radiograph (case 1).
Fig. 2 Illustration of the installation of the first cement portion, against the inner wall of the sinus cavity.
Fig. 3 Sinus cavity completely filled with VitalOs, after installation of the implant.
It anchors well onto the bone and implant surfaces, providing suitable stability between bone and implant; it hardens, acquiring its own mechanical stability; it shows adequate radiopacity allowing radiographic monitoring of its resorption.

This paper aims at understanding the resorption process of PD VitalOs Cement and simultaneous bone neoformation, when the cement is used in osseous defects around implants. Two cases were selected to show how the cement degradation can be followed up through radiographic monitoring. To illustrate and provide explanation to some of the findings, results of an in vivo study run by the author are presented as well. In this study, the resorption of various bone substitutes and simultaneous bone neoformation were followed up with the help of fluorescent bone markers.

Case I

The first case is a 30-year-old male patient, in good health condition, non-smoker, presenting with absence of tooth 25, with pneumatization of the maxillary sinus. This situation required the regeneration of osseous tissue for the installation of a dental implant. As there was enough bone structure to provide the implant with primary stability, a sinus floor elevation was proposed, with installation of the implant in the same surgery.

Surgical protocol

After administration of a local anesthetic, a lateral access to the sinus cavity was performed through a lateral window. Then, the sinus membrane was carefully lifted up to the desired height. The drilling of the site was performed according to the recommendations of the manufacturer. During this phase, particular attention was paid to avoid perforating the membrane with the drills. A gaze humidified with a vasoconstrictor solution was placed into the sinus cavity for five minutes to control the bleeding as efficiently as possible. Indeed, blood flow must be avoided during injection of the PD VitalOs cement in order to allow contact between cement and bone and anchoring of VitalOs onto the sinus wall. Then, the cement was injected into the sinus to fill the space between the inner (nasal) wall of the cavity and the space prepared for the installation of the implant (Fig. 2). Once the cement was hard, the implant was installed as planned. Finally, the rest of the cavity was filled up with VitalOs up to the edges of the lateral window, without using a membrane. Indeed, the use of a membrane is not necessary with VitalOs, since the product acts itself also as a barrier.

Radiographic control

Thanks to the adequate radiopacity of the cement, it is easy to perform a radiographic follow-up and analysis of the resorption and new bone growth. Radiographs were therefore taken at 30, 60, 120 and 180 days after implant installation (Figs. 4–7), as well as 12 months (Fig. 8) after installation of the implant.
restoration, to follow-up the behavior of the new bone after loading of the implant.

**Case II**

The second case is a 40-year-old female patient in good health condition, non-smoking, presenting with a fracture of tooth 14 and presence of a large radicular cyst (Fig. 9) requiring removal of the root. The site presented a very large loss of bony structure, with fenestration of the buccal side of the socket wall. Therefore, the implant could only be stabilized in the inferior wall of the sinus cavity with the use of Summers’ osteotomes.

**Surgical protocol**

After administration of a local anesthetic, the access to the remaining portion of the root was performed and the root was carefully extracted, paying attention to preserve as much as possible the remaining osseous tissue. Thorough curettage of the apical portion was made to remove completely the cyst and the soft tissues. The implantation bed was then prepared with the osteotomes in order to gain sufficient stability. The implant was installed (Fig. 10) and the large bony defect was filled with the injectable VitalOs Cement (Fig. 11). At this stage it is important to control bleeding during and after injection until the cement is set. This is done with the help of a suction canula in the vicinity of the hardening cement. Although the cement itself promotes to a certain extent hemostasis in the site where it is injected, the bleeding occurring in the neighboring soft tissues must be controlled during the five minutes necessary for the cement to set.

**Radiographic control**

The control performed followed the same schedule described for the case I, i.e., radiographs were taken at 30, 60, 120 and 180 days after implant installation (Figs. 12–15), as well as 12 months (Fig. 16) after installation of the restoration.

**Histological study in animal model**

It is quite interesting to link the radiographic control of the presented cases with histological results obtained from a study where various bone substitutes were implanted in rabbit tibiae. The bone neoformation in the augmented cylindrical defects (4 mm diameter) was followed up by injecting polyfluorochromic bone markers during the first weeks following surgery: alizarin after 14 and 21 days, calcine after 28 and 35 days, tetracycline after 42 and 49 days. The animals were sacrificed at day 56. The investigated products were two granular bone substitutes (Geistlich Bio-Oss® and Straumann® Bone Ceramic) and a bone cement (PD VitalOs Cement®).

**Results**

The radiographs taken 30 days after surgery show the beginning of resorption at the periphery of the implants.
Legacy™3 Implant System
Conical Connection with Platform Switching
All-in-one Packaging Including Implant, Cover Screw, Healing Collar, Transfer and Final Preparable Abutment

THE NEXT GENERATION
in Implant Technology

FEATURES AND BENEFITS

1. **Body Design:**
   Evenly Tapered with self-tapping grooves from mid-point to apex.

2. **Thread Design:**
   Micro-threads with progressively deeper, double-lead, buttress threads.

3. **Surface Options - 17 year history:**
   SBM - medium rough texture; HA Coating below micro-threads.

4. **Diameter Options:**
   Six Diameters from 3.2mmD to 5.7mmD in 0.5mmD increments.

5. **Conical Connection:**
   Lead-in Bevel above Internal Hex - 23 year history (Niznick #4,960,381)
   Color Coded for easy Identification; Platform Switching on 3.2mmD, 4.2mmD and 5.2mmD Implants.

6. **Fixture-mount Packaging:**
   Fixture-mount standardizes insertion tools, serves as Transfer and can be shortened for use as Final Abutment.

Prosthetic compatibility with Screw-Vent, BioHorizons and MIS Implants
VitalOs cement. This process progresses towards the center of the cement mass. In the sinus lift case, the progress also takes place from the coronal towards the apical region around the implant (yellow arrows on Figure 4). Across the different times of analysis, it is possible to see that the material is gradually being replaced and the osseous tissue is being formed, apparently without incorporation of cement particles. At the end of the neoformation process, the newly formed bone displays radiographic characteristics which are similar to those of the native surrounding bone. This is also confirmed after 12 months of loading, where the bone around the implant has not resorbed, indicating a good preservation of the new osseous volume around the implant.

The results of the histological study show that the injectable brushite cement resorbs much more uniformly and rapidly than the granular materials (Figs. 17–19). It is also possible to state that the ossification process in the defects filled with VitalOs starts in the first weeks after augmentation: intensive brick-red areas due to alizarin indicate bone deposition between the second and the fourth week, and green areas due to calcein reveal the osteogenesis activity after 4–6 weeks. The coloration of the osteogenesis areas is more intense at every follow-up stage in the defects filled with the cement than in those filled with granular materials. This shows the maturity of the bone in the sites grafted with the brushite cement. Moreover, the organization of this newly formed bone tissue presents a structure which is more suitable for subsequent implant installation, i.e. a more organized and lamellar bone, with presence of osteons (yellow arrows, Fig. 17).

New bone formation is also seen in the defects grafted with granules. However, the bone deposition is stronger in the calcein phase (green fluorescence, see Figs. 18–19) than in the alizarin one (red), which indicates a delayed bone formation, starting four weeks after surgery. The fluorescence of the new bone tissue is weaker than in the sites filled with PD VitalOs Cement, which is a sign for lower maturity of the new bone tissue.

_Discussion_

The radiographic follow-up of the augmented sites shows clearly that the resorption takes place from the cement periphery towards its center until it is completely substituted by newly formed bone.

Brushite cements are known to resorb from their outer surface in contact with bone, towards the inner core of the injected mass. The resorption of brushite takes place through a combination of dissolution and cell-mediated processes. The whole process happens at the bone-cement interface, called resorption front (Theiss 2005, Lu 2002). This resorption front can always be easily located on radiographs and allows accurate determination of the moment when the cement is completely resorbed and replaced by bone. An earlier publication of the same author in the first implants edition of this year showed that the newly formed tissue does not contain any cement remnants and displays a lamellar, well organized structure (Gehrke 2009).

As we could observe in the radiographic sequences presented here, the injectable VitalOs cement presents a good radiopacity during the whole duration of the resorption process. The cement is easily located on the radiographs because it produces white homogeneous areas that distinguish clearly from the surrounding bone structure. This difference is easily recognizable until the very end of the resorption process. The radiopacity of this cement was shown to be equivalent to that of cortical bone (Pittet 2002). These features definitely facilitate the follow-up of its resorption and help the professional determining when the implant can be loaded.

Why is it so important to determine accurately the right time for loading the implant? There are several reasons for this. First, to make sure that the structure surrounding the implant is able to withstand the stresses applied through the crown-implant system. The residual bone used as bed for implantation provides the initial stability to the implant, but in most cases its quantity and structure are not sufficient to support the stresses applied on the implant once it is in function. The presence of newly formed bone around the implant is therefore re-
required to provide the required mechanical stability. Now, the time necessary for substitution of the graft material by newly formed bone may vary from one case to another. In a maxillary sinus as in case I, the cement generally resorbs completely in six months. Nevertheless, this period can be longer in older patients or in patients presenting metabolic disturbances. In these cases it is necessary to wait longer before starting the prosthetic phase.

In addition to that, this time period also depends on the volume or thickness of material applied: the larger it is, the longer it takes to resorb and be replaced by new bone tissue. These various factors influencing the resorption time of the graft material make it impossible to predict accurately when the augmented bone site will be ready to withstand the load applied onto the implant.

For all these reasons, the ease of location of the grafting material on radiographs is of great advantage: it allows following each patient individually to estimate the level of new bone reconstruction. This is important since this process is very much patient-dependent and should be assessed individually in order to choose the right time for installing the restoration. Finally, it is also the predictability of the treatment success which is increased through accurate radiographic monitoring.

The radiographic monitoring is more difficult to achieve with granular materials because their appearance on radiographs is closer to that of bone and the grafted area becomes eventually a mix of granules and bony tissue. To emphasize this, Figures 20 and 21 show radiographs of sites augmented with granular materials (Bone Ceramic), 180 days after augmentation and one year after crown installation, respectively. It can be seen that the newly formed bone is not as well defined as in the cases filled with the brushite cement (Fig. 15). The resorption mechanism of granular materials is also accompanied by new bone formation, but this takes place between the granules, most of them remaining eventually embedded in the new bone structure.

On the histologies, the bone markers showed that the substitution of the cement by bone starts during the first weeks after implantation. The bone tissue eventually obtained after eight weeks in the model is well organized, displaying a much more suitable structure than the bone growing in the defects filled with granular materials. The results obtained here are in accordance with results from literature showing that brushite cements exhibit fast resorption (Theiss 2005, Lu 2002), despite their lack of macroporosity. This good capacity of resorption and substitution by osseous tissue confirms the osteogenic potential of PD VitalOs Cement.

**Conclusion**

This paper evidences and discusses the different resorption kinetics and mechanisms occurring between a brushite calcium phosphate cement, VitalOs Cement, and granular bone substitutes. We showed that the good radiopacity of VitalOs allows more precise and secure monitoring of the resorption and substitution of the cement by new bone tissue. The neoformation kinetics is also patient-dependent and the ease to monitor radiographically the resorption process adds reliability to the overall treatment.

In summary, it was shown here that:

- The radiopacity of the cement allows easy and reliable monitoring of the resorption process;
- Osteogenesis starts earlier in the cement than in the granular substitutes presented here;
- The bone neoformation is more intensive in contact with VitalOs than with granular products;
- The cement is fully resorbed and its substitution by new bone tissue requires less time than with the granular substitutes.

_The literature list can be requested from the author._

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DGZI Adria Congress on the Isle of Vis

authors Rolf Vollmer, Rainer Valentin, Germany

The sixth VIS Congress was held on the Isle of Vis in the picturesque village of Komlás from July 12 to 13, 2009. The Adria Congress was the second meeting, which was organized together with DGZI (German Association for Dental Implantology). Komlás is only a two-hour flight from Germany, and another two hours by ferry from Split, and one soon stands in an untouched countryside, where time seems to stand still. This statement is certainly right with regard to the beautiful nature, but not when you have a look at the highly sophisticated scientific program! The program was dedicated to general dentistry, with special focus on dental implantology and oral surgery. Internationally well-known speakers such as Dr Rainer Valentin and Dr Rolf Vollmer of the DGZI Board of Directors, enhanced the scientific program. Dr Istvan Urban gave an impressive report on modern possibilities for bone regeneration. Dr Istvan Urban works at the Sankt Istvan Hospital in Budapest, and he carries out scientific research in cooperation with the Loma Linda University in California. The oral surgeon Dr Darko Slovša—the recently graduated first specialist in implantology of DGZI in Croatia—explained in detail the anatomical basics for a successful implantation, and showed the possibilities of how to avoid making mistakes. Added together, it was an outstanding program in a picturesque environment.

DGZI wants to express their special thanks for the excellent organization to Prof Dr Bozidara Pavelica of the University of Zagreb, Dr Matko Bôzic and Dr stom. Željko Ferić from Split, as well as to Dr Rainer Valentin, member of the DGZI Board of Directors. It was generally agreed to hold next year’s VIS Congress again on the occasion of the Corpus Christi holiday (June 3, 2010).

DGZI plans a stronger commitment next year, in order to offer a special curriculum during the Adria Congress, and in cooperation with the VIS Congress, to interested colleagues. As soon as we know more details, we will inform our members about it.
39TH
INTERNATIONAL
ANNUAL CONGRESS OF THE DGZI
Complications, Risks and Failures in the Oral Implantology

October 9–10, 2009, in Munich
Hilton Munich Park

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www.dgzi.de
www.event-dgzi.de

SPECIAL PODIUM
„Periimplantitis – a challenge of the future” with talk of experts
5th Arab German Implantology Meeting DGZI in Damascus

The Arab German Implantology Meeting DGZI was held this year from April 8–10. The fifth meeting was organized together with the Dental Association in Damascus and in cooperation with the 3rd Syrian Annual Implantology Meeting.

Because the first four Arab-German meetings in Dubai had been perceived as very successful, the neighbouring countries voiced the desire to now hold this meeting of highly qualified implantologists.

Thus, an agreement was reached with the Syrian Dental Association, with the result that the congress could be held in Damascus. Due to the interest of the Syrian Dental Association, DGZI decided to organize this year’s meeting, together with the local Dental Association, in the megacity of Damascus. It was the aim of the organizers to offer a qualified congress close to other home countries, e.g. for colleagues from Iraq. The success of the meeting confirms that this was the right decision. The Syrian Dental Association counts 16,000 members. At the moment there are 2,200 colleagues, who are working as dental specialists in different fields. Many of them obtained their dental education in Germany or in other European countries. Dr Safoh Al Buni is their acting President. As in former years, he invited the presidents of the dental associations of all Arab countries to this congress, in order to talk about future plans, and the foundation of an Arab Implantology Association. DGZI can attest to setting standards in this region.

The following colleagues flew in from Germany: The DGZI members of the board Dr Rolf Vollmer, Dr Rainer Valentin, and Dr Roland Hille. They were accompanied by the speakers Prof Dr Werner Götz, Dr Martina Vollmer, Dr Dr Werner Stermann, Prof Dr Maike Vesper, Dr Peuthen, Dr Manfred Schreiber, Dr Gerd Rosenkranz, and Dr Robert Laux. Well-known speakers from Lebanon, Jordan, Egypt, Sudan, and Saudi Arabia completed the program.

A complete dental implantology spectrum was covered, ranging from sinuslift to immediate loading, complications, smoking, bone substitute materials, and aesthetics. The conference room was fully occupied from morning until night with over 450 participants. There was enough time for discussions, and the speakers gave answers to all kinds of interesting questions even during the breaks. The joint evening events took place in typical Syrian style, and atmosphere. A dental fair completed the event, and everyone involved agreed that the success of the event will continue in the region into 2010.
We thank also especially Dr Iyad Ghoneim for his excellent support of the organisation committee.

**Information about Syria**

The Syrian Arab Republic is a state in the Middle East, which was of major importance in the ancient world that was especially characterized thanks to an early advanced civilization. Due to its location on the Mediterranean Sea, and the plateaus with its adjacent deserts inside the country, Syria covers a wide range of different terrains. One can walk on the old paths of the ancient world, and have a look at the history of civilized mankind on-site. The country’s surface extends over 185,000 km², there are 19 million inhabitants, and the majority of them speak Arabic. Since the seventies, Syria has to cope with rapid population growth, which causes many health and social problems. The relations with its direct neighbour Israel, and also with the United States of America are still very tense. Thanks to its oil resources, Syria is in a quite good situation, because the shortage of oil will lead to an enormous increase in value of "black gold" in the coming decades. Syria is certainly a country one should get to know.

**Damascus**

Damascus is often praised as the "lively", "the mysterious", "the pearl of the Middle East", and sometimes even "the mother of all cities". Damascus is the oldest human settlement. Visitors will find it one of the most fascinating cities on earth, with 4,000 years of history, and populated 12,000 years ago. The old Aramaic name of the city was "Darméseq", which means "place with abundant water". This allusion sounds a bit strange for a place that is situated in direct vicinity to the Arabian desert, but it refers to the fact that Damascus is located in one of the most fertile oases in the Middle East called the Oasis Ghouta region, a vital line, which is watered by the Barada river. One can visit the Damascus citadel, the former Roman main street Via Recto, the Christian Quarter with the famous House of Saint Ananias, the Chapel of Saint Paul, and of course the magnificent Umayyad Mosque, which once was planned to be the most wonderful and greatest Islamic mosque. In Damascus you can also observe something that has become very rare lately. Here, especially in the old part of the city, Muslims, Christians and Jews live together in peace. Churches and mosques stand side by side, and one is warmly welcomed by friendly, hospitable, and interested people. The city's atmosphere cannot be more oriental than it is already.

In the endless souqs of the old town one feels like being back in the times of the tales from 1001 Arabian Nights. One can admire the impressive caravanserais, visit the Hamams, pass by the madrasas, and then reach the city palaces. Lush gardens, luxury hotels, vivid coffee houses, and charming old houses make a visit to this city become a feast for the senses.

Our conclusion: A safe country, Arabic lifestyle, kind and helpful citizens and colleagues.
Prince Miteb bin Abdullah bin Abdulaziz, Deputy Commander of the National Guard for Military Affairs, patronized the 20th Saudi Dental Society and Second National Guard Conference and a Workshop themed ‘New Dental Era’ from April 20 to 22 in Riyadh.

Internationally renowned speakers with topics on Orthodontics, Dental Aesthetics, Dental Implants and Endodontics were the highlights of the scientific sessions. A special award was given to the best research project conducted by undergraduate and postgraduate students and another award for the best presentation in the poster session. Workshops were conducted by local and international speakers. International manufacturing dental and medical companies participated in the exhibition. A compilation of approved abstracts is published in the Journal of Dental Research under the IADR Saudi Arabian Section.

"The remarkable success of the last conference held in Jeddah of March 2008 is promised to be repeated, if not, paralleled as I invite you once again for this year’s conferences. This conference is accredited by the Saudi Commission for Health Specialties", said Prof Yousef F. Talic, President of the Saudi Dental Society. Through this outcome the National Guard Health Affairs and the Saudi Dental Society continued to contemplate the interest of the growing population of dentists in the Kingdom. The international conference was organized to keep our colleagues well-informed of the advances and challenges in the global society for this particular field of speciality. It offers a unique and dynamic experience for educational career advancement where topics of interest from various dental specialities were presented and discussed, Prof Yousef F. Talic summarized.
Dr Ali Al Ehaideb, Chairman, Dental Services—Central Region, National Guard Health Affairs pointed out:

The scientific lectures and continuing education (CE) courses are designed to widen and enhance our knowledge through the shared ideas and experiences of our distinguished international and local invited speakers who are well-respected and pioneers in their field of speciality. New approach, different techniques utilizing the state of the art modern technology waits for our colleagues. It will also highlight the display of modern instruments, supplies and equipment by internationally known participating dental & medical companies.

Thus, it is our privilege to encourage your participation for the success of this meaningful scientific event.

The German Association of Dental Implantology (DGZI) was invited as a cooperation partner and was represented by its 1. Vice-president and Treasurer Dr Rolf Vollmer as well as Dr Mazen Tamimi, the international representative of DGZI. Almost 1,500 dentist participated in the meeting and agreed on the high level of education in Saudi Arabia.

_Information Saudi–Arabia_

70 years ago the Kingdom of Saudi–Arabia was founded. On 23rd September, 1932, King Abd Al Asis III. Ibn Saud unified his Kingdoms Nejd and Hejaz to one big state. The constitution is based on the Quran.

Saudi Arabia is located on the Arabian Peninsula, and borders with Iraq, Jordan, Kuwait, Oman, Quatar, the United Arab Emirates, Yemen, the Red Sea, and the Persian Gulf. It is a desert country with extreme temperatures. From the middle of April until the middle of October daily temperatures of 45 °C and more can be reached. Whereas in the winter months January and February temperatures of 15 °C and less can be expected. At night the temperature in the desert can even be below the freezing point. The coastal zones have humid climate. Saudi Arabia is six times larger than Germany. Its size is approximately 2,15 million
square kilometres. Approximately 80 per cent of the Arabian Peninsula belongs to the kingdom, though only 20 million inhabitants live there, i.e. a quarter of the total German population. Colleagues from over 85 countries work as dentists. Education and training programs are highly ranked in Saudi Arabia. Every dentist has to prove his/her CME points regularly. A dentist is only allowed to place dental implants when he/she has obtained a so called license issued by the Ministry of Health.

Saudi Arabia is governed by an Islamic authoritarian monarchy under the reign of King Fahd bin Abd al-Aziz Al Saud, who is also the Prime Minister. The King and his widely ramified family stand at the top of the pyramid, followed by the tribal leaders and urban salesmen. The capital is the ancient Nejd capital Riyadh.

Women have to wear long, black garments, and they have to wear a veil over their face. Driving is forbidden for women. It is also forbidden to eat pork and drink alcohol throughout Saudi Arabia. The import of drugs is strictly forbidden, and will be punished with draconian penalty.

The prosperity of the country is due to its oil production since 1933. In 1960 Saudi Arabia founded the Organization of Petroleum Exporting Countries (OPEC), together with Iran, Iraq, Kuwait, and Venezuela. The aim of the coalition of the five most important oil exporting countries was a joint oil policy, and the stabilization of the world market price. Later on more oil producing countries joined this association. In the meantime Saudi Arabia has become the most important oil producing country in the world, and it controls approximately 26 per cent of all known oil reserves. Riyadh [Arabic “the gardens”] is the capital of the Kingdom of Saudi Arabia, and its same-named province. The city counts 4,6 million inhabitants. The city is the economic, administrative, and cultural center of the country. It hosts numerous universities, museums, and monuments. Riyadh is the seat of the Saudi Arabian government, the parliament, all national boards, and numerous diplomatic missions.

Riyadh has been the capital of Saudi Arabia since 1932 when the country gained its independence. From the historical point of view the capital is an important transit city for the Arabian area, the pilgrim routes to Mecca and Medina, which are the most significant Islamic pilgrimage sites. The main palace of the royal dynasty has been located in Riyadh since 1824.
## Selected Events 2009/2010

### September 2009

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<tr>
<td>September 2–5</td>
<td>FDI Singapore</td>
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<td>Web: <a href="http://www.fdiworlddental.org">www.fdiworlddental.org</a></td>
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<td>September 4–5</td>
<td>6th Forum of Innovations in Dentistry</td>
<td>Leipzig, Germany</td>
<td>Web: <a href="http://www.event-fiz.de">www.event-fiz.de</a></td>
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### October 2009

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<tr>
<td>October 9–10</td>
<td>39th International Congress of DGZI</td>
<td>Munich, Germany</td>
<td>Phone: +49-3 41/4 84 74-3 08 Fax: +49-3 41/4 84 74-2 90 Web: <a href="http://www.event-dgzi.de">www.event-dgzi.de</a></td>
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<tr>
<td>October 11–15</td>
<td>58th Annual Meeting of AAID</td>
<td>New Orleans, Louisiana, USA</td>
<td>Web: aaaid.com</td>
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<tr>
<td>October 16–17</td>
<td>4th International Computer Aided Implantology Academy Congress</td>
<td>Istanbul, Turkey</td>
<td>Web: <a href="http://www.tpidakademi.com">www.tpidakademi.com</a></td>
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<tr>
<td>October 23–25</td>
<td>1st European Implant Direct Congress</td>
<td>Barcelona, Spain</td>
<td>Web: <a href="http://www.implantdirect.eu">www.implantdirect.eu</a></td>
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### November 2009

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<tr>
<td>November 27 – December 2</td>
<td>GNYDM Greater New York Dental Meeting</td>
<td>New York, USA</td>
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### March 2010

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<tr>
<td>March 2–4</td>
<td>Biannual WFLD World Congress in Conjunction with UAE International Dental Conference &amp; Arab Dental Exhibition</td>
<td>Dubai, UAE</td>
<td>Web: <a href="http://www.aeedc.com">www.aeedc.com</a></td>
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### April 2010

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<tr>
<td>April 15–18</td>
<td>IDEM International Dental Exhibition and Meeting</td>
<td>Singapore</td>
<td>Web: <a href="http://www.idem-singapore.com">www.idem-singapore.com</a></td>
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Congratulations and Happy Birthday to all DGZI-members around the world.
Implant Direct

1st European Implant Direct Congress, October, 23–25 2009!

Implant Direct will hold its first European Congress in the beautiful and central Mediterranean city of Barcelona from October 23rd to 25th 2009. The congress will focus on the daily challenges in the dental practice with the lead title “Challenging conventional thinking in Implant Dentistry”. Outstanding clinical program with Live Surgeries. The scientific program features leading speakers such as Dr Gerald Niznick, Dr Marius Steigmann, Dr Aragoneses, Dr Fernando D’Avenia and many others, which will present the state of the art techniques of implant surgery and prosthetics. In addition, two live surgeries will be held at the leading dental training center COEC in Barcelona.

Dr Pedro Pena and Dr Marius Steigmann illustrating the latest techniques of soft tissue management and bone augmentation.

Case competition—Send us your best clinical studies!

Implant Direct has initiated a case competition contest in which doctors from across Europe are invited to submit their cases. The top three entrants will be invited to our 1st European Implant Direct Congress in Barcelona where their cases will be presented. The top ten cases will be published in the Implant Direct casebook as well as be submitted for publication in selected European Implant Journal. Learn, relax and enjoy with an entertaining social and cultural program! Join us during our congress and register as soon as possible as places are limited to 350 attendants. Registrations via www.implantdirect.eu, and tollfree infoline 00800 4030 4030.

Dentaurum

The first surgical tray for cleaning and disinfecting machines

In the field of implantology up to now, rotating instruments with angled shafts and other additional components required longer and more time-consuming cleaning treatments after implantation. In manual cleaning and disinfecting methods, each instrument must be treated individually. Until now, it was not possible to clean the implantological instruments directly in the surgical tray in an automatic process, due to the difficulty in immersing the instruments in sufficient amounts of water and cleaning liquid. Now the new tioLogic® easyClean is available to save time and improve hygiene. The rotating instruments and additional components are placed back into their relevant positions directly after use and remain there during the entire cleaning and disinfecting process. Their position within the tray enables them to be thoroughly immersed and rinsed. The concept for the ideal retention fixture for rinsing rotating instruments, was developed in collaboration with the companies Miele Professional and Dentaurum Implants GmbH. During this collaboration, the new Miele tray insert for general use in dental practices was developed and the specialised tioLogic® easyClean for the implant system from Dentaurum Implants evolved. The latter comprises all surgical instruments and all essential components required for preparing an implant site and for inserting tioLogic® implants. The components are arranged according to requirements during the course of surgery. The difference to other similar solutions is that every instrument is attached to a spring so that the contact surface area is minimal. This is why rose head burs and depth drills, surface cutter and conical formers can be entirely immersed in water and cleaning liquid. These can remain in their relevant positions throughout the whole treatment, including sterilisation. The Miele tray insert for dental practices comprises two areas with space for 30 drills with contra-angle handpiece ISO shafts and 15 positions for turbine shafts. This allows rotating instruments to be automatically treated whilst in the tray.

Dentaurum Implants GmbH
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Since our beginnings, we have always been focused on quality and innovation toward the battle against cross-contamination and infections.
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**Mono Drape “No Panic”**

Liquid and bacterial repellent patient drape with adhesive triangular shape 11 x 10 cm and integrated transparent plastic eye-shield. This mono drape, called “no panic”, offers to the patient the opportunity to have visible communication with the operator during any kind of oral surgery, implantology or maxillofacial surgery. The wide dimension of the drape makes it suitable for advanced surgery. Two velcro loops fix in a safety and easy way surgical instruments and aspirators, avoiding random contamination.

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Straumann

**Cheap implants: more expensive in the long run?**

In order to reduce costs, low-price manufacturers conduct little or no clinical research to prove that their implants will deliver long-term function and esthetics. They also do little or nothing to advance dental education. Neither do they do anything to develop innovations that enhance the standard of patient care. If they did, the price of their products would rise dramatically.

Dental patients are rarely able to make an informed choice about the products that go into their mouths. They do not know, for instance, if the implant design has been tested to ensure that it will not fail or lead to bone and soft tissue loss over time. Nor do they know what level of training their dentist has in implant dentistry. Patients are also unaware that the increase in low-cost manufacturers in recent years has led to a rise in the number of implants that need servicing but have to be removed because their origin cannot be traced.

Therefore, every patient should ask three important questions: ‘Has the implant system been thoroughly tested?’; ‘How long is it guaranteed?’ and ‘Will replacement parts be available in more than 10 years’ time?’. If not, there is a higher risk that the implant may have to be removed, which is unpleasant and costly.

Peace of mind comes at a price. The Straumann dental implant system is backed by long-term published clinical studies and a 10-year guarantee. It is supported by a broad education program for dental professionals so that patients receive a high standard of care. Premium solutions may appear expensive initially, but they could well be cheaper in the long run. The question is: Can doctors and patients afford to save in the short term?

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CAMLOG

**Bite registration posts for more safety even in difficult situations**

With the new bite registration posts, the CAMLOG® Implant System now offers even more options for fabrication of precision prosthetic restorations on implants. The CAMLOG bite registration posts have been developed for implant-supported measurement and transfer of the arch relations. They allow the fabrication of a precision bite registration even with limited occlusal space conditions and with inadequate residual teeth.

The color-coded bite registration posts have a prosthetic height of 8.1 mm. For bite registration, the bite registration caps are used or a splinted bite registration without caps is fabricated. Splinting the bite registration posts with plastic is recommended in particular for significantly reduced residual teeth or for edentulous patients in order to achieve optimized support of the bite registration. Splinting is also possible with divergent implants, which is achieved thanks to a shortened post/implant connection in comparison to the Tube-in-Tube™ connection. The bite registration posts have a reduced diameter shoulder support and are also suitable for platform switching in conjunction with CAMLOG® SCREWLINE implants (not for implant diameter 3.3 mm).

Thanks to the exact fit of all connecting parts, the CAMLOG bite registration posts ensure high precision when measuring and transferring the arch relations. They offer a strong plus for safety when determining the arch relations on implants even in difficult situations.

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**Sybron Implant Solutions**

**New sales and marketing subsidiary in Barcelona**

As a direct result of the outstanding sales development with the German-made Dental Implant Systems line BEGO Semados® implantology products and the constantly growing user base in Spain and Portugal, BEGO Implant Systems in Bremen, Germany, has decided to build up a new company in Spain, called BEGO Implant Systems Iberia, S.L.

“The new company has been formed to increase our market share and competitive strength in Spain and Portugal and to offer our customers a more direct interface to BEGO Implant Systems sales and support people working close to our customers in the Iberia market”, says Walter Esinger, General Manager of BEGO Implant Systems.

The logistic partner remains unchanged—MAB Dental/Dentimail will continue handling customers orders, as in the past. The direct sales representatives will visit customers personally and regularly in the future to introduce the BEGO Implant Systems product portfolio including the latest novelties from IDS 2009 in Cologne. The direct sales and support team in Spain and Portugal will be headed by Mrs Silvia Albaci. Silvia Albaci is already very well known in the Iberian Dental Implants Industry working previously for Klockner Dental Implants in Spain. General Manager of the new subsidiary will be Walter Esinger who is also General Manager of BEGO Implant Systems GmbH & Co. KG in Bremen, Germany. To celebrate the formation of the new company Silvia Albaci and her team recently come with some key users of the BEGO Implant System to a training course in Bremen and at the dental clinics of Dr. Stefan Ries, in Wertheim where they participated in the course “Implantology meets Nouvelle Cuisine”.

**Sybron Implant Solutions**

**The new Zirconium Abutment—for a naturally perfect result**

The Zirconium abutment is a partially standardized abutment designed for the Pitt-Easy implant system. With this prosthetic abutment, the user will be able to comply with high aesthetic demands for an implant-supported denture. The abutment consists of two parts, so that the modified Zirconium part is bonded with the titanium in processing. The patented Zirconium abutment consists of the following parts:

- abutment base made of titanium (grade 4)
- internal threads, preventing loss of the screw
- abutment blank made of high-quality TZP-A Zirconium oxide ceramics
- retaining screw for processing the abutment in the laboratory or in the dental practice (marking groove at the screw shaft)

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**Xign®**

The Allfit® Xign® implant system is designed for enossal dental implantations.

The Xign® implant system is suitable for single-step and dual-step implant protocols.

It is made of Grade 4 pure titanium.

Xign® implants are abraded with aluminium oxide and hot-etched and have an osmotically active nanocoating. All accessory parts are colour-coded.

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